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$ightharpoonup \underline{B}$ COMMISSION IMPLEMENTING REGULATION (EU) 2017/2470

of 20 December 2017

establishing the Union list of novel foods in accordance with Regulation (EU) 2015/2283 of the European Parliament and of the Council on novel foods

(Text with EEA relevance)

(OJ L 351, 30.12.2017, p. 72)

Amended by:

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		No	page	date
<u>M1</u>	Commission Implementing Regulation (EU) 2018/460 of 20 March 2018	L 78	2	21.3.2018
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► <u>M3</u>	Commission Implementing Regulation (EU) 2018/462 of 20 March 2018	L 78	11	21.3.2018
► <u>M4</u>	Commission Implementing Regulation (EU) 2018/469 of 21 March 2018	L 79	11	22.3.2018
► <u>M5</u>	Commission Implementing Regulation (EU) 2018/991 of 12 July 2018	L 177	9	13.7.2018
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► <u>M13</u>	Commission Implementing Regulation (EU) 2018/1133 of 13 August 2018	L 205	18	14.8.2018
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► <u>M21</u>	Commission Implementing Regulation (EU) 2018/2016 of 18 December 2018	L 323	1	19.12.2018
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► <u>M23</u>	Commission Implementing Regulation (EU) 2019/108 of 24 January 2019	L 23	4	25.1.2019
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► <u>M26</u>	Commission Implementing Regulation (EU) 2019/387 of 11 March 2019	L 70	17	12.3.2019
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► <u>M36</u>	Commission implementing Regulation (EU) 2019/1979 of 26 November 2019	L 308	62	29.11.2019
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► <u>M38</u>	Commission Implementing Regulation (EU) 2020/16 of 10 January 2020	L 7	6	13.1.2020
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► <u>M41</u>	Commission Implementing Regulation (EU) 2020/443 of 25 March 2020	L 92	7	26.3.2020
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► <u>M44</u>	amended by Commission Implementing Regulation (EU) 2021/1318 of 9 August 2021	L 286	5	10.8.2021
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11117	2020	L 250	•	7.0.2020
► <u>M50</u>	Commission Implementing Regulation (EU) 2020/1559 of 26 October 2020	L 357	7	27.10.2020
► <u>M51</u>	Commission Implementing Regulation (EU) 2020/1634 of 4 November 2020	L 367	39	5.11.2020
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► <u>M65</u>	Commission Implementing Regulation (EU) 2021/912 of 4 June 2021	L 199	10	7.6.2021
► <u>M66</u>	Commission Implementing Regulation (EU) 2021/1318 of 9 August	L 286	5	10.8.2021
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► <u>M77</u>	Commission Implementing Regulation (EU) 2022/168 of 8 February 2022	L 28	5	9.2.2022
► <u>M78</u>	Commission Implementing Regulation (EU) 2022/169 of 8 February 2022	L 28	10	9.2.2022

COMMISSION IMPLEMENTING REGULATION (EU) 2017/2470

of 20 December 2017

establishing the Union list of novel foods in accordance with Regulation (EU) 2015/2283 of the European Parliament and of the Council on novel foods

(Text with EEA relevance)

Article 1

Union list of authorised novel foods

The Union list of novel foods authorised to be placed on the market within the Union as referred to in Article 6(1) of Regulation (EU) 2015/2283 is hereby established and set out in the Annex to this Regulation.

Article 2

This Regulation shall enter into force on the twentieth day following that of its publication in the Official Journal of the European Union.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

ANNEX

UNION LIST OF NOVEL FOODS

Content of the list

- 1. The Union list shall consist of Tables 1 and 2.
- 2. Table 1 includes the authorised novel foods and contains the following information:
 - Column 1: Authorised novel food
 - Column 2: Conditions under which the novel food may be used. This column is further subdivided into two: Specified food category and Maximum levels
 - Column 3: Additional specific labelling requirements
 - Column 4: Other requirements
- 3. Table 2 includes the specifications on novel foods and contains the following information:
 - Column 1: Authorised novel food
 - Column 2: Specifications

Table 1: Authorised novel foods

Authorised novel food	Conditions under which the novel food may be used		Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
N-Acetyl-D-neur- aminic acid	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Infant and follow-on formulae as defined by Regulation (EU) No 609/2013 (1)	0,05 g/L of reconstituted formula	containing it shall be 'N-acetyl-D-neuraminic acid' Food supplements containing N-acetyl-D-neuraminic acid shall bear a statement that the food supplement		
	Processed cereal-based foods and baby foods for infants and young children as defined by Regulation (EU) No 609/2013	0,05 g/kg for solid foods	should not be given to infants, young children and children under 10 years of age where they consume breast milk or other foods with added <i>N</i> -acetyl-D-neuraminic		
	Foods for special medical purposes for infants and young children as defined by Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the infants and young children for whom the products are intended but in any case not higher than the maximum levels specified for the category mentioned in the table corresponding to the products.	acid within the same twenty four hour period.		
	Total diet replacement foods for weight control as defined by Regulation (EU) No 609/2013	0,2 g/L (drinks) 1,7 g/kg (bars)			
	Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014 (²)	1,25 g/kg			
	Unflavoured pasteurised and sterilised (including UHT) milk-based products	0,05 g/L			

Authorised novel food	Conditions under which the novel food may be used		Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Unflavoured fermented milk-based products, heat treated after fermentation, flavoured fermented milk products including heat-treated products	0,05 g/L (beverages) 0,4 g/kg (solids)			
	Dairy analogues, including beverage whiteners	0,05 g/L (beverages) 0,25 g/kg (solids)			
	Cereal bars	0,5 g/kg			
	Table top sweeteners	8,3 g/kg			
	Fruit and vegetable-based drinks	0,05 g/L			
	Flavoured drinks	0,05 g/L			
	Speciality coffee, tea, herbal and fruit infusions, chicory; tea, herbal and fruit infusions and chicory extracts; tea, plant, fruit and cereal preparations for infusions	0,2 g/kg			
	Food Supplements as defined in Directive 2002/46/EC (3)	300 mg/day for general population older than 10 years 55 mg/day for infants 130 mg/day for young children 250 mg/day for children between 3 to 10 years of age			
Adansonia digitata (Baobab) dried fruit pulp	Not specified		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Baobab fruit pulp'		

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	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Ajuga reptans extract from cell	Specified food category	Maximum levels			
	cultures	Food Supplements as defined in Directive 2002/46/EC	In line with normal use in food supplements of a similar extract of the flowering aerial parts of <i>Ajuga reptans</i>			
▼ <u>M77</u>						
	Akkermansia muci- niphila (pasteurised)	Foods for special medical purposes as defined under Regulation (EU) No 609/2013 for the adult population, excluding pregnant and lactating women	3.4×10^{10} cells/day	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'pasteurised Akkermansia muciniphila'.		Authorised on 1 Marc 2022. This inclusion is based on proprietar scientific evidence an scientific data protected is
		Food supplements as defined in Directive 2002/46/EC for the adult population, excluding pregnant and lactating women	3.4×10^{10} cells/day	The labelling of food supplements containing pasteurised Akkermansia muciniphila shall bear a statement that they should be consumed by adults only, excluding pregnant and lactating women.		accordance with Article 2 of Regulation (EU) 2015 2283. Applicant: A-Mansia Biotec S.A., rue Granbonpré, 11 Bâtiment H, 1435 Mont Saint-Guibert. Belgium During the period of dat protection, the novel foo pasteurised Akkermansi muciniphila is authorised to placing on the market withit the Union only by A-Mansi Biotech S.A., unless subsequent applicant obtain authorisation for the nove food without reference to the proprietary scientific dat protected in accordance with Article 26 of Regulatio (EU) 2015/2283 or with the agreement of Mansia Biotec S.A End date of the dat protection: 1 March 2027.

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
L-Alanyl-L- Glutamine	Specified food category	Maximum levels			
	Food Supplements as defined in Directive 2002/46/EC				
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013 excluding foods for infants and young children				
	Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen				
Algal oil from the microalgae <i>Ulkenia</i> sp.	Specified food category	Maximum levels of DHA	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Oil from the		
	Bakery products (breads, rolls and sweet biscuits)	200 mg/100 g	micro-algae Ulkenia sp.'		
	Cereal bars	500 mg/100 g			
	Non-alcoholic beverages (including milk based beverages)	60 mg/100 ml			

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
<u>M25</u>						
	Allanblackia seed oil	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
		Yellow fat spreads and cream based spreads	30 g/100 g	containing it shall be 'Allanblackia seed oil'		
		Mixtures of vegetable oils (*) and milk (falling under the food category: Dairy analogues, including beverage whiteners)	30 g/100 g			
		(*) Except olive oils and olive pomace oils as Regulation (EU) No 1308/2013.	defined in Part VIII of Annex VII of			
<u>M9</u>						
	Aloe macroclada Baker leaf extract	Specified food category	Maximum levels			
		Food Supplements as defined in Directive 2002/46/EC	In line with normal use in food supplements of the similar gel derived from Aloe vera (L.) Burm.			
	Antarctic Krill oil from Euphausia superba	Specified food category	Maximum levels of combined DHA and EPA	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Lipid extract from the crustacean Antarctic Krill (Euphausia superba)'		
		Dairy products except milk-based drinks	200 mg/100 g or for cheese products 600 mg/100 g			
		Dairy analogues except drinks	200 mg/100 g or for analogues to cheese products 600 mg/			

Authorised novel food	Conditions under which the no	Conditions under which the novel food may be used		Other requirements	► <u>M29</u> Data Protection ◀
	Non-alcoholic beverages Milk-based drinks Dairy analogue drinks	80 mg/100 ml			
	Spreadable fat and dressings	600 mg/100 g			
	Cooking fats	360 mg/100 ml			
	Breakfast cereals	500 mg/100 g			
	Bakery products (breads, rolls and sweet biscuits)	200 mg/100 g			
	Nutrition bars/cereal bars	500 mg/100 g			
	Food Supplements as defined in Directive 2002/46/EC	3 000 mg/day for the general population 450 mg/day for pregnant and lactating women			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀	
	Processed cereal-based food and baby food intended for infants and young children covered by Regulation (EU) No 609/2013	200 mg/100 ml				
	Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen					
	Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014					
Antarctic Krill oil rich in phosp- holipids from	Specified food category	Maximum levels of combined DHA and EPA	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Lipid extract			
noupias trom Euphausia superba	Dairy products except milk-based drinks	200 mg/100 g or for cheese products 600 mg/100 g	C d	(Euphausia superba)'		
	Dairy analogues except drinks	200 mg/100 g or for analogues to cheese products 600 mg/				
	Non-alcoholic beverages Milk-based drinks Dairy analogue drinks	80 mg/100 ml				
	Spreadable fat and dressings	600 mg/100 g				
	Cooking fats	360 mg/100 ml				
	Breakfast cereals	500 mg/100 g				
	Bakery products (breads, rolls and sweet biscuits)	200 mg/100 g				
	Nutrition bars/cereal bars	500 mg/100 g				

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Food Supplements as defined in Directive 2002/46/EC	3 000 mg/day for the general population 450 mg/day for pregnant and lactating women			
		Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
		Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			
		Processed cereal-based food and baby food intended for infants and young children covered by Regulation (EU) No 609/2013	200 mg/100 ml			
		Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen				
		Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014				
<u>M66</u>						
	Arachidonic acid-rich oil from	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	acid-rich oil from the fungus <i>Mortierella alpina</i>	Infant formula and follow-on formula as defined in Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013	containing it shall be 'Oil from Mortierella alpina' or 'Mortierella alpina oil'		
		Foods for special medical purposes for infants as defined in Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013			

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Argan oil from	Specified food category	Maximum levels	The designation of the novel food		
	Argania spinosa	As seasonings	Not specified	on the labelling of the foodstuffs containing it shall be 'Argan oil'		
		Food Supplements as defined in Directive 2002/46/EC	In line with normal food use of vegetable oils	and if used as seasoning 'Vegetable oil only for seasoning' shall be mentioned on the label		
<u>M69</u>	Astaxanthin-rich oleoresin from	Specified food category Food Supplements as defined in Directive	Maximum levels 40-80 mg/day of oleoresin, resulting	The designation of the novel food on the labelling of the foodstuffs		
	Haematococcus pluvialis algae	2002/46/EC, excluding infants, young children, children, and adolescents younger than14 years	in ≤ 8 mg astaxanthin per day	containing it shall be 'Astaxanthin- rich oleoresin from <i>Haematococcus</i> pluvialis algae' The labelling of food supplements containing Astaxanthin-rich oleoresin from <i>Haematococcus</i> pluvialis algae shall bear a statement that they should not be consumed by infants, children, and adolescents younger than 14 years.		
<u>M9</u>	Basil seeds (Ocimum basilicum)	Specified food category Fruit juice and fruit/vegetable blend beverages	Maximum levels 3 g/200 ml for addition of whole basil seeds (Ocimum basilicum)			
<u>M32</u>	Betaine	Specified food category	Maximum levels (⁷)	The designation of the novel food		Authorised on 22 Augus
	Betaine	Drink powders, isotonic and energy drinks intended for sportsmen Protein and cereal bars intended for sportsmen	60 mg/100 g 500 mg/100 g	on the labelling of the foodstuffs containing it shall be 'betaine'. The labelling of foods containing betaine shall bear a statement that		2019. This inclusion i based on proprietary scien tific evidence and scientifi data protected in accordance
		Meal replacements intended for sportsmen	20 mg/100 g	the foods should not be used if food supplements containing		with Article 26 of Regulation (EU) 2015/2283.
		Total diet replacement for weight control as defined under Regulation (EU) No 609/2013	500 mg/100 g (bar) 136 mg/100 g (soup) 188 mg/100 g (porridge) 60 mg/100 g (beverages)	betaine are consumed the same day.		Applicant: DuPont Nutritio Biosciences ApS, Lange brogade 1 Copenhagen k DK-1411, Denmark. Durin the period of data protection
		Foods for Special Medical Purposes as defined under Regulation (EU) No 609/2013 for adults	400 mg/day			the novel food betaine is authorised for placing of the market within the Unionly by DuPont Nutrition Biosciences ApS unless subsequent applicant obtain

▼<u>M32</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	►M29 Data Protection ◀
			,		·	authorisation for the novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of DuPont Nutrition Biosciences ApS, End date of the data protection: 22 August 2024.
▼ <u>M9</u>						
	Fermented black bean extract	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Fermented black bean (Soya) extract' or 'Fermented Soya extract'		
		Food Supplements as defined in Directive 2002/46/EC	4,5 g/day			
	Bovine lactoferrin	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Lactoferrin		
		Infant formula and follow-on formula as defined in Regulation (EU) No 609/2013 (ready to drink)	100 mg/100 ml	from cows' milk'		
		Foods on dairy basis intended for young children (ready to eat/drink)	200 mg/100 g			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Processed cereal food (solid)	670 mg/100 g			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	Depending on the needs of the individual up to 3 g/day			
	Beverages based on milk	200 mg/100 g			
	Powdered drink mixes based on milk (ready to drink)	330 mg/100 g			
	Beverages based on fermented milk (including yoghurt drinks)	50 mg/100 g			
	Non-alcoholic drinks	120 mg/100 g			
	Products based on yoghurt	80 mg/100 g			
	Products based on cheese	2 000 mg/100 g			
	Ice cream	130 mg/100 g			
	Cakes and pastries	1 000 mg/100 g			
	Candies	750 mg/100 g			
	Chewing gum	3 000 mg/100 g			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Bovine milk basic whey protein isolate	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		Authorised on 20 Novem 2018. This inclusion is bar
	Infant formulae as defined in Regulation (EU) No 609/2013 Follow-on formulae as defined in Regulation (EU) No 609/2013 Total diet replacement foods for weight control as defined by Regulation (EU) No 609/2013 Foods for special medical purposes as defined in Regulation (EU) No 609/2013 Food Supplements as defined in Directive 2002/46/EC	30 mg/100 g (powder) 3,9 mg/100 mL (reconstituted) 30 mg/100 g (powder) 4,2 mg/100 mL (reconstituted) 300 mg/day 30 mg/100 g (powder formula for infants during the first months of life until the introduction of appropriate complementary feeding) 3,9 mg/100 mL (reconstituted formula for infants during the first months of life until the introduction of appropriate complementary feeding) 30 mg/100 g (powder formula for infants when appropriate complementary feeding) 30 mg/100 g (powder formula for infants when appropriate complementary feeding is introduced) 4,2 mg/100 mL (reconstituted formula for infants when appropriate complementary feeding is introduced) 58 mg/day for young children 380 mg/day for children and adolescents from 3 to 18 years of age 610 mg/day for or children and adolescents from 3 to 18 years of age 610 mg/day for children and adolescents from 3 to 18 years of age 610 mg/day for adults	containing it shall be 'Milk whey protein isolate'. Food supplements containing bovine milk basic whey protein isolate shall bear the following statement: 'This food supplement should not be consumed by infants/children/ adolescents under the age of one/ three/eighteen (*) years' (*) Depending on the age group the food supplement is intended for.		on proprietary scient evidence and scientific of protected in accordance warticle 26 of Regulation (I 2015/2283. Applicant: Arr Protéines S.A.S., 19 bis, de la Libération 35-Saint-Brice-en- Cog France. During the period data protection the not food bovine milk bawhey protein isolate is at orised for placing on market within the Un only by Armor Protéis S.A.S. unless a subsequapplicant obtains authoration for the novel fewithout reference to proprietary scient evidence or scientific of protected in accordance warticle 26 of Regulation (I 2015/2283 or with agreement of Arr Protéines S.A.S. End date the data protection 20 November 2023.

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Buglossoides arvensis seed oil	Specified food category	Maximum levels of stearidonic acid (STA)	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Refined		
	Dairy products and analogues	250 mg/100 g	Buglossoides oil'		
		75 mg/100 g for drinks			
	Cheese and cheese products	750 mg/100 g			
	Butter and other fat and oil emulsions including spreads (not for cooking or frying purposes)	750 mg/100 g			
	Breakfast cereals	625 mg/100 g			
	Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children	500 mg/day			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013, excluding foods for special medical purposes intended for infants and young children	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			

▼M9

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Calanus finmarchicus oil	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Food supplements as defined in Directive 2002/46/EC	2,3 g/day	containing it shall be 'oil from Calanus finmarchicus (crustacean)'		
74					
Calcium fructoborate	Specified food category Food supplements as defined in Directive 2002/46/EC for the adult population, excluding food supplements for pregnant and lactating women	Maximum levels 220 mg/day	1. The designation of the novel food on the labelling of the foodstuffs containing it shall be 'calcium fructoborate'. 2. The labelling of food supplements containing calcium fructoborate shall bear a statement that those food supplements should not be consumed by population under 18 years of age and by pregnant and lactating women.		Authorised on 23 December 2021. This inclusion is basson proprietary scientification and protected in accordance with Article 26 of Regulation (EU) 2015/2283. Applicant: VDF Future Ceuticals, Inc., 300 Wester Momence, Illino 60954, the United States. During the period of daprotection, the novel for calcium fructoborate is authorised for placing on the market within the United States. The control of the calcium fructoborate is authorised for placing on the market within the United States. The control of the novel for calcium fructoborate is authorised for placing on the market within the United States. The control of the novel food without reference the proprietary scientification of the novel food without reference the proprietary scientification of the proprietary scientif

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	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	▶ <u>M29</u> Data Protection ◀
	Calcium L-Methylfolate	Specified food category	Maximum levels (expressed as folic acid)	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Calcium L-Methylfolate'		
		Foods for special medical purposes and total diet replacement for weight control as defined in Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013			
		Infant formulae and follow-on formula as defined by Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013			
		Processed cereal-based foods and baby foods for infants and young children as defined by Regulation (EU) No 609/2013				
		Food supplements as defined in Directive 2002/46/EC excluding food supplements for infants and young children	In accordance with Directive 2002/ 46/EC			
		Food fortified in accordance with Regulation (EC) No 1925/2006	In accordance with Regulation (EU) No 1925/2006			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Chewing gum base (monomethoxypoly- ethylene glycol)	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Gum base (including 1,3-butadiene, 2-methyl-		
	Chewing gum	8 %	homopolymer, maleated, esters with polyethylene glycol mono-Me ether)' or 'Gum base (including CAS No: 1246080-53-4)'		
Chewing gum base (Methyl vinyl ether-maleic	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Gum base		
anhydride copolymer)	Chewing gum	2 %	containing it shall be 'Gum base (including methyl vinyl ether-maleic anhydride copolymer)' or 'Gum base (including CAS No 9011-16-9)'		
Chia oil from Salvia hispanica	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Chia oil		
	Fats and oils	10 %	(Salvia hispanica)		
	Pure chia oil	2 g/day			
	Food Supplements as defined in Directive 2002/46/EC	2 g/day			

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	Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
▼ <u>M61</u>						
	Chia seeds (Salvia hispanica)	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	mspanica)	Bread products	5 % (whole or ground chia seeds)	containing it shall be 'Chia seeds (Salvia hispanica)'		
		Baked products	10 % whole chia seeds			
		Breakfast cereals	10 % whole chia seeds			
		Sterilised ready to eat meals based on cereal grains, pseudocereal grains and/or pulses	5 % whole chia seeds			
		Fruit, nut and seed mixes				
		Chia seeds as such				
		Confectionery (including chocolate and chocolate products), excluding chewing gums				
		Dairy products (including yoghurt) and analogues				
		Edible ices				
		Fruit and vegetables products (including fruit spreads, compotes with/without cereals, fruit-preparations to underlay or to be mixed with dairy products, fruit desserts, mixed fruits with coconut milk for a twin pot)				
		Non-alcoholic beverages (including fruit juice and fruit/vegetable blend beverages)				
		Puddings that do not require heat treatment at or above 120 °C in their manufacture, processing or preparation				

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► M29 Data Protection ◀
Chitin-glucan from Aspergillus niger	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
Aspergulus niger	Food Supplements as defined in Directive 2002/46/EC	5 g/day	containing it shall be 'Chitin-glucan from Aspergillus niger'		
Chitin-glucan	Specified food category	Maximum levels	The designation of the novel food		
complex from Fomes fomentarius	Food Supplements as defined in Directive 2002/46/EC	5 g/day	on the labelling of the foodstuffs containing it shall be 'Chitin-glucan from Fomes fomentarius'		
Chitosan extract	Specified food category	Maximum levels	The designation of the novel food on		
from fungi (Agaricus bisporus; Aspergillus niger)	Food Supplements as defined in Directive 2002/46/EC	In line with normal use in food supplements of chitosan from crustaceans	the labelling of the foodstuffs containing it shall be 'Chitosan extract from <i>Agaricus bisporus</i> ' or 'Chitosan extract from <i>Aspergillus niger</i> '		
Chondroitin	Specified food category	Maximum levels	The designation of the novel food		
sulphate	Food supplements as defined in Directive 2002/46/EC for adult population, excluding pregnant and lactating women	1 200 mg/day	on the labelling of the foodstuffs containing it shall be 'Chondroitin sulphate derived from microbial fermentation and sulphation'		
Chromium	Specified food category	Maximum levels of total chromium	The designation of the novel food		
Picolinate	Foods covered by Regulation (EU) No 609/2013	9/ 250 μg/day I	on the labelling of the foodstuffs containing it shall be 'Chromium Picolinate'		
	Foods fortified in accordance with Regulation (EC) No 1925/2006 (4)				
Chromium-	Specified food category	Maximum levels	The designation of the novel food on		
containing yeast (Yarrowia lipolytica) biomass	Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children	2 g/day for children from 3 to 9 years of age, resulting in 46 µg of chromium per day 4 g/day for children from 10 years of age, adolescents and adults, resulting in 92 µg of chromium per day	the labelling of the foodstuffs containing it shall be 'chromium-containing yeast (<i>Yarrowia lipolytica</i>) biomass' The labelling of food supplements containing chromium-containing yeast (<i>Yarrowia lipolytica</i>) biomass shall bear a statement that the food supplements should not be consumed by infants and young children (children under 3 years of age)/children from 3 to 9 years of age (12).		

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Cistus incanus L. Pandalis herb	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Cistus		
	Herbal infusions	Intended daily intake: 3 g herbs/day (2 cups/day)	incanus L. Pandalis herb'		
Citicoline	Specified food category	Maximum levels			
	Food Supplements as defined in Directive 2002/46/EC	500 mg/day			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	250 mg per serving and a maximum daily consumption level of 1 000 mg			
Clostridium butyricum	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Clostridium'		
	Food Supplements as defined in Directive 2002/46/EC	1,35 × 10 ⁸ CFU/day	butyricum MIYAIRI 588 (CBM 588)' or 'Clostridium butyricum (CBM 588)'		

	Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
▼ <u>M76</u>						
	Coffea arabica L. and/or Coffea canephora Pierre ex A.Froehner dried cherry pulp and its infusion (Traditional food from a third country)	Specified food category Coffee cherry pulp from Coffea arabica L. and/or Coffea canephora Pierre ex A.Froehner for the preparation of infusions	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'coffee cherry pulp' and/or 'cascara (coffee cherry pulp)', and/or 'coffee cherry pulp infusion' and/or 'coffee cherry pulp dried infusion'. If the product containing the novel food a participation of the food and the food a		
		Coffee, coffee and chicory extracts, instant coffee, tea, herbal- and fruit-infusions, coffee substitutes, coffee mixes and instant mixes for hot beverages (and their flavoured counterparts).		food contains more than 150 mg/l of caffeine (as such or after reconstitution), it shall be labelled with the following indication: 'High caffeine content. Not recommended for children or pregnant or breast-feeding women' in the same field of vision as the name of the food,		
		Flavoured and unflavoured non-alcoholic ready-to-drink beverages		followed by the caffeine content expressed in mg per 100 ml. Typical infusion preparations are prepared with up to 6 g of coffee cherry pulp per 100 ml of hot water (> 75 °C). For the coffee cherry pulp placed on the market as such for the preparation of infusions, instructions shall be given to the consumer on the preparation.		

▼M9

	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Specified food category	Maximum levels	The designation of the novel food		Authorised on 16 Ap
Cereal bars	0,20 g/100 g	containing it shall be 'D-ribose'. The labelling of foods containing D-ribose shall bear a statement that		2019. This inclusion based on proprieta scientific evidence as
Fine bakery wares	0,31 g/100 g			scientific data protected accordance with Article
Chocolate confectionery (excluding chocolate bars)	0,17 g/100 g	supplements containing D-ribose are consumed the same day.	supplements containing D-ribose are	of Regulation (EU) 2015 2283. Applicant: Bioenergy Lif
Milk-based drinks (excluding malts and shakes)	0,08 g/100 g			Science, Inc., 1384 Johnson St. N. Minneapolis, Minnesot
Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen, isotonic and energy drinks	0,80 g/100 g		55304 period the n author the m only Science subsec	55304, USA. During the period of data protection, the novel food D-ribose is authorised for placing on the market within the Union only by Bioenergy Life Science, Inc. unless a subsequent applicant obtains
Bars intended to meet the expenditure of intense muscular effort especially for sportsmen	3,3 g/100 g			
Meal replacement for weight control (as drinks)'	0,13 g/100 g			authorisation for the nov- food without reference the proprietary scientif
Meal replacement for weight control (as bars)	3,30 g/100 g			evidence or scientific dat protected in accordance wit
Confectionery	0,20 g/100 g		20	Article 26 of Regulation (EU 2015/2283 or with the agreement of Bioenergy Life
Tea and infusions (in powder form to be reconstituted)	0,23 g/100 g			Science, Inc. End date of the dat protection: 16 April 2024 (years).
	Cereal bars Fine bakery wares Chocolate confectionery (excluding chocolate bars) Milk-based drinks (excluding malts and shakes) Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen, isotonic and energy drinks Bars intended to meet the expenditure of intense muscular effort especially for sportsmen Meal replacement for weight control (as drinks)' Meal replacement for weight control (as bars) Confectionery Tea and infusions (in powder form to be	Cereal bars O,20 g/100 g Fine bakery wares O,31 g/100 g Chocolate confectionery (excluding chocolate bars) Milk-based drinks (excluding malts and shakes) Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen, isotonic and energy drinks Bars intended to meet the expenditure of intense muscular effort especially for sportsmen Meal replacement for weight control (as drinks)' Meal replacement for weight control (as bars) O,20 g/100 g Confectionery O,20 g/100 g Tea and infusions (in powder form to be 0,23 g/100 g	Cereal bars 0,20 g/100 g Fine bakery wares 0,31 g/100 g Chocolate confectionery (excluding chocolate bars) Milk-based drinks (excluding malts and shakes) Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen, isotonic and energy drinks Bars intended to meet the expenditure of sportsmen Meal replacement for weight control (as drinks)' Meal replacement for weight control (as bars) Confectionery 0,20 g/100 g on the labelling of the foodstuffs containing it shall be 'D-ribose'. The labelling of foods containing D-ribose shall bear a statement that the foods should not be used if food supplements containing D-ribose are consumed the same day. 0,08 g/100 g 0,80 g/100 g 3,3 g/100 g Meal replacement for weight control (as bars) 3,30 g/100 g Confectionery 0,20 g/100 g Tea and infusions (in powder form to be 0,23 g/100 g	Cereal bars 0,20 g/100 g Fine bakery wares 0,31 g/100 g Chocolate confectionery (excluding chocolate bars) Milk-based drinks (excluding malts and shakes) Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen. Meal replacement for weight control (as drinks) Meal replacement for weight control (as bars) Confectionery 0,20 g/100 g on the labelling of the foodstuffs containing is shall be 'D-ribose'. The labelling of foods containing D-ribose shall bear a statement that the foods should not be used if food supplements containing D-ribose are consumed the same day. 0,08 g/100 g 0,80 g/100 g 1,33 g/100 g Meal replacement for weight control (as drinks)' Meal replacement for weight control (as bars) 3,30 g/100 g Confectionery 0,20 g/100 g Tea and infusions (in powder form to be 0,23 g/100 g

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Extract of defatted cocoa powder	Specified food category	Maximum levels	Consumers shall be instructed not to consume more than 600 mg polyphenols corresponding to 1,1 g of		
	Nutrition bars	1 g/day and 300 mg polyphenols corresponding to not more than 550 mg of extract of defatted	extract of defatted cocoa powder per day		
	Milk based beverages	cocoa powder in one portion of food (or food supplement)			
	Any other foods (including food supplements as defined in Directive 2002/46/EC) which have become established vehicles for functional ingredients and which are typically positioned for consumption by health conscious adults				
Low fat cocoa	Specified food category	Maximum levels	Consumers shall be instructed not to consume more than 600 mg of cocoa flavanols per day		
	Foods including food supplements as defined in Directive 2002/46/EC	730 mg per serving and around 1,2 g/day			
Coriander seed oil from Coriandrum sativum	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Coriander		
	Food Supplements as defined in Directive 2002/46/EC	600 mg/day	seed oil'		

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		Authorised on 20 November 2018. This inclusion is based on proprietary scientific
	Food Supplements as defined in Directive 2002/46/EC for the adult population	350 mg/day	extract powder'		evidence and scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283.
					Applicant: Ocean Spray Cranberries Inc. One Ocean Spray Drive Lakeville- Middleboro, MA, 02349, USA.
					During the period of data protection the novel food, cranberry extract powder, is authorised for placing on the market within the Union only by Ocean Spray Cranberries Inc. unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of Ocean Spray Cranberries Inc.
					End date of the data protection: 20 November 2023.
	Authorised novel food Cranberry extract powder	Cranberry extract Specified food category Food Supplements as defined in Directive	Cranberry extract powder Specified food category Maximum levels Food Supplements as defined in Directive 350 mg/day	Cranberry extract powder Specified food category Food Supplements as defined in Directive 350 mg/day Maximum levels Maximum levels The designation of the novel food on the labelling of the foodstuffs containing it shall be 'cranberry extract powder'	Cranberry extract powder Specified food category Maximum levels The designation of the novel food on the labelling of the foodstuffs containing it shall be 'cranberry extract powder' Food Supplements as defined in Directive 350 mg/day

	Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Crataegus pinna- tifida dried fruit	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
		Herbal infusions	In line with normal food use of Crataegus laevigata	containing it shall be 'Crataegus pinnatifida dried fruit'		
		Jams and jellies in accordance with Directive 2001/113/EC (5)				
		Compotes				
	α-cyclodextrin	Not specified		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Alpha-cyclodextrin' or 'α-cyclodextrin'		
	γ-cyclodextrin	Not specified	Not specified			
▼ <u>M21</u>	Decorticated grains of Digitaria exilis (Kippist) Stapf (Traditional food from a third country)	Not specified		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'decorticated fonio (Digitaria exilis) grains'		
▼ <u>M9</u>						
	Dextran preparation produced	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	by Leuconostoc mesenteroides	Bakery products	5 %	containing it shall be 'Dextran'		

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Diacylglycerol oil of plant origin	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
piant origin	Cooking oils		containing it shall be 'Diacylglycerol oil of plant origin (at least 80 %		
	Fat spreads		diacylglycerols)'		
	Salad dressings				
	Mayonnaise				
	Meal replacement for weight control (as drinks)				
	Bakery products				
	Yoghurt type products				
Dihydrocapsiate (DHC)	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Dihydrocapsiate' Food supplements containing synthetic dihydrocapsiate will be labelled as 'not intended for children up to 4.5 years'		
(DHC)	Cereal bars	9 mg/100 g			
	Biscuits, cookies and crackers	9 mg/100 g			
	Rice based snacks	12 mg/100 g			
	Carbonated drinks, dilutable drinks, fruit juice based beverages	1,5 mg/100 ml	emilien up to 115 years		
	Vegetable drinks	2 mg/100 ml			
	Coffee based drinks, tea based drinks	1,5 mg/100 ml			
	Flavoured water — still	1 mg/100 ml			
	Precooked oatmeal cereal	2,5 mg/100 g			
	Other cereals	4,5 mg/100 g			
	Ice cream, dairy desserts	4 mg/100 g			
	Pudding mixes (ready to eat)	2 mg/100 g	7		

•	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
·		Products based on yoghurt	2 mg/100 g			
		Chocolate confectionery	7,5 mg/100 g			
		Hard candy	27 mg/100 g			
		Sugar-free gum	115 mg/100 g			
		Whitener/creamer	40 mg/100 g			
		Sweeteners	200 mg/100 g			
		Soup (ready to eat)	1,1 mg/100 g			
		Salad dressing	16 mg/100 g			
		Vegetable protein	5 mg/100 g			
		Ready to eat meals	3 mg/meal			
		Meal replacements for weight control	3 mg/meal			
		Meal replacement for weight control (as drinks)	1 mg/100 ml			
		Food Supplements as defined in Directive 2002/46/EC	3 mg/single intake 9 mg/day			
		Non-alcoholic powdered drink mixes	14,5 mg/kg equivalent to 1,5 mg/100 ml			
	Dried Euglena	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'dried biomass of Euglena gracilis algae'. The labelling of food supplements containing dried Euglena gracilis shall bear a statement that those food supplements should not be consumed by infants/children under		Authorised on 23 December
į	gracilis	Breakfast cereal bars, granola bars and protein bars	630 mg/100 g			2020. This inclusion is based on proprietary scientific evidence and scientific data protected in accordance with
		Yoghurt	150 mg/100 g			
		Yoghurt Beverages	95 mg/100 g			Article 26 of Regulation (EU) 2015/2283.
		Fruit and vegetable juices, nectars, fruit/vegetable blend beverages	120 mg/100 g			Applicant: Kemin Foods L.C., 2100 Maury Street
		Fruit-Flavoured Drinks	40 mg/100 g	3 years of age/children under 10 years of age/children and		Des Moines, IA 50317, USA.
		Meal replacement beverages	75 mg/100 g	adolescents under 18 years of		CO/I.
		Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants	100 mg/day for young children 150 mg/day for children from 3 to 9 years of age 225 mg/day for children from 10 years of age and adolescents (to 17 years of age) 375 mg/day for adults	age (¹²).		

▼<u>M52</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Total diet replacement for weight control as defined by Regulation (EU) No 609/2013	190 mg/meal			During the period of data protection, the novel food is authorised for placing on the market within the Union only by Kemin Foods L.C. unless a subsequent applicant obtains authorisation for that novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of Kemin Foods L.C. End date of the data protection: 23 December 2025.
▼ <u>M13</u>	Dried aerial parts of Hoodia parviflora	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'deied coriel		Authorised on 3 September 2018. This inclusion is based
		Food Supplements as defined in Directive 2002/46/EC for adult population	9,4 mg/day	containing it shall be 'dried aerial parts of Hoodia parviflora'		on proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283. Applicant: Desert Labs, Ltd Kibbutz Yotvata, 88820 Israel.

▼<u>M13</u>

	Authorised novel food Conditions under which the novel		vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
						During the period of data protection the novel food dried aerial parts of <i>Hoodia parviflora</i> is authorised for placing on the market within the Union only by Desert Labs, Ltd unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of Desert Labs, Ltd. End date of the data protection: 3 September 2023.
▼ <u>M9</u>	Dried extract of Lippia citriodora from cell cultures	Specified food category Food Supplements as defined in Directive 2002/46/EC	Maximum levels In line with normal use in food supplements of a similar extract from the leaves of Lippia citriodora	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'dried extract of <i>Lippia citriodora</i> from cell cultures HTN®Vb'		
	Echinacea angus- tifolia extract from cell cultures	Specified food category Food Supplements as defined in Directive 2002/46/EC	Maximum levels In line with normal use in food supplements of a similar extract from the root of Echinacea angustifolia			

▼ <u>M9</u>						
	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Echinacea purpurea extract from cell cultures	Specified food category Food Supplements as defined in Directive 2002/46/EC	Maximum levels In line with normal use in food supplements of a similar extract from florets within the flower head of Echinacea purpurea	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'dried extract of <i>Echinacea purpurea</i> from cell cultures EchiPure-PC TM '		
	Echium plan- tagineum oil	Specified food category	Maximum levels of stearidonic acid (STA)	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Refined		
		Milk-based products and drinkable yoghurt products delivered in a single dose	250 mg/100 g; 75 mg/100 g for drinks	echium oil'		
		Cheese preparations	750 mg/100 g			
		Spreadable fat and dressings	750 mg/100 g			
		Breakfast cereals	625 mg/100 g			
		Food supplements as defined in Directive 2002/46/EC	500 mg/day			
		Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
		Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			

▼<u>M50</u>

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
O Ecklonia cava phlorotannins	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Ecklonia cava Phlorotannins'.		
	Food supplements as defined in Directive 2002/46/EC intended for the general population, excluding children under the age of 12 years	163 mg/day for adolescents from 12 to 14 years of age	Food supplements containing <i>Ecklonia cava</i> phlorotannins shall bear the following statement:		
		230 mg/day for adolescents above 14 years of age 263 mg/day for adults	(a) This food supplement should not be consumed by children/adolescents under the age of twelve/fourteen/eighteen(*) years.		
			(b) This food supplement should not be consumed by persons with thyroid disease or by persons who are aware of or have been identified as being at risk of developing thyroid disease.		
			(c) This food supplement should not be consumed if other food supplements containing iodine are also consumed.		
			(*) Depending on the age group the food supplement is intended for.		

▼M9

<u>v19</u>					
Authorised novel food	Conditions under which the novel food may be used		Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
<u> 118</u>					
Egg membrane hydrolysate	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'egg		Authorised on 25 Novemb 2018. This inclusion is base on proprietary scientification
	Food Supplements as defined in Directive 2002/46/EC intended for the general adult population 450 mg/day	membrane hydrolysate'.		evidence and scientific da protected in accordance wi Article 26 of Regulation (E 2015/2283.	
					Applicant: Biova, LLC 5800 Merle Hay Rd, Sur 14 PO Box 394 Johnst 50131, Iowa USA. During the period of data protection the novel food expendence of the market within the Unit only by Biova, LLC. unless subsequent applicant obtain authorisation for the novel food without reference the proprietary scientific day to the control of the proprietary scientification of the protected in accordance with the control of the protected of Regulation (El 2015/2283) or with the agreement of Biova, LLC.
					End date of the da protection: 25 Novemb 2023

	Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Epigallocatechin gallate as a purified extract from green tea leaves (Camellia sinensis)	Specified food category	Maximum levels	The labelling shall bear a statement that consumers should not consume more than 300 mg of extract per day		
		Foods including food supplements as defined in Directive 2002/46/EC	150 mg of extract in one portion of food or food supplement	more than 500 mg of extract per day		
<u>M50</u>						
	L-ergothioneine	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'L-ergothioneine'		
		Alcohol-free beverages	0,025 g/kg			
		Milk-based drinks	0,025 g/kg			
		'Fresh' milk products(*)	0,040 g/kg			
		Cereal bars	0,2 g/kg			
		Chocolate confectionery	0,25 g/kg			
		Food supplements as defined in Directive 2002/46/EC	30 mg/day for general population (excluding pregnant and lactating women)			
			20 mg/day for children older than 3 years			
		(*) When used in milk products L-ergothioneir any milk constituent	ne may not replace in whole or in part,			

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
▼ <u>M50</u>						
	Extract of three herbal roots	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	(Cynanchum wilfordii Hemsley, Phlomis umbrosa Turcz. and Angelica gigas Nakai)	Food supplements as defined in Directive 2002/46/EC for adult population	175 mg/day	containing it shall be 'extract of three herbal roots (Cynanchum wilfordii Hemsley, Phlomis umbrosa Turcz. and Angelica gigas Nakai)'.		
				The labelling of food supplements containing the extract of mixture of the three herbal roots shall bear a statement in close proximity to the list of ingredients indicating that it should not be consumed by individuals with known celery allergy.		
<u>M9</u>						
	Ferric Sodium EDTA	Specified food category	Maximum levels (expressed as anhydrous EDTA)	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Ferric		
		Food supplements as defined in Directive 2002/46/EC	18 mg/day for children	Sodium EDTA'		
			75 mg/day for adults			
		Foods covered by Regulation (EU) No 609/2013	12 mg/100 g			
		Foods fortified in accordance with Regulation (EC) No 1925/2006				
	Ferrous ammonium	Specified food category	Maximum levels	The designation of the novel food		
	Phosphate Food supplements as define 2002/46/EC	Food supplements as defined in Directive 2002/46/EC	Directive 2002/46/EC, Regulation (EU) No 609/2013 and/or Regu-	on the labelling of the foodstuffs containing it shall be 'Ferrous ammonium phosphate'		
		Foods covered by Regulation (EU) No 609/2013				
		Foods fortified in accordance with Regulation (EC) No 1925/2006				
		-				

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► M29 Data Protection ◀
Fish peptides from Sardinops sagax	Specified food category	Maximum levels fish peptide product	The designation of the novel food on the labelling of the foodstuffs		
Surumops sugux	Foods based on yoghurt, yoghurt drinks, fermented milk products, and powdered milk	0,48 g/100 g (ready to eat/drink)	containing it shall be 'Fish (Sardinops sagax) peptides'		
	Flavoured water, and vegetable-based drinks	0,3 g/100 g (ready to drink)			
	Breakfast cereals	2 g/100 g			
	Soups, stews and soup powders	0,3 g/100 g (ready to eat)			
Flavonoids from Glycyrrhiza glabra	Specified food category	Maximum levels of flavonoids from Glycyrrhiza glabra	1. The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Flavonoids from <i>Glycyrrhiza glabra L</i> .' 2. The labelling of the foods where the product was added as a novel food ingredient shall bear a statement that: (a) the product should not be consumed by pregnant and breast feeding women, children and young adolescents; and (b) people taking prescription drugs should only consume the product under medical supervision;	Beverages containing flavonoids shall be presented to the final consumer as single portions.	
	Beverages based on milk	120 mg/day			
	Beverages based on yoghurt	_			
	Beverages based on fruit or vegetables				
	Food Supplements as defined in Directive 2002/46/EC	120 mg/day			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013	120 mg/day			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	120 mg/day			
			(c) a maximum of 120 mg of flavonoids per day should be consumed.		
			3. The amount of flavonoids in the final food shall be indicated on the labelling of the food containing it.		

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
▼ <u>M40</u>	Fruit pulp, pulp juice, concentrated pulp juice from Theobroma cacao L. (Traditional food from a third country)	Not specified	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'cocoa (<i>Theobroma cacao</i> L.) pulp', 'cocoa (<i>Theobroma cacao</i> L.) pulp juice' or 'cocoa (<i>Theobroma cacao</i> L.) concentrated pulp juice'			
▼ <u>M9</u>			depending on the form used.			
	Fucoidan extract from the seaweed Fucus vesiculosus	Specified food category Foods including food supplements as defined in Directive 2002/46/EC for the general population	Maximum levels 250 mg/day	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Fucoidan extract from seaweed <i>Fucus vesiculosus</i> '.		
	Fucoidan extract from the seaweed Undaria pinnatifida	Specified food category Foods including food supplements as defined in Directive 2002/46/EC for the general population	Maximum levels 250 mg/day	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Fucoidan extract from seaweed Undaria pinna- tifida'		
	2'-Fucosyllactose	Specified food category Unflavoured pasteurised and sterilised (including UHT) milk-based products Unflavoured fermented milk-based products	Maximum levels 1,2 g/l 1,2 g/l beverages 19,2 g/kg products other than beverages	 The designation of the novel food on the labelling of the foodstuffs containing it shall be '2'-fucosyllactose'. The labelling of food supplements containing 2'-fucosyllactose shall bear a statement that the supplements should not be used if other foods with 		
		Flavoured fermented milk-based products including heat-treated products	1,2 g/l beverages 19,2 g/kg products other than beverages	added 2'-fucosyllactose are consumed the same day.		
		Dairy analogues, including beverage whiteners	1,2 g/l beverages			

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		12 g/kg for products other than beverages	3. The labelling of food supplements containing 2'-fuco-		
		400 g/kg for whitener	syllactose intended for young children shall bear a statement that the supplements should not be used if breast milk or other		
	Cereal bars	12 g/kg	lactose are consumed the same day.		
	Table-top sweeteners	200 g/kg			
	Infant formula as defined in Regulation (EU) No 609/2013	1,2 g/l alone or in combination with up to 0,6 g/l of lacto- <i>N</i> -neotetraose at a ratio of 2:1 in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			
	Follow-on formula as defined in Regulation (EU) No 609/2013	1,2 g/l alone or in combination with up to 0,6 g/l of lacto- <i>N</i> -neotetraose at a ratio of 2:1 in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			
	Processed cereal-based food and baby food for infants and young children as defined in Regulation (EU) No 609/2013	12 g/kg for products other than beverages			
		1,2 g/l for liquid food ready for use, marketed as such or reconstituted as instructed by the manufacturer			

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Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Milk-based drinks and similar products intended for young children	1,2 g/l for milk-based drinks and similar products added alone or in combination with up to 0,6 g/l lacto- <i>N</i> -neotetraose, at a ratio of 2:1 in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013	4,8 g/l for drinks			
		40 g/kg for bars			
	Bread and pasta products bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014	60 g/kg			
	Flavoured drinks	1,2 g/l			
	Coffee, tea (excluding black tea), herbal and fruit infusions, chicory; tea, herbal and fruit infusions and chicory extracts; tea, plant, fruit and cereal preparations for infusions, as well as mixes and instant mixes of these products	9,6 g/l — the maximum level refers to the products ready to use			
	Food supplements as defined in Directive 2002/46/EC, excluding food supplements for	3,0 g/day for general population			
	infants	1,2 g/day for young children			

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
<u> 136</u>						
	2'-Fucosyllactose/ Difucosyllactose	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		Authorised on 19.12.2019 This inclusion is based or
	mixture ('2'-FL/ DFL')	Unflavoured pasteurised and unflavoured sterilised (including UHT) milk products	2,0 g/L	containing it shall be '2'-Fucosyllactose/Difucosyllactose mixture'. The labelling of food supplements containing the 2'-Fucosyllactose/Difucosyllactose mixture shall bear	lactose/Difucosyllactose mixture'.	proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283.
	(microbial source)	Unflavoured fermented milk-based products	2,0 g/L (beverages)			
			20 g/kg (products other than beverages)			Applicant: Glycom A/S Kogle Allé 4, DK-2970 Hørsholm, Denmark. During
		Flavoured fermented milk-based products including heat-treated products	2,0 g/L (beverages) 20 g/kg (products other than beverages)	and/or Difucosyllactose are consumed the same day.		the period of data protection the novel food 2'-Fucosyl lactose/Difucosyllactose mixture is authorised fo placing on the marke
		Beverages (flavoured drinks)	2,0 g/L	-		within the Union only by Glycom A/S, unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific
		Cereal bars	20 g/kg			
		Infant formula as defined under Regulation (EU) No 609/2013	1,6 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer		evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of Glycom A/S.	
		Follow-on formula as defined under Regulation (EU) No 609/2013	1,2 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			End date of the date protection: 19.12.2024.
		Processed cereal-based food and baby food for infants and young children as defined under Regulation (EU) No 609/2013	1,2 g/L (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			
			10 g/kg for products other than beverages			

▼<u>M36</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Total diet replacement foods for weight control as defined under Regulation (EU) No 609/2013	4,0 g/L (beverages) 40 g/kg (products other than beverages)			
		Food for special medical purposes as defined under Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
		Food Supplements as defined in Directive 2002/46/EC intended for the general population excluding infants	4,0 g/day			
▼ <u>M56</u>		Milk-based drinks and similar products intended for young children	1,2 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			
▼ <u>M72</u>						
	3-Fucosyllactose (3-FL)	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		Authorised on 12 December 2021. This inclusion is based
	(microbial source)	Unflavoured pasteurised and unflavoured sterilised (including UHT) milk products	0,85 g/L	containing it shall be '3-Fucosyllactose'.		on proprietary scientific evidence and scientific data protected in accordance with
		Unflavoured and flavoured fermented milk- based products including heat-treated products	0,5 g/L (beverages) 5,0 g/kg (products other than beverages)	The labelling of food supplements containing 3-Fucosyllactose (3-FL) shall bear a statement that they should not be consumed:		Article 26 of Regulation (EU) 2015/2283.
		Dairy analogues	0,85 g/L (beverages) 8,5 g/kg (products other than beverages)	a) if foods containing added 3-Fucosyllactose are consumed on the same day;b) by infants and children under 3 years of age.		

▼<u>M72</u>

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Flavoured drinks, energy and sports drinks	1,0 g/L			Applicant: DuPont Nutrition & Biosciences ApS Lange-
	Cereal bars	30,0 g/kg			brogade 1, 1001 Copenhagen K, Denmark. During the
	Infant formula as defined under Regulation (EU) No 609/2013	0,85 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			period of data protection, the novel food 3-Fucosyl- lactose is authorised for placing on the market within the Union only by
	Follow-on formula as defined under Regulation (EU) No 609/2013	0,85 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			DuPont Nutrition & Biosciences ApS, unless subsequent applicant obtain authorisation for the nove food without reference to the proprietary scientification.
	Milk-based drinks and similar products intended for young children	0,85 g/L (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of DuPont
	Processed cereal-based food and baby food for infants and young children as defined under Regulation (EU) No 609/2013	0,3 g/L (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			Nutrition & Biosciences ApS. End date of the data protection: 12 December
		3,0 g/kg for products other than beverages			2026.
	Total diet replacement foods for weight control as defined under Regulation (EU)	2,0 g/L (beverages)			
	No 609/2013	30,0 g/kg (products other than beverages)			
	Foods for special medical purposes as defined under Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Food Supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children	5,0 g/day			

▼M9

<u>17</u>					
Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
<u>64</u>					
Galacto-oligos- accharide	Specified food category	Maximum levels (expressed as ratio kg galacto-oligosaccharide/kg final food)			
	Food Supplements as defined in Directive 2002/46/EC	0,333			
	Food supplements as defined in Directive 2002/46/EC, excluding infants and young children	0,450 (corresponding to 5,4 g galacto-oligosaccharide/serving; maximum 3 servings/day up to a maximum of 16,2 g/day)			
	Milk	0,020			
	Milk drinks	0,030			
	Meal replacement for weight control (as drinks)	0,020			
	Dairy analogue drinks	0,020			
	Yoghurt	0,033			
	Dairy based desserts	0,043			

▼<u>M64</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
_		Frozen dairy desserts	0,043			
		Fruit drinks and energy drinks	0,021			
		Infant meal replacement drinks	0,012			
		Baby juice	0,025			
		Baby yogurt drink	0,024			
		Baby dessert	0,027			
		Baby snack	0,143			
		Baby cereals	0,027			
		Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen	0,013			
		Juice	0,021			
		Fruit pie filling	0,059			
		Fruit preparations	0,125			
		Bars	0,125			
		Cereals	0,125			
		Infant formula and follow-on formula as defined in Regulation (EU) No 609/2013	0,008			
<u>19</u>						
(Glucosamine HCl	Specified food category	Maximum levels			
		Food Supplements as defined in Directive 2002/46/EC	In line with normal food use of glucosamine from shell fish			
		Foods covered by Regulation (EU) No 609/2013				
		Meal replacement for weight control				

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► M29 Data Protection ◀
	Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen				
	Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014				
Glucosamine	Specified food category	Maximum levels			
sulphate KCl	Food Supplements as defined in Directive 2002/46/EC	In line with normal food use of glucosamine from shell fish			
Glucosamine	Specified food category	Maximum levels			
sulphate NaCl	Food Supplements as defined in Directive 2002/46/EC	In line with normal food use of glucosamine from shell fish			
Guar Gum	Specified food category	Maximum levels	1. The designation of the novel food		
	Fresh dairy products such as yogurts, fermented milks, fresh cheeses and other dairy-based desserts.	1,5 g/100 g	on the labelling of the foodstuffs containing it shall be 'Guar Gum'. 2. A specific mention of the possible risks of digestive discomfort linked to the exposure of children aged under 8 to guar gum must be visible on		
	Fruit or vegetable-based liquid foodstuffs (of the 'smoothie' variety)	1,8 g/100 g			
	Fruit or vegetable-based compotes	3,25 g/100 g			
	Cereals accompanied by a dairy product, in packaging containing two compartments	10 g/100 g in the cereals None in the accompanying dairy product 1 g/100 g in the product when ready to eat	the label of any foodstuffs containing it. For example, 'Excessive consumption of these products may cause digestive discomfort, especially for children under 8 years of age'. 3. In the case of products with two compartments containing dairy and cereal products respectively, the instructions for use must clearly specify the need to mix the cereal and the dairy product		

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
			before consumption, in order to take into account the potential risk of gastro-intestinal obstruction.		
Heat-treated milk products fermented	Specified food category	Maximum levels			
with Bacteroides cylanisolvens	Fermented milk products (in liquid, semi-liquid and spray-dried powder forms)				
Hydroxytyrosol	Specified food category	Maximum levels	The designation of the novel food on the labelling of the food		
	Fish and vegetable oils, (except olive oils and olive pomace oils as defined in Part VIII of Annex VII of Regulation (EU) No 1308/2013 (6)), placed as such on the market	0,215 g/kg	products containing it shall be 'hydroxytyrosol'.		
			The labelling of the food products containing hydroxytyrosol shall		
	Spreadable fats as defined in Part VII of Annex VII of Regulation (EU) No 1308/2013, placed as such on the market	0,175 g/kg	containing hydroxytyrosol shall bear the following statements: (a) This food product should not be consumed by children under the age of three years, pregnant women, and lactating women; (b) This food product should not be used for cooking, baking or frying'		
ce Structuring	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
HPLC 12	Edible ices	0,01 %	containing it shall be 'Ice Structuring Protein'		
Aqueous extracts of dried leaves of leaves	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
guayusa	Herbal infusions	In line with normal use in herbal infusions and food supplements of	containing it shall be 'Extracts of		
	Food Supplements as defined in Directive 2002/46/EC	a similar aqueous extract of dried leaves of <i>Ilex paraguariensis</i>			

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
M47						
	Infusion from coffee leaves of <i>Coffea</i>	Specified food category	Maximum levels	The designation of the novel food		
	arabica L. and/or Coffea canephora Pierre ex A. Froehner (Traditional food from a third country)	Herbal infusions		on the labelling of the foodstuffs containing it shall be 'Infusion from coffee leaves of Coffea arabica and/or Coffea canephora'.		
<u> 19</u>						
	Isomalto-oligos-	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Isom-		
	accharide	Energy-Reduced Soft Drinks	6,5 %			
		Energy Drinks	5,0 %	altooligosaccharide'.		
		Foods intended to meet the expenditure of intense muscular efforts, especially for sportsmen (including isotonic drinks)	6,5 %	2. Foods containing the novel ingredient must be labelled as 'a source of glucose'.		
		Fruit Juices	5 %			
		Processed Vegetables and Vegetable Juices	5 %			
		Other Soft Drinks	5 %			
		Cereals Bars	10 %			
		Cookies, Biscuits	20 %			
		Breakfast Cereal Bars	25 %			
		Hard Candies	97 %			
		Soft Candies/Chocolate Bars	25 %			
		Meal replacement for weight control (as bars or milk based)	20 %			
	Isomaltulose	Not specified		1. The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Isomaltulose'.		

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	►M29 Data Protection ◀
				2. The designation of the novel food on the labelling shall be accompanied by indication that the 'Isomaltulose is a source of glucose and fructose'.		
▼ <u>M14</u>						
	Lactitol	Specified food category	Maximum levels	The designation of the novel food		
		Food Supplements as defined in Directive 2002/46/EC (capsules, tablets or powder) intended for the adult population	20 g/day	on the labelling of the food supplements containing it shall be 'Lactitol'		
7 <u>M9</u>						
	Lacto-N-neotetraose	Specified food category	Maximum levels	1. The designation of the novel food		
		Unflavoured pasteurised and sterilised (including UHT) milk-based products	0,6 g/l	on the labelling of the foodstuffs containing it shall be 'lacto- <i>N</i> -neotetraose'.		
		Unflavoured fermented milk-based products	0,6 g/l for beverages	2. The labelling of food		
			9,6 g/kg for products other than beverages	supplements containing lacto- <i>N</i> - neotetraose shall bear a statement that the supplements		
		Flavoured fermented milk-based products	0,6 g/l for beverages	should not be used if other		
		including heat-treated products	9,6 g/kg for products other than beverages	foods with added lacto- <i>N</i> -neotetraose are consumed the same day.		
		Dairy analogues, including beverage	0,6 g/l for beverages	3. The labelling of food		
		whiteners	6 g/kg for products other than beverages	supplements containing lacto-N- neotetraose intended for young children shall bear a statement		
			200 g/kg for whitener	that the supplements should not be used if breast milk or other foods with added lacto-N-		
		Cereal bars	6 g/kg			
		Table-top sweeteners	100 g/kg	neotetraose are consumed the same day.		
		Infant formula as defined in Regulation (EU) No 609/2013	0,6 g/l in combination with up to 1,2 g/l of 2'-fucosyllactose at a ratio of 1:2 in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			

Follow-on formula as defined in Regulation (EU) No 609/2013 in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer Processed cereal-based food and haby food for infants and young children as defined in Regulation (EU) No 609/2013 intended for young children as defined in intended for young children as defined in the final products as a such or reconstituted as instructed by the manufacturer Mill-based drinks and similar products intended for young children as a such or reconstituted as instructed by the manufacturer Mill-based drinks and similar products intended for young children as a such or reconstituted as instructed by the manufacturer. Foods for special medical purposes as defined in Regulation (EU) No 609/2013 In accordance with the particular nutritional requirements of the persons for whom the products are intended. Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 Bread and pasta products bearing statements on the absence or reduced presence of glutten in accordance with the requirements of Commission Implementing Regulation (EU) Flavoured drinks O, 6 gl for iliquid food ready for use, marketed as such or reconstituted as instructed by the manufacturer. In accordance with the particular nutritional requirements of the persons for whom the products are intended. Jog 2 kg for bars 30 g/kg Flavoured drinks	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
for infants and young children as defined in Regulation (EU) No 609/2013 Milk-based drinks and similar products intended for young children Milk-based drinks and similar products intended for young children Milk-based drinks and similar products intended for young children Milk-based drinks and similar products intended for young children Milk-based drinks and similar products art artio of 1:2 in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer Foods for special medical purposes as defined in Regulation (EU) No 609/2013 In accordance with the particular nutritional requirements of the persons for whom the products are intended Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 Bread and pasta products bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014			1,2 g/l of 2'-fucosyllactose at a ratio of 1:2 in the final product ready for use, marketed as such or reconstituted as instructed by the			
intended for young children similar products added alone or in combination with 2'-fucosyllactose, at a ratio of 1:2 in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer Foods for special medical purposes as defined in Regulation (EU) No 609/2013 In accordance with the particular nutritional requirements of the persons for whom the products are intended Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 Bread and pasta products bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014		for infants and young children as defined in	beverages 0,6 g/l for liquid food ready for use, marketed as such or reconstituted as			
in Regulation (EU) No 609/2013 Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 Bread and pasta products bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014 nutritional requirements of the persons for whom the products are intended 2,4 g/l for drinks 20 g/kg for bars 30 g/kg			similar products added alone or in combination with 2'-fucosyllactose, at a ratio of 1:2 in the final product ready for use, marketed as such or reconstituted as instructed			
defined in Regulation (EU) No 609/2013 20 g/kg for bars Bread and pasta products bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014 30 g/kg			nutritional requirements of the persons for whom the products are			
on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014						
Flavoured drinks 0,6 g/l		on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU)	30 g/kg			
		Flavoured drinks	0,6 g/l			

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Coffee, tea (excluding black tea), herbal and fruit infusions, chicory; tea, herbal and fruit infusions and chicory extracts; tea, plant, fruit and cereal preparations for infusions, as well as mixes and instant mixes of these products	4,8 g/l — the maximum level refers to the products ready to use			
		Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants	1,5 g/day for general population 0,6 g/day for young children			
▼ <u>M43</u>						
	Lacto-N-tetraose ('LNT')	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		Authorised on 23.4.2020. This inclusion is based on
	(microbial source)	Unflavoured pasteurised and unflavoured sterilised (including UHT) milk products	1,0 g/l	tetraose are consumed the same day.		proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation
		Unflavoured fermented milk-based products	1,0 g/l (beverages) 10 g/kg (products other than beverages)			(EU) 2015/2283. Applicant: Glycom A/S, Kogle Allé 4, DK-2970 Hørsholm, Denmark. During the period of data protection,
		Flavoured fermented milk-based products including heat-treated products	1,0 g/l (beverages) 10 g/kg (products other than beverages)			the novel food lacto- <i>N</i> -tetraose is authorised for placing on the market within the Union only by Glycom A/S, unless a subsequent applicant obtains
		Beverages (flavoured drinks)	1,0 g/l			authorisation for the novel food without reference to the proprietary scientific
		Cereal bars	10 g/kg			evidence or scientific data protected in accordance with Article 26 of Regulation
		Infant formula as defined under Regulation (EU) No 609/2013	0,8 g/l in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			(EU) 2015/2283 or with the agreement of Glycom A/S.

▼<u>M43</u>

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Follow-on formula as defined under Regulation (EU) No 609/2013	0,6 g/l in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			End date of the data protection: 23.4.2025.
	Processed cereal-based food, baby food for infants and young children as defined under Regulation (EU) No 609/2013	0,6 g/l (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer 5 g/kg for products other than beverages			
	Milk based drinks and similar products intended for young children	0,6 g/l (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer 5 g/kg for products other than beverages			
	Total diet replacement foods for weight control as defined under Regulation (EU) No 609/2013	2,0 g/l (beverages) 20 g/kg (products other than beverages)			
	Food for special medical purposes as defined under Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Food Supplements as defined in Directive 2002/46/EC, excluding infants	2,0 g/day for young children, children, adolescents, and adults			

V 1V17						
	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
▼ <u>M20</u>	Lonicera caerulea L. berries (haskap) (Traditional food from a third country)	Not specified		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'haskap (Lonicera caerulea) berries'		
▼ <u>M9</u>	Lucerne leaf extract from Medicago sativa	Specified food category Food supplements as defined in Directive 2002/46/EC	Maximum levels 10 g/day	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Lucerne (Medicago sativa) protein' or 'Alfalfa (Medicago sativa) protein'.		
	Lycopene	Specified food category Fruit/vegetable juice-based drinks (including concentrates)	Maximum levels 2,5 mg/100 g	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Lycopene'		
		Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen	2,5 mg/100 g			
		Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	8 mg/meal			
		Breakfast cereals	5 mg/100 g			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Fats and dressings	10 mg/100 g			
	Soups other than tomato soups	1 mg/100 g			
	Bread (including crispy breads)	3 mg/100 g			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Food supplements as defined in Directive 2002/46/EC	15 mg/day			
Lycopene from Blakeslea trispora	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Lycopene'		
biakesieu irispora	Fruit/vegetable juice-based drinks (including concentrates)	2,5 mg/100 g			
	Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen	2,5 mg/100 g			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	8 mg/meal			
	Breakfast cereals	5 mg/100 g			
	Fats and dressings	10 mg/100 g			
	Soups other than tomato soups	1 mg/100 g			
	Bread (including crispy breads)	3 mg/100 g			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Food supplements as defined in Directive 2002/46/EC	15 mg/day			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Lycopene from tomatoes	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Fruit/vegetable juice-based drinks (including concentrates)	2,5 mg/100 g	containing it shall be 'Lycopene'		
	Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen	2,5 mg/100 g			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	8 mg/meal			
	Breakfast cereals	5 mg/100 g			
	Fats and dressings	10 mg/100 g			
S	Soups other than tomato soups	1 mg/100 g			
	Bread (including crispy breads)	3 mg/100 g			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Food supplements as defined in Directive 2002/46/EC	15 mg/day			
Lycopene oleoresin from tomatoes	Specified food category	Maximum levels of lycopene	The designation of the novel food on the labelling of the foodstuffs		_
	Fruit/vegetable juice-based drinks (including concentrates)	2,5 mg/100 g	containing it shall be 'Lycopene oleoresin from tomatoes'		
i	Drinks intended to meet the expenditure of intense muscular effort especially for sportsmen	2,5 mg/100 g			

Authorised novel fo	Od Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Total diet replacement for weight control covered by Regulation (EU) No 609/2013 and meal replacements for weight control	8 mg/meal			
	Breakfast cereals	5 mg/100 g			
	Fats and dressings	10 mg/100 g			
	Soups other than tomato soups	1 mg/100 g			
	Bread (including crispy breads)	3 mg/100 g			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
50					
Hen egg white	Specified food category	Maximum levels	The designation of the novel food		
lysozyme hydrolysate	Food supplements as defined in Directive 2002/46/EC intended for adult population	1000 mg/day	on the labelling of food supplements containing it shall be 'Hen egg white lysozyme hydrolysate'.		
9					
– Magnesium citrate	Specified food category	Maximum levels	The designation of the novel food		
malate	Food Supplements as defined in Directive 2002/46/EC		on the labelling of the foodstuffs containing it shall be 'Magnesium citrate malate'		
Magnolia Bark	Specified food category	Maximum levels	The designation of the novel food		
Extract	Mints (confectionary products)	0,2 % for breath freshening	on the labelling of the foodstuffs containing it shall be 'Magnolia		
	Chewing gum	purposes. Based on a 0,2 % maximum incorporation level and a maximum gum/mint size of 1,5 g each, each gum or mint serving will contain no more than 3 mg of magnolia bark extract.	Bark Extract'		
Maize-germ oil hig	sh Specified food category	Maximum levels	The designation of the novel food		
in unsaponifiable matter	Food Supplements as defined in Directive 2002/46/EC	2 g/day	on the labelling of the foodstuffs containing it shall be 'Maize-germ oil extract'		
	Chewing gum	2 %			

_	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	►M29 Data Protection ◀
M	I ethylcellulose	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs	Methylcellulose is not to be	
		Edible ices	2 %	containing it shall be 'Methylcel- lulose'	used in foods specially prepared for	
		Flavoured drinks			young children	
		Flavoured or unflavoured fermented milk products				
		Cold desserts (dairy, fat, fruit, cereal, egg-based products)				
		Fruit preparations (pulps, purees or compotes)				
_		Soups and broths				
<u>M11</u>						
	-Methylnicoti- amide chloride	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		Authorised on 2 Septembe 2018. This inclusion is based
		Food Supplements as defined in Directive 2002/46/EC for the adult population excluding pregnant and lactating women	58 mg/day	containing it shall be '1- Methylnicotinamide chloride'. Food supplements containing 1- Methylnicotinamide shall bear the following statement: This food supplement should be consumed by adults only excluding pregnant and lactating women		on proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation (EU 2015/2283. Applicant: Pharmena SA Wolczanska 178, 90 530 Lodz, Poland. During the period of data protection the

▼<u>M11</u>

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
					novel food 1-methylnicotinamide chloride is authorised for placing on the market within the Union only by Pharmena S.A. unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of Pharmena S.A. End date of the data
					protection: 2 September 2023
(6S)-5-methyltet- rahydrofolic acid, glucosamine salt	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be '(6S)-5-		
			methyltetrahydrofolic acid, gluco- samine salt' or '5MTHF-gluco- samine'		
	Food Supplements as defined in Directive 2002/46/EC as a source of folate				
Monomethylsil- anetriol (Organic	Specified food category	Maximum levels of silicon	The designation of the novel food on the labelling of the food supplements containing it shall be 'Organic silicon (monomethylsilanetriol)'		
Silicon)	Food Supplements as defined in Directive 2002/46/EC for adult population (in liquid form)	10,40 mg/day			

V 1V12						
	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Mycelial extract from Shiitake mushroom (Lentinula edodes)	Specified food category	Maximum levels	The designation of the novel food		
		Bread products	2 ml/100 g	on the labelling of the foodstuffs containing it shall be 'extract from		
		Soft drinks	0,5 ml/100 ml	the mushroom Lentinula edodes' or		
		Ready prepared meals	2,5 ml per meal	'extract from Shiitake mushroom'		
		Foods based on yoghurt	1,5 ml/100 ml			
		Food supplements as defined in Directive 2002/46/EC	2,5 ml per day dose			
▼ <u>M38</u>	Nicotinamide riboside chloride	Specified food category	Maximum levels	mihagida ahlamida		Authorised on 20 February 2020. This inclusion is
	riboside chioride	Food Supplements as defined in Directive 2002/46/EC	300 mg/day for the general adult population, excluding pregnant and lactating women 230 mg/day for pregnant and lactating women		containing it shall be 'Nicotinamide riboside chloride'	
						Applicant: ChromaDex Inc., 10900 Wilshire Boulevard Suite 600, Los Angeles, CA 90024 USA. During the period of data protection, the novel food is authorised for placing on the market within the Union only by ChromaDex Inc. unless a subsequent applicant obtains authorisation for that novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of ChromaDex Inc.
						End date of the data protection: 20 February 2025.

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Noni fruit juice (Morinda citrifolia)	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Pasteurised fruit and fruit nectar based drinks	30 ml with one serving (up to 100 % noni juice) or 20 ml twice a day, not more than 40 ml per day	containing it shall be 'Noni juice' or 'Juice of Morinda citrifolia'		
Noni fruit juice powder (Morinda citrifolia)	Food supplements as defined in Directive 2002/46/EC	6,6 g/day (equivalent to 30 ml of noni juice)	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Noni juice powder' or 'Juice powder of Morinda citrifolia'		
Noni fruit puree and concentrate	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be: For fruit puree: 'Morinda citrifolia fruit puree' or 'Noni fruit puree'		
(Morinda citrifolia)		Fruit puree			
	Candy/confectionery	45 g/100 g			
	Cereal bars	53 g/100 g	For fruit concentrate: 'Movinda citrifolia fruit concentrate'		
	Powdered nutritional drink mixes (dry weight)	53 g/100 g	'Morinda citrifolia fruit concentrate' or 'Noni fruit concentrate'		
	Carbonated beverages	11 g/100 g			
	Ice cream & sorbet	31 g/100 g			
	Yoghurt	12 g/100 g			
	Biscuits	53 g/100 g			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Buns, cakes and pastries	53 g/100 g			
	Breakfast cereals (wholegrain)	88 g/100 g			
	Jams and jellies in accordance with Directive 2001/113/EC	133 g/100 g			
	2001/113/EC	Based on pre-processing quantity to produce final 100 g product			
	Sweet spreads, fillings and icings	31 g/100 g			
	Savoury sauces, pickles, gravies and condiments	88 g/100 g			
	Food Supplements as defined in Directive 2002/46/EC	26 g/day			
		Fruit concentrate			
	Candy/Confectionery	10 g/100 g			
	Cereal bars	12 g/100 g			
	Powdered nutritional drink mixes (dry weight)	12 g/100 g			
	Carbonated beverages	3 g/100 g			
	Ice cream & sorbet	7 g/100 g			
	Yoghurt	3 g/100 g			
	Biscuits	12 g/100 g			
	Buns, cakes and pastries	12 g/100 g			
	Breakfast cereals (wholegrain)	20 g/100 g			
	Jams and jellies in accordance with Directive 2001/113/EC	30 g/100 g			
	Sweet spreads, fillings and icings	7 g/100 g			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Savoury sauces, pickles, gravies and condiments	20 g/100 g			
	Food Supplements as defined in Directive 2002/46/EC	6 g/day			
Noni leaves (Morinda citrifolia)	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
(caractural caragonal)	For the preparation of infusions	A cup of infusion to be consumed shall not be prepared with more than 1 g of dried and roasted leaves of Morinda citrifolia	containing it shall be 'Noni leaves' or 'leaves of <i>Morinda citrifolia</i> '. 2. Instructions shall be given to the consumer that a cup of infusion should not be prepared with more than 1 g of dried and roasted leaves of Morinda citrifolia.		
Noni fruit powder (<i>Morinda citrifolia</i>)	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Morinda citrifolia fruit powder' or 'Noni fruit powder'		
(ezorman caryona)	Food Supplements as defined in Directive 2002/46/EC	2,4 g per/day			
Odontella aurita microalgae	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
micioaigae	Flavoured pasta	1,5 %	containing it shall be 'Odontella aurita microalgae'		
	Fish soups	1 %			
	Marine terrines	0,5 %			
	Broth preparations	1 %			
	Crackers	1,5 %			
	Frozen breaded fish	1,5 %			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Oil enriched with phytosterols/phytostanols	Specified food category	Maximum levels of phytosterols/ phytostanols	In accordance with Annex III.5 to Regulation (EU) No 1169/2011		
	Spreadable fats as defined in Annex VII, Part VII and Appendix II, points B and C of Regulation (EU) No 1308/2013, and excluding cooking and frying fats and spreads based on butter or other animal fat	novel food ingredient shall be presented in such a manner that they can be easily divided into portions that contain either a maximum of 3 g (in case of one portion per day) or a maximum			
	Milk based products, such as products based on semi-skimmed and skimmed milk products, possibly with the addition of fruits and/or cereals, products based on fermented milk such as yoghurt and cheese based products (fat content ≤ 12 g per 100 g), where possibly the milk fat has been reduced and the fat or protein has been partly or fully replaced by vegetable fat or protein	of 1 g (in case of three portions per day) of added phytosterols/ phytostanols. 2. The amount of phytosterols/ phytostanols added to a container of beverages shall not exceed 3 g. 3. Salad dressings, mayonnaise and spicy sauces shall be packed as single portions.			
	Soya drinks				
	Salad dressings, mayonnaise and spicy sauces				

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Oil extracted from squids	Specified food category	Maximum levels of DHA and EPA combined	The designation of the novel food on the labelling of the foodstuffs		
		Dairy products except milk-based beverages	200 mg/100 g or for cheese products 600 mg/100 g	containing it shall be 'Squid oil'.		
		Dairy analogues except drinks	200 mg/100 g or for analogues to cheese products 600 mg/			
		Spreadable fat and dressings	600 mg/100 g			
		Breakfast cereals	500 mg/100 g			
		Bakery products (breads and bread rolls)	200 mg/100 g			
		Cereal bars	500 mg/100 g			
		Non-alcoholic beverages (including milk-based beverages)	60 mg/100 ml			
		Food Supplements as defined in Directive 2002/46/EC	3 000 mg/day for general population 450 mg/day for pregnant and lactating women			
		Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products intended			
		Total diet replacement for weight control defined in Regulation (EU) No 609/2013 and meal replacements for weight control	200 mg/meal			
▼ <u>M53</u>	Extract from <i>Panax</i>	Specified food category	Maximum levels	The designation of the novel food		Authorised on 23 December
	notoginseng and Astragalus membra- naceus	Food supplements as defined in Directive 2002/46/EC for the general adult population, excluding food supplements for pregnant women	35 mg/day	on the labelling of the foodstuffs containing it shall be 'Extract from Panax notoginseng and Astragalus membranaceus' The labelling of food supplements containing extract from Panax notoginseng and Astragalus membranaceus shall bear a statement that those food supplements should not be consumed by the population under 18 years of age and by pregnant women.		2020. This inclusion is based on proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283.

▼<u>M53</u>

	Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
						Applicant: NuLiv Science, 1050 W. Central Ave., Building C, Brea, CA 92821, USA.
						During the period of data protection, the novel food is authorised for placing on the market within the Union only by NuLiv Science, unless a subsequent applicant obtains authorisation for that novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of NuLiv Science.
						End date of the data protection: 23 December 2025.
▼ <u>M45</u>						
	Partially defatted chia seed (Salvia	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	hispanica) powders	Powder with high pro	otein content	containing it shall be 'Partially defatted chia seed (Salvia hispanica) powder'		
		Unflavoured fermented milk products, including natural unflavoured buttermilk (excluding sterilised buttermilk) non-heat-treated after fermentation	0,7 %	powder		
		Unflavoured fermented milk products, heat-treated after fermentation	0,7 %			
		Flavoured fermented milk products including heat-treated products	0,7 %			

▼<u>M45</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection •
-		Confectionery	10 %			
		Fruit juices as defined by Directive 2001/112/ EC (8) and vegetable juices	2,5 %			
		Fruit nectars as defined by Directive 2001/ 112/EC and vegetable nectars and similar products	2,5 %			
		Flavoured drinks	3 %			
		Food supplements as defined in Directive 2002/46/EC excluding food supplements for infants and young children	7,5 g/day			
		Powder with high fi	bre content			
		Confectionery	4 %			
		Fruit juices as defined by Directive 2001/112/ EC and vegetable juices	2,5 %			
		Fruit nectars as defined by Directive 2001/ 112/EC and vegetable nectars and similar products	4 %			
		Flavoured drinks	4 %			
		Food supplements as defined in Directive 2002/46/EC excluding food supplements for infants and young children	12 g/day			
<u>60</u>						
	Partially defatted	Specified food category	Maximum levels	The designation of the novel food		
	rapeseed powder from <i>Brassica rapa</i>	Cereal bars mixed	20 g/100 g	on the labelling of the foodstuffs containing it shall be 'Partially		
	L. and	Muesli and similar breakfast cereals	20 g/100 g	defatted Rapeseed powder'.		
i	Brassica napus L.	Extruded breakfast cereal products	20 g/100 g	Any foodstuff containing 'Partially		
		Snacks (excluding potato crisps)	15 g/100 g	defatted Rapeseed powder' from Brassica rapa L. and Brassica		
		Breads and rolls with added special ingredients (such as seeds, raisins, herbs)	7 g/100 g	napus L.' shall bear a statement that this ingredient may cause allergic reaction to consumers who		
		Brown breads bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014	7 g/100 g	are allergic to mustard and products thereof. That statement shall appear in close proximity to the list of ingredients.		

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Multigrain bread and rolls	7 g/100 g			
		Meat substitutes	10 g/100 g			
		Meat balls	10 g/100 g			
▼ <u>M9</u>	Pasteurised	Specified food category	Maximum levels	The wording 'pasteurised by		
	fruit-based prep- arations produced using high-pressure treatment	Types of fruit: apple, apricot, banana, blackberry, blueberry, cherry, coconut, fig, grape, grapefruit, mandarin, mango, melon, peach, pear, pineapple, prune, raspberry, rhubarb, strawberry		high-pressure treatment' shall be displayed next to the name of the fruit preparations as such and in any product in which it is used		
▼ <u>M35</u>						
	Phenylcapsaicin	Specified food category	Maximum levels	The designation of the novel food		Authorised on 19 December
		Foods for special medical purposes as defined under Regulation (EU) No 609/2013 excluding foods for infants, young children and children under the age of 11 years	2,5 mg/day	on the labelling of the foodstuffs containing it shall be 'phenylcapsaicin'.	containing it shall be 'phenylcap- saicin'.	2019. This inclusion is based on proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation (EU)
		Food supplements as defined in Directive 2002/46/EC intended for the general population, excluding children under the age of 11 years	2,5 mg/day			Applicant: aXichem AB, Södergatan 26, SE 211 34, Malmö Sweden. During the period of data protection, the novel food phenylcapsaicin is authorised for placing on the market within the Union only by aXichem AB, unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of aXichem AB.

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Phosphated maize starch	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Baked bakery products	15 %	containing it shall be 'Phosphated maize starch'		
	Pasta				
	Breakfast cereals				
	Cereal bars				
Phosphatidylserine from fish phosp- holipids	Specified food category	Maximum levels of phosphati- dylserine	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Fish phos-		
	Beverages based on yoghurt	50 mg/100 ml	phatidylserine'		
	Powders based on milk powders	3 500 mg/100 g (equivalent to 40 mg/100 ml ready to drink)			
	Foods based on yoghurt	80 mg/100 g			
	Cereal bars	350 mg/100 g			
	Chocolate based confectionary	200 mg/100 g			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In compliance with Regulation (EU) No 609/2013			
	Food supplements as defined in Directive 2002/46/EC	300 mg/day			

Authorised novel food	Conditions under which the no	Conditions under which the novel food may be used Addit		Other requirements	► <u>M29</u> Data Protection ◀
Phosphatidylserine from soya phosp- holipids	Specified food category	Maximum levels of phosphati- dylserine	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Soya phos-		
nonpius	Beverages based on yoghurt	50 mg/100 ml	phatidylserine'		
	Powders based on milk powder	3,5 g/100 g (equivalent to 40 mg/100 ml ready to drink)	to		
	Foods based on yoghurt	80 mg/100 g			
	Cereal bars	350 mg/100 g			
	Chocolate based confectionary	200 mg/100 g			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In compliance with Regulation (EU) No 609/2013			
Phospholipid product containing equal amounts of	Specified food category	Maximum levels of phosphati- dylserine	The designation of the novel food on the labelling of the foodstuffs containing shall be 'Soy phosphatidylserine and phosphatidic acid'	not intended to	
phosphatidylserine and phosphatidic	Breakfast cereals	80 mg/100 g			
acid	Cereal bars	350 mg/100 g			
	Foods based on yogurt	80 mg/100 g			
	Soy-based yogurt-like products	80 mg/100 g			
	Yogurt based-drinks	50 mg/100 g			
	Soy-based yogurt-like drinks	50 mg/100 g			
	Powders based on milk powder	3,5 g/100 g (equivalent to 40 mg/100 ml ready-to drink)			
	Food Supplements as defined in Directive 2002/46/EC	800 mg/day			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In compliance with Regulation (EU) No 609/2013			

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Phospholipides	Specified food category	Maximum levels			
from egg yolk	Not specified				
Phytoglycogen	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Phytog- lycogen'		
	Processed foods	25 %			
Phytosterols/	Specified food category	Maximum levels	In accordance with Annex III.5 of		
phytostanols	Rice drinks	1. They shall be presented in such a manner that they can be easily divided into portions that contain either a maximum of 3 g (in case of 1 portion/day) or a maximum of 1 g (in case of 3 portions/day) of added phytosterols/phytostanols. The amount of phytosterols/phytostanols added to a container of beverages shall not exceed 3 g. Salad dressings, mayonnaise and spicy sauces shall be packed as single portions	Regulation (EU) No 1169/2011		
	Rye bread with flour containing ≥ 50 % rye (wholemeal rye flour, whole or cracked rye kernels and rye flakes) and ≤ 30 % wheat; and with ≤ 4 % added sugar but no fat added.				
	Salad dressings, mayonnaise and spicy sauces.				
	Soya drink				
	Milk type products, such as semi-skimmed and skimmed milk type products, possibly with the addition of fruits and/or cereals, where possibly the milk fat has been reduced, or where milk fat and/or protein has been partly or fully replaced by vegetable fat and/or protein.				
	Products based on fermented milk such as yoghurt and cheese type products (fat content < 12 % per 100 g), where possibly the milk fat has been reduced, or where milk fat and/or protein has been partly or fully replaced by vegetable fat and/or protein				
	Spreadable fats as defined in Annex VII, Part VII and Appendix II, points B and C of Regulation (EU) No 1308/2013, and excluding cooking and frying fats and spreads based on butter or other animal fat.				
	Food Supplements as defined in Directive 2002/46/EC	3 g/day			

Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Specified food category	Maximum levels			
For frying and as seasoning	In line with normal food use of vegetable oils			
Not specified		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Potato protein'		
Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
Food Supplements as defined in Directive 2002/46/EC for general adult population	120 PPU/day (2,7 g of enzyme preparation/day) (2 × 10 ⁶ PPI/day)	containing it shall be 'Prolyl oligo- peptidase'		
	PPU – Prolyl Peptidase Units or Proline Protease Units			
	PPI – Protease Picomole International			
Specified food category	Maximum levels			
2002/46/EC e.e.	equalising 12,6 mg pig kidney extract a day Diamine oxidase			
Food for special medical purposes as defined in Regulation (EU) No 609/2013	capsules or 3 tablets with a content of DAO of 0,3 mg/capsule or 0,3 mg/tablet)			
	Specified food category For frying and as seasoning Not specified Specified food category Food Supplements as defined in Directive 2002/46/EC for general adult population Specified food category Food Supplements as defined in Directive 2002/46/EC	For frying and as seasoning In line with normal food use of vegetable oils Not specified Maximum levels	Specified food category Maximum levels	Specified food category Maximum levels

A	Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
<u>M10</u>						
Py qu sal	vrroloquinoline ninone disodium lt	Food Supplements as defined in Directive 2002/46/EC intended for the adult population, excluding pregnant and lactating women	Maximum levels 20 mg/day	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Pyrroloquinoline quinone disodium salt'. Food supplements containing Pyrroloquinoline quinone disodium salt shall bear the following statement: This food supplement should be consumed by adults only excluding pregnant and lactating women		Authorised on 2 Septemb 2018. This inclusion is based on proprietary scientific evidence are scientific data protected accordance with Article 2 of Regulation (EU) 2012283. Applicant: Mitsubishi Granding Company, Inc. Mitsubishi Building 5. Marunouchi 2-chom Chiyoda-ku, Tokyo 1008324, Japan. During the period of data protection the novel food Pyrroloquinolin quinone disodium salt is authorised for placing on the market within the Union on by Mitsubishi Gas Chemic Company, Inc., unless subsequent applicant obtain authorisation for the novel food without reference to the proprietary scientific evidence or scientific data protected accordance with Article 26 Regulation (EU) 2015/228 or with the agreement Mitsubishi Gas Chemic Company, Inc. End date of the da protection: 2 september 202

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Rapeseed oil high in unsaponifiable	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	matter	Food Supplements as defined in Directive 2002/46/EC	1,5 g per portion recommended for daily consumption	containing it shall be 'Rapeseed oil extract'		
	Rapeseed Protein	As a vegetable protein source in foods except in infant formula and follow-on formula		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Rapeseed protein'.		
				2. Any foodstuff containing 'rapeseed protein' shall bear a statement that this ingredient may cause allergic reaction to consumers who are allergic to mustard and products thereof. Where relevant, this statement shall appear in close proximity to the list of ingredients.		
▼ <u>M17</u>	Refined shrimp	Specified food category	Maximum levels	The designation of the novel food		Authorised on 20 November
	peptide concentrate	Food Supplements as defined in Directive 2002/46/EC for the adult population	1 200 mg/day	on the labelling of the foodstuffs containing it shall be 'refined shrimp peptide concentrate'.		2018. This inclusion is based on proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283.
						Applicant: Marealis AS., Stortorget 1, Kystens Hus, 2nd floor, N-9008 Tromsø Postal address: P.O. Box 1065, 9261 Tromsø, Norway. During the period of data protection the novel food refined shrimp peptide

▼<u>M17</u>

Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
				concentrate is authorised find placing on the market with the Union only by Marea AS unless a subseque applicant obtains authorization for the novel for without reference to the proprietary scientific day rotected in accordance with Article 26 of Regulation (El 2015/2283) or with the agreement of Marealis A End date of the day protection: 20 November 2023.
Specified food category	Maximum levels	1. The designation of the novel food		
Food supplements as defined in Directive 2002/46/EC for the adult population	150 mg/day	on the labelling of the food supplements containing it shall be ' <i>Trans</i> -resveratrol'. 2. The labelling of food supplements containing <i>trans</i> -resveratrol shall bear a statement that people using medicines should only consume the product under medical supervision.		
-	Specified food category Food supplements as defined in Directive	Food supplements as defined in Directive 150 mg/day	Specified food category Maximum levels 1. The designation of the novel food on the labelling of the food supplements containing it shall be 'Trans-resveratrol'. 150 mg/day 2. The labelling of food supplements containing trans-resveratrol shall bear a statement that people using medicines should only consume the product under medical	Specified food category Maximum levels 1. The designation of the novel food on the labelling of the food supplements containing it shall be 'Trans-resveratrol'. 2. The labelling of food supplements containing trans-resveratrol shall bear a statement that people using medicines should only consume the product under medical

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► M29 Data Protection ◀
Trans-resveratrol	Specified food category	Maximum levels	1. The designation of the novel food		
(microbial source)	Food supplements as defined in Directive 2002/46/EC	In line with normal use in food supplements of resveratrol extracted from Japanese knotweed (Fallopia japonica)	on the labelling of the food supplements containing it shall be ' <i>Trans</i> -resveratrol'.		
			2. The labelling of food supplements containing trans-resveratrol shall bear a statement that people using medicines should only consume the product under medical supervision.		
Rooster comb	Specified food category	Maximum levels	on the labelling of the foodstuffs on the labelling of the foodstuffs containing it shall be 'Rooster		
extract	Milk-based drinks	40 mg/100 g or mg/100 ml			
	Milk based fermented drinks	80 mg/100 g or mg/100 ml	comb extract' or 'Cockerel comb extract'		
	Yoghurt-type products	65 mg/100 g or mg/100 ml			
	Fromage frais	110 mg/100 g or mg/100 ml			
Sacha inchi oil from	Specified food category	Maximum levels	The designation of the novel food		
Plukenetia volubilis	As for linseed oil	In line with normal food use of linseed oil	on the labelling of the foodstuffs containing it shall be 'Sacha inchi oil (Plukenetia volubilis)'		
Salatrims	Specified food category	Maximum levels	1. The designation of the novel food		
	Bakery products and confectionary		on the labelling of the foodstuffs containing it shall be 'reduced energy fat (salatrims)'.		
			2. There shall be a statement that excessive consumption may lead to gastro-intestinal disturbance.		
			3. There shall be a statement that the products are not intended for use by children.		

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Schizochytrium sp. oil rich in DHA and EPA	Specified food category	Maximum levels of DHA and EPA combined:	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'DHA and		
	Food Supplements as defined in Directive 2002/46/EC for adult population excluding pregnant and lactating women	3 000 mg/day	EPA-rich oil from the microalgae Schizochytrium sp.'		
	Food Supplements as defined in Directive 2002/46/EC for pregnant and lactating women	450 mg/day			
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			
	Milk-based drinks and similar products intended for young children	200 mg/100 g			
	Processed cereal based food and baby food for infants and young children as defined in Regulation (EU) No 609/2013	-			
	Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen				
	Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014				
	Bakery Products (Breads, Rolls and Sweet Biscuits)	200 mg/100 g			
	Breakfast Cereals	500 mg/100 g			
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	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Cooking Fats	360 mg/100 g			
		Dairy Analogues except drinks	600 mg/100 g for cheese; 200 mg/ 100 g for soy and imitation milk products (excluding drinks)			
		Dairy Products except milk-based drinks	600 mg/100 g for cheese; 200 mg/ 100 g for milk products (including milk, fromage frais and yoghurt products; excluding drinks)			
		Non-alcoholic Beverages (including dairy analogue and milk-based drinks)	80 mg/100 g			
		Cereal/Nutrition Bars	500 mg/100 g			
		Spreadable Fats and Dressings	600 mg/100 g			
▼ <u>M26</u>						
	Schizochytrium sp. (ATCC PTA-9695)	Specified food category	Maximum levels of DHA	The designation of the novel food on the labelling of the foodstuffs		
	oil	Dairy products except milk-based drinks	200 mg/100 g or for cheese products $600 mg/100 g$	containing it shall be 'Oil from the microalgae <i>Schizochytrium</i> sp.'		
		Dairy analogues except drinks	200 mg/100 g or for analogues to cheese products 600 mg/100 g			
		Spreadable fats and dressings	600 mg/100 g			
		Breakfast cereals	500 mg/100 g			
		Food Supplements as defined in Directive 2002/46/EC	250 mg DHA/day for general population			
			450 mg DHA/day for pregnant and lactating women			
		Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			

▼<u>M26</u>

Authorised novel foo	od Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► M29 Data Protection ◀
	Milk-based drinks and similar products intended for young children	200 mg/100 g			
	Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen				
	Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014				
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutri- tional requirements of the persons for whom the products are intended			
	Bakery products (breads, rolls, and sweet biscuits)	200 mg/100 g			
	Cereal bars	500 mg/100 g			
	Cooking fats	360 mg/100 g			
	Non-alcoholic beverages (including dairy analogue and milk-based drinks)	80 mg/100 ml			
	Infant formula and follow-on formula as defined in Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013			
	Processed cereal-based foods and baby foods for infants and young children as defined in Regulation (EU) No 609/2013	200 mg/100 g			
	Fruit/vegetable puree	100 mg/100 g			
Schizochytrium sp.	Specified food category	Maximum levels of DHA	The designation of the novel food		
(FCC-3204) oil	Infant formula and follow-on formula as defined in Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013	on the labelling of the foodstuffs containing it shall be 'Oil from the microalgae <i>Schizochytrium</i> sp.'.		
	Food supplements as defined in Directive 2002/46/EC for the general population above 3 years of age	1 g/day	The labelling of food supplements containing <i>Schizochytrium</i> sp. (FCC-3204) oil shall bear a statement that they should not be consumed by infants and children under 3 years of age.		

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
▼ <u>M24</u>						
	Schizochytrium sp.	Specified food category	Maximum levels of DHA	The designation of the novel food on the labelling of the foodstuffs		
		Dairy products except milk-based drinks	200 mg/100 g or for cheese products 600 mg/100 g	containing it shall be 'Oil from the microalgae <i>Schizochytrium</i> sp.'		
		Dairy analogues except drinks	200 mg/100 g or for analogues to cheese products 600 mg/100 g			
		Spreadable fat and dressings	600 mg/100 g			
		Breakfast cereals	500 mg/100 g			
		Food Supplements as defined in Directive 2002/46/EC	250 mg DHA/day for general population			
			450 mg DHA/day for pregnant and lactating women			
		Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			
		Milk-based drinks and similar products intended for young children	200 mg/100 g			
		Processed cereal-based foods and baby foods for infants and young children as defined in Regulation (EU) No 609/2013				
		Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen				
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▼<u>M24</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Implementing Regulation (EU) No 828/2014				
		Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
		Bakery products (breads, rolls, and, sweet biscuits)	200 mg/100 g			
		Cereal bars	500 mg/100 g			
		Cooking fats	360 mg/100 g			
		Non-alcoholic beverages (including dairy analogue and milk-based drinks)	80 mg/100 ml			
		Fruit/vegetable puree	100 mg/100 g			
▼ <u>M50</u>						
	Schizochytrium sp. (T18) oil	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	()	Dairy products except milk-based drinks	200 mg/100 g or for cheese products 600 mg/100 g	containing it shall be 'Oil from the microalgae <i>Schizochytrium</i> sp.'.		
		Dairy analogues except drinks	200 mg/100 g or for analogues to cheese products 600 mg/100 g			
		Spreadable fats and dressings	600 mg/100 g			
		Breakfast cereals	500 mg/100 g			

▼<u>M50</u>

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Food supplements as defined in Directive 2002/46/EC	250 mg DHA/day for general population			
		450 mg DHA/day for pregnant and lactating women			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 and meal replacements for weight control	250 mg/meal			
	Milk-based drinks and similar products intended for young children	200 mg/100 g			
	Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen				
	Foods bearing statements on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014				
	Foods for special medical purposes as defined in Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
	Bakery products (breads, rolls and, sweet biscuits)	200 mg/100 g			
	Cereal bars	500 mg/100g			

▼<u>M50</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Cooking fats	360 mg/100 g			
		Non-alcoholic beverages (including dairy analogue and milk-based drinks)	80 mg/100 ml			
		Infant formula and follow-on formula as defined in Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013			
		Processed cereal-based foods and baby foods for infants and young children as defined in Regulation (EU) No 609/2013	200 mg/100 g			
		Fruit/vegetable puree	100 mg/100 g			
▼ <u>M62</u>						
	Schizochytrium sp. (WZU477) oil	Specified food category	Maximum levels of DHA	The designation of the novel food on the labelling of the foodstuffs		Authorised on 16 May 2021. This inclusion is based on
	(WZ0477) UII	Infant formula and follow-on formula as defined in Regulation (EU) No 609/2013	In accordance with Regulation (EU) No 609/2013	containing it shall be 'Oil from the microalgae <i>Schizochytrium</i> sp.'		proprietary scientific evidence and scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283.
						Applicant: Progress Biotech bv, Canaalstaete, Kanaalweg 33, 2903LR Capelle aan den Ijssel, the Netherlands.
						During the period of data protection, the novel food is authorised for placing on the market within the Union only by Progress Biotech by unless a subsequent applicant obtains authorisation for that novel food without reference to the proprietary scientific evidence or scientific data

▼<u>M62</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
						protected in accordance with Article 26 of Regulation (EU 2015/2283 or with the agreement of Progress Biotech bv. End date of the data protection: 16 May 2026 (5 years).
7 <u>M55</u>						
	Selenium-containing yeast (Yarrowia lipolytica) biomass	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'selenium-containing yeast (Yarrowia lipolytica) biomass'.		
		Food supplements as defined in Directive 2002/46/EC (³), excluding food supplements for infants and children under 4 years of age	50 mg/day for children from 4 to 6 years of age, resulting in 10 μg of selenium per day 100 mg/day for children from 7 to 10 years of age, resulting in 20 μg of selenium per day 500 mg/day for adolescents from 11 to 17 years of age, resulting in 100 μg of selenium per day 800 mg/day for adults, resulting in	The labelling of food supplements containing selenium-containing yeast (<i>Yarrowia lipolytica</i>) biomass shall bear a statement that the food supplements should not be consumed by infants and children under 4 years of age/children under 7 years of age/children under 11 years of age/children and adolescents under 18 years of age (12).		
			800 mg/day for adults, resulting in 160 μg of selenium per day			

▼ M9

▼ <u>M9</u>						
	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
▼ <u>M59</u>	3'-Sialyllactose (3'-SL) sodium salt	Specified food category	Maximum levels (expressed as 3'- Sialyllactose)	The designation of the novel food on the labelling of the foodstuffs		Authorised on 18 February 2021. This inclusion is
	(microbial source)	Unflavoured pasteurised and unflavoured sterilised (including UHT) milk products	0,25 g/L	containing it shall be '3'-Sialyl-lactose sodium salt'.		based on proprietary scientific evidence and scientific data protected in
		Flavoured fermented milk-based products including heat-treated products	0,25 g/L (beverages) 0,5 g/kg (products other than beverages)	than than than The labelling of food supplements containing 3'-Sialyllactose sodium salt shall bear a statement that they should not be consumed: a) if foods containing added 3'-Sialyllactose sodium salt are consumed the same day. b) by infants and young children ady for reconmanu- ready reconmanu-	um ney 3'-	accordance with Article 26 of Regulation (EU) 2015, 2283.
		Unflavoured fermented milk-based products	0,25 g/L (beverages) 2,5 g/kg (products other than beverages)			Applicant: Glycom A/S Kogle Allé 4, DK-2970 Hørsholm, Denmark. During the period of data protection
		Beverages (flavoured drinks, excluding drinks with a pH less than 5)	0,25 g/L		the novel food 3'-sialyl- lactose sodium salt is auth- orised for placing on the	
		Cereal bars	2,5 g/kg			market within the Unior
		Infant formula as defined under Regulation (EU) No 609/2013	0,2 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer		only by Glycom A/S, unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283 or with the agreement of Glycom A/S.	
		Follow-on formula as defined under Regulation (EU) No 609/2013	0,15 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			
		Processed cereal-based food and baby food for infants and young children as defined under Regulation (EU) No 609/2013 0,15 g/L (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer		End date of the data protection: 18 February 2026.		
			1,25 g/kg for products other than beverages			
		Milk-based drinks and similar products intended for young children	0,15 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			

other

than

0,5 g/L (beverages)

5 g/kg (products beverages)

Total diet replacement foods for weight control as defined under Regulation (EU) No 609/2013

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Food for special medical purposes as defined under Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
		Food Supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children	0,5 g/day			
▼M58						
	6'-Sialyllactose (6'-SL) sodium salt	Specified food category	Maximum levels (expressed as 6'-Sialyllactose)	The designation of the novel food on the labelling of the foodstuffs	ibelling of the foodstuffs	Authorised on 17 February 2021. This inclusion i
	(microbial source)	Unflavoured pasteurised and unflavoured sterilised (including UHT) milk products	0,5 g/L	containing it shall be '6'-Sialyl-lactose sodium salt'.		based on proprietary scie ntific evidence and scientific data protected in accordance
		Unflavoured fermented milk-based products	0,5 g/L (beverages)	The labelling of food supplements containing 6'-Sialyllactose (6'-SL)	with Article 2 lation (EU) 201: Applicant: Gl. Kogle Allé 4 Hørsholm, Denr the period of da the novel foo	with Article 26 of Regulation (EU) 2015/2283. Applicant: Glycom A/S, Kogle Allé 4, DK-2970 Hørsholm, Denmark. During
			2,5 g/kg (products other than beverages)	sodium salt shall bear a statement that they should not be consumed:		
		Flavoured fermented milk-based products	0,5 g/L (beverages)	a) if foods containing added 6'-		
		including heat-treated products	5,0 g/kg (products other than beverages)	Sialyllactose sodium salt are consumed on the same day.		ti ti
		Beverages (flavoured drinks, excluding drinks with a pH less than 5)	0,5 g/L	b) by infants and young children		lactose sodium salt is authorised for placing on the market within the Union only by Glycom A/S, unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific evidence or scientific data
		Cereal bars	5,0 g/kg			
		Infant formula as defined under Regulation (EU) No 609/2013	0,4 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer		obtains authoris novel food with to the propriet evidence or so protected in acc Article 26 of Re 2015/2283 or agreement of G	
		Follow-on formula as defined under Regulation (EU) No 609/2013	0,3 g/L in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			protected in accordance wit Article 26 of Regulation (EU 2015/2283 or with th agreement of Glycom A/S.
		Processed cereal-based food and baby food for infants and young children as defined under Regulation (EU) No 609/2013	0,3 g/L (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			End date of the data protection: 17 February 2026.
			2,5 g/kg for products other than beverages			

▼<u>M58</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Milk based drinks and similar products intended for young children	0,3 g/L (beverages) in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer			
		Total diet replacement foods for weight control as defined under Regulation (EU)	1,0 g/L (beverages)			
		No 609/2013	10,0 g/kg (products other than beverages)			
		Food for special medical purposes as defined under Regulation (EU) No 609/2013	In accordance with the particular nutritional requirements of the persons for whom the products are intended			
		Food Supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children	1,0 g/day			
▼ <u>M22</u>						
	Syrup from Sorghum bicolor (L.) Moench	Not specified		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Sorghum		
	(Traditional food from a third country)			(Sorghum bicolor) syrup'		
▼ <u>M9</u>						
	Fermented soybean extract	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	CAUACI	Food Supplements as defined in Directive 2002/46/EC (capsules, tablets or powder form) intended for the adult population, excluding pregnant and lactating women	100 mg/day	containing it shall be 'Fermented soybean extract'. 2. The labelling of food supplements containing fermented soybean extract shall bear a statement that persons taking medication should only consume the product under medical supervision.		

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► M29 Data Protection ◀
Spermidine-rich wheat germ extract	Specified food category	Maximum levels	The designation of the novel food		
(Triticum aestivum)	Food Supplements as defined in Directive 2002/46/EC intended for the adult population, excluding pregnant and lactating women	Equivalent of max. 6 mg/day spermidine	on the labelling of the food supplements containing it shall be 'spermidine-rich wheat germ extract'		
Sucromalt	Specified food category	Maximum levels	1. The designation of the novel food		
	Not specified		on the labelling of the foodstuffs containing it shall be 'Sucromalt'.		
			2. The designation of the novel food on the labelling shall be accompanied by indication that the product is a source of glucose and fructose.		
Sugar cane fibre	Specified food category	Maximum levels			
	Bread	8 %			
	Bakery goods	5 %			
	Meat and muscle products	3 %			
	Seasonings and spices	3 %			
	Grated cheeses	2 %			
	Special diet foods	5 %			
	Sauces	2 %			
	Beverages	5 %			
Sugars obtained from cocoa (<i>Theobroma cacao</i> L.) pulp	Not specified		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'sugars obtained from cocoa (<i>Theobroma cacao</i> L.) pulp', 'Glucose obtained from cocoa (<i>Theobroma cacao</i> L.) pulp' or 'Fructose obtained from cocoa (<i>Theobroma cacao</i> L.) pulp', depending on the form used.		

▼M9

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Sunflower oil extract	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Food Supplements as defined in Directive 2002/46/EC	1,1 g/day	containing it shall be 'Sunflower oil extract'		
70					
Synsepalum dulcificum dried fruits	Specified food category Food Supplements as defined in Directive 2002/46/EC for adult population excluding pregnant and lactating women	Maximum levels 0,7 g/day	1. The designation of the novel food on the labelling of food supplements containing it shall be 'dried Synsepalum dulcificum fruits' 2. The labelling of food supplements containing Synsepalum dulcificum dried fruits shall bear a statement that this food supplement should be consumed by adults only excluding pregnant and lactating women.		Authorised on 5 Decem 2021. This inclusion based on proprie scientific evidence scientific data protected accordance with Article of Regulation (EU) 202283. Applicant: Medicinal Gards.L. Marqués de Urquijo 1° D, Office 1, Mad 28008, Spain. During the period of oprotection, the novel food authorised for placing on market within the Union oby Medicinal Gardens sunless a subsequent applie obtains authorisation for novel food without refere to the proprietary scient evidence or scientific oprotected in accordance of Article 26 of Regulation (I 2015/2283) or with agreement of Medic Gardens S.L. End date of the data protect 5 December 2026.

<u>.</u>					
Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Dried <i>Tenebrio</i>	Specified food category	Maximum levels	The designation of the novel food		Authorised on 22 June 20
molitor larva (yellow mealworm)	Dried <i>Tenebrio molitor</i> larva, whole or in powder		on the labelling of the foodstuffs containing it shall be 'Dried Tenebrio molitor larva (yellow mealworm)'. 2. The labelling of the foodstuffs containing dried Tenebrio molitor larva (yellow mealworm) shall bear a statement that this ingredient may cause allergic reactions to consumers with known allergies to crustaceans and products thereof, and to dust mites. This statement shall appear in close proximity to the list of ingredients.		This inclusion is based proprietary scientific evide and scientific data protecte accordance with Article 2 Regulation (EU) 2015/228
	Protein products	10 g/100 g			Applicant: SAS EAP Group, 35 Boulevard du Libre Échange, 31650 Saint-Orens-
	Biscuits	10 g/100 g			de-Gameville, France. During the period of protection, the novel foo
	Legumes-based dishes	10 g/100 g			authorised for placing on market within the Union by SAS EAP Group, unle
	Pasta-based products	10 g/100 g			subsequent applicant ob authorisation for that r food without reference to proprietary scientific evid or scientific data protects accordance with Article 2 Regulation (EU) 2015/2 or with the agreement of EAP Group.
					End date of the data protect 22 June 2026.

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► M29 Data Protection ◀
Dried Tetraselmis chuii microalgae	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Sauces	20 % or 250mg/day	containing it shall be 'Dried microalgae <i>Tetraselmis chuii</i> ' or 'Dried microalgae <i>T. chuii</i> '		
	Special salts	1 %	Food supplements containing dried microalgae <i>Tetraselmis chuii</i> shall		
	Condiment	250 mg/day	bear the following statement: 'Contains negligible amounts of iodine'		
	Food Supplements as defined in Directive 2002/46/EC	250 mg/day			
Therapon barcoo/ Scortum	Intended use identical to that of the salmon, n products and dishes, including cooked, raw, s				
D-Tagatose	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
	Not specified		containing it shall be 'D- Tagatose'.		
			2. The labelling of any product where the level of D-Tagatose exceeds 15 g per serving and all beverages containing greater than 1 % D-Tagatose (as consumed) shall bear a statement 'excessive consumption may produce laxative effects'.		
. Taxifolin-rich	Specified food category	Maximum levels	The designation of the novel food		
extract	Yogurt plain/Yogurt with fruits ^(*)	0,020 g/kg	on the labelling of the foodstuffs containing it shall be 'taxifolin-rich		
	Kephir ^(*)	0,008 g/kg	extract'		
	Buttermilk ^(*)	0,005 g/kg	1		

▼<u>M50</u>

	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
		Milk powder ^(*)	0,052 g/kg			
		Cream ^(*)	0,070 g/kg			
		Sour cream ^(*)	0,050 g/kg			
		Cheese ^(*)	0,090 g/kg			
		Butter ^(*)	0,164 g/kg			
		Chocolate confectionery	0,070 g/kg			
		Non-alcoholic beverages	0,020 g/L			
		Food supplements as defined in Directive 2002/46/EC intended for the general population, excluding infants, young children, children and adolescents younger than 14 years	100 mg/day			
		(*) When used in milk products Taxifolin-rich part, any milk constituent	extract may not replace in whole or in			
▼ <u>M9</u>						
	Trehalose	Specified food category	Maximum levels	The designation of the novel food on the labelling of the foodstuffs		
		Not specified		containing it shall be 'Trehalose' and shall be displayed on the labelling of the product as such or in the list of ingredients of foodstuffs containing it. 2. The designation of the novel food on the labelling shall be accompanied by indication that the 'Trehalose is a source of glucose'.		

▼<u>M50</u>

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
UV-treated mushrooms (Agaricus bisporus)	Specified food category Mushrooms (Agaricus bisporus)	Maximum levels of vitamin D_2 20 µg of vitamin $D_2/100$ g fresh weight	 The designation on the label of the novel food as such or of the foodstuffs containing it shall be 'UV-treated mushrooms (Agaricus bisporus)'. The designation on the label of the novel food as such or of the foodstuffs containing it shall be accompanied by indication that a 'controlled light treatment was used to increase vitamin D levels' or 'UV treatment was used to increase vitamin D₂ levels'. 		
UV-treated baker's yeast (Saccharomyces cerevisiae)	Specified food category Yeast-leavened breads and rolls Yeast-leavened fine bakery wares Food supplements as defined in Directive 2002/46/EC	Maximum levels of vitamin D ₂ 5 μg of vitamin D ₂ /100 g 5 μg of vitamin D ₂ /100 g	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Vitamin D yeast' or 'Vitamin D ₂ yeast'		
	Pre-packed fresh or dry yeast for home baking	45 μg/100 g for fresh yeast 200 μg/100 g for dried yeast	 The designation of the novel food on the labelling of the foodstuffs shall be 'Vitamin D yeast' or 'Vitamin D₂ yeast'. The labelling of the novel food shall bear a statement that the foodstuff is only intended for baking and that it should not be eaten raw. The labelling of the novel food shall bear instructions for use for the final consumers so that a maximum concentration of 5 μg/100 g of vitamin D₂ in final home-baked products is not exceeded. 		

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
UV-treated bread	Specified food category	Maximum levels of vitamin D_2	The designation on the label of the novel food shall be accompanied by 'contains vitamin D produced by UV-treatment'		
	Yeast leavened bread and rolls (without toppings)	3 μg vitamin D ₂ /100 g			
UV-treated milk	Specified food category	Maximum levels of vitamin D_3	The designation on the label of the novel food shall be 'UV-treated'.		
	Pasteurised whole milk as defined in Regulation (EU) No 1308/2013 to be consumed as such		accordance with Point 2 of Part A of Annex XIII to Regulation (EU) No 1169/2011 of the European Parliament and of the Council, the designation for the labelling shall be accom-		
	Pasteurised semi-skimmed milk as defined in Regulation (EU) No 1308/2013 to be consumed as such	1-15 μg/kg for general population excluding infants			

7 <u>M49</u>	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	►M29 Data Protection ◀
▼ <u>M49</u>				, , ,	other requirements	Data Protection
	Vitamin D ₂ mushroom powder	Specified food category	Maximum levels of vitamin D_2 (11)	on the labelling of the foodstuffs containing it shall be 'UV-treated mushroom powder containing vitamin D' or 'UV-treated mushroom powder containing vitamin D ₂ ' The labelling of food supplements containing vitamin D ₂ mushroom powder shall bear a statement that they should not be consumed by infants		Authorised on 27 August 2020. This inclusion is based on proprietary
	mushroom powder	Breakfast cereals	2,25 μg of vitamin $D_2/100$ g			
		Yeast-leavened bread and pastries	2,25 μg of vitamin D ₂ /100 g			scientific evidence and scientific data protected in
		Grain products and pastas	2,25 μg of vitamin D ₂ /100 g			accordance with Article 26 of Regulation (EU) 2015/
		Fruit juice and fruit/vegetable blend beverages	1,125 μg of vitamin $D_2/100$ mL			2283. Applicant: Oakshire Naturals,
		Milk and dairy products (excluding fluid milks)	2,25 μg of vitamin $D_2/100$ g/1,125 μg of vitamin $D_2/100$ mL (beverages)		they should not be consumed by infants	LP., PO Box 388 Kennett Square, Pennsylvania 19348, United States. During the period of data protection, the novel food vitamin D ₂ mushroom powder is authorised for placing on the market within the Union only by Oakshire Naturals, LP., unless a subsequent applicant obtains authorisation for the novel food without reference to the proprietary scientific evidence or scientific data
		Cheese (excluding cottage cheese, ricotta cheese, and hard-grating cheeses)	2,25 μg of vitamin $D_2/100~g$		n o o n o o L a a a w w p p e e p A	
		Meal replacement bars and beverages	2,25 μg of vitamin $D_2/100$ g/1,125 μg of vitamin $D_2/100$ mL (beverages)			
		Dairy analogues	2,25 μg of vitamin $D_2/100$ g/1,125 μg of vitamin $D_2/100$ mL (beverages)			
		Meat analogues	2,25 μg of vitamin $D_2/100$ g			protected in accordance with Article 26 of Regulation
		Soups and broths	2,25 μg of vitamin D ₂ /100 g			(EU) 2015/2283 or with the agreement of Oakshire
		Extruded vegetable snacks	2,25 μg of vitamin D ₂ /100 g			Naturals, LP.
		Foods for Special Medical Purposes as defined under Regulation (EU) No 609/2013 excluding those intended for infants	15 μg/day			
		Food supplements as defined in Directive 2002/46/EC intended for the general population excluding infants	15 μg/day			

1117								
	Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀		
<u>M73</u>								
	Vitamin D ₂ mushroom powder	Specified food category Breakfast cereals	Maximum levels of vitamin D_2 2,1 µg/100 g	1. The designation of the novel food on the labelling of the foodstuffs containing it shall be 'UV-treated		Authorised on 19 Decemb 2021. This inclusion is base on proprietary scientif		
		Yeast leavened bread and similar pastries	2,1 μg/100 g	mushroom powder containing vitamin D ₂ '		evidence and scientific da protected in accordance wi		
		Grain products and pasta and similar products	2,1 μg/100 g	2. The labelling of food		Article 26 of Regulation (EU) 2015/2283.		
		Fruit/vegetable juices and nectars	1,1 µg/100 ml (marketed as such or reconstituted as instructed by the manufacturer)	supplements containing vitamin D_2 mushroom powder shall bear a statement that they should not be consumed by infants and		Applicant: MBio, Monagha Mushrooms, Tullygon Tyholland, Co. Monagha		
		Dairy products and analogues other than beverages	2,1 µg/100 g (marketed as such or reconstituted as instructed by the manufacturer)	children under 3 years of age.		Ireland. During the period data protection, the nov food vitamin D ₂ mushroo powder is authorised f		
		Dairy products and analogues as beverages	1,1 µg/100 ml (marketed as such or reconstituted as instructed by the manufacturer)			placing on the mark within the Union only MBio, Monagh Mushrooms, unless subsequent applicant obtai authorisation for the nov food without reference		
		Milk and dairy powders	21,3 μg/100 g (marketed as such or reconstituted as instructed by the manufacturer)		subsequent appli authorisation for food without r the proprietary evidence or sci protected in acco Article 26 of (EU) 2015/2283 agreement			
		Meat analogues	2,1 μg/100 g					evidence or scientific da protected in accordance wi
		Soups	2,1 µg/100 ml (marketed as such or reconstituted as instructed by the manufacturer)			Article 26 of Regulation (EU) 2015/2283 or with t		
		Extruded vegetable snack	2,1 μg/100 g				End date of the da	
		Meal replacement for weight control	2,1 μg/100 g		protection: 19			
		Foods for Special Medical Purposes as defined under Regulation (EU) No 609/2013 excluding those intended for infants	In accordance with the particular nutritional requirements of the persons for whom the products are intended			2020.		
		Food supplements as defined in Directive 2002/46/EC excluding food supplements intended for infants and young children	15 μg of vitamin D ₂ /day					

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Vitamin K ₂ (menaquinone)	To be used in compliance with Directive 2002 and/or Regulation (EC) No 1925/2006	/46/EC, Regulation (EU) No 609/2013	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Menaquinone' or 'Vitamin K ₂ '		
Wheat bran extract	Specified food category	Maximum levels	The designation of the novel food	The 'Wheat	
	Beer and substitutes	0,4 g/100 g	1 2	Bran Extract' may not be	
	Ready to eat cereals	9 g/100 g	extract'	introduced onto	
	Dairy products	2,4 g/100 g]	the market as a food supple-	
	Fruit and vegetable juices	0,6 g/100 g]	ment or food	
	Soft drinks	0,6 g/100 g]	supplement ingredient. Nor	
	Meat preparations	2 g/100 g		may it be added to infant formula.	
5					
Wolffia arrhiza and/	Specified food category	Maximum levels	The designation of the novel food		
or Wolffia globosa fresh plants (Tradi- tional food from a third country)	Wolffia arrhiza and/or Wolffia globosa fresh plants as such		on the labelling of the foodstuffs containing it shall be 'Wolffia arrhiza and Wolffia globosa' or 'Wolffia arrhiza' or 'Wolffia globosa' depending on the plant used.		
<u>6</u>					
Xylo-oligos- accharides	Specified food category	Maximum levels (10)	The designation of the novel food on the labelling of the foodstuffs		
accharacs	White bread	14 g/kg	containing it shall be 'Xylo-oligos-		
	Wholemeal bread	14 g/kg	accharides'		
	Breakfast cereals	14 g/kg	_		
	Biscuits	14 g/kg			
	Soy drink	3,5 g/kg	_		
	Yoghurt (9)	3,5 g/kg			
	Fruit spreads	30 g/kg]		
	Chocolate confectionery	30 g/kg]		
	Food supplements as defined in Directive 2002/46/EC for the general adult population	2 g/day			

Authorised novel food	Conditions under which the nov	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
Yarrowia lipolytica	Specified food category	Maximum levels	The designation of the novel food		
cast biolilass	Food Supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children	6 g/day for children from 10 years of age, adolescents and general adult population	containing it shall be 'Yarrowia lipolytica yeast heat-killed biomass'		
		3 g/day for children from 3 to 9 years of age			
Yeast beta-glucans	Specified food category	Maximum levels of pure beta-glucans from yeast (Sacchar- omyces cervisiae)	The designation of the novel food on the labelling of the foodstuffs containing it shall be 'Yeast		
	Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children	1,275 g/day for children older than 12 years and general adult population	glucans'		
		0,675 g/day for children younger than 12 years			
	Total diet replacement for weight control as defined in Regulation (EU) No 609/2013	1,275 g/day			
	Food for special medical purposes as defined in Regulation (EU) No 609/2013, excluding food for special medical purposes intended for infants and young children	1,275 g/day			
	Beverages based on fruit and/or vegetable juices including concentrate and dehydrated juices	1,3 g/kg			
	Fruit-flavoured drinks	0,8 g/kg			
	Cocoa beverages preparation powder	38,3 g/kg (powder)			
	Other beverages	0,8 g/kg (ready to drink)			
		7 g/kg (powder)			
	Cereal bars	6 g/kg			
	Breakfast cereals	15,3 g/kg			
	Varrowia lipolytica reast biomass	Food Supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 Food for special medical purposes as defined in Regulation (EU) No 609/2013, excluding food for special medical purposes intended for infants and young children Beverages based on fruit and/or vegetable juices including concentrate and dehydrated juices Fruit-flavoured drinks Cocoa beverages preparation powder Other beverages Cereal bars	Food Supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children Specified food category Food Supplements as defined in Directive adult population 3 g/day for children from 3 to 9 years of age Maximum levels of pure beta-glucans from yeast (Saccharomyces cervisiae) Food supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children Total diet replacement for weight control as defined in Regulation (EU) No 609/2013 Food for special medical purposes as defined in Regulation (EU) No 609/2013, excluding food for special medical purposes intended for infants and young children Beverages based on fruit and/or vegetable juices including concentrate and dehydrated juices Fruit-flavoured drinks O,8 g/kg Cocoa beverages preparation powder Other beverages Oceal bars Oscilier in Directive of age, adolescents and general adult popusation and general adult popusation of age, adolescents and general adult population of age, adolescents and general adult popusation of age, adolescents adult popusation of age, adol	Specified food category Section Food Supplements as defined in Directive 2002/46/EC, excluding food supplements for infants and young children Specified food category Section Section	Specified food category

Authorised novel food	Conditions under which the no	vel food may be used	Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Wholegrain and high fibre instant hot breakfast cereals	1,5 g/kg			
	Cookie-type biscuits	6,7 g/kg			
	Cracker-type biscuits	6,7 g/kg			
	Milk based beverages	3,8 g/kg			
	Fermented milk products	3,8 g/kg			
	Milk product analogues	3,8 g/kg			
	Dried milk/milk powder	25,5 g/kg			
	Soups and soup mixes	0,9 g/kg (ready to eat)			
		1,8 g/kg (condensed)			
		6,3 g/kg (powder)			
	Chocolate and confectionery	4 g/kg			
	Protein bars and powders	19,1 g/kg			
	Jam, marmalade and other fruit spreads	11,3 g/kg			
12					
Zeaxanthin	Specified food category	Maximum levels	The designation of the novel food		
	Food Supplements as defined in Directive 2002/46/EC	2 mg/day	on the labelling of the foodstuffs containing it shall be 'Zeaxanthin'.		
9					
Zinc L-pidolate	Specified food category	Maximum levels	The designation of the novel food		
	Foods covered by Regulation (EU) No 609/2013	3 g/day	on the labelling of the foodstuffs containing it shall be 'Zinc L-pidolate'		
	Milk based drinks and similar products intended for young children				
	Meal replacement for weight control				
	Foods intended to meet the expenditure of intense muscular effort, especially for sportsmen				

Authorised novel food	Conditions under which the novel food may be used A		Additional specific labelling requirements	Other requirements	► <u>M29</u> Data Protection ◀
	Food bearing statement on the absence or reduced presence of gluten in accordance with the requirements of Commission Implementing Regulation (EU) No 828/2014				
	Food Supplements as defined in Directive 2002/46/EC				

- (1) Regulation (EU) No 609/2013 of the European Parliament and of the Council of 12 June 2013 on food intended for infants and young children, food for special medical purposes, and total diet replacement for weight control and repealing Council Directive 92/52/EEC, Commission Directives 96/8/EC, 1999/21/EC, 2006/125/EC and 2006/141/EC, Directive 2009/39/EC of the European Parliament and of the Council and Commission Regulations (EC) No 41/2009 and (EC) No 953/2009 (OJ L 181, 29.6.2013, p. 35).
- (2) Commission Implementing Regulation (EU) No 828/2014 of 30 July 2014 on the requirements for the provision of information to consumers on the absence or reduced presence of gluten in food (OJ L 228, 31.7.2014, p. 5).
- (3) Directive 2002/46/EC of the European Parliament and of the Council of 10 June 2002 on the approximation of the laws of the Member States relating to food supplements (OJ L 183, 12.7.2002, p. 51).
- (4) Regulation (EC) No 1925/2006 of the European Parliament and of the Council of 20 December 2006 on the addition of vitamins and minerals and of certain other substances to foods (OJ L 404, 30.12.2006, p. 26).
- (5) Council Directive 2001/113/EC of 20 December 2001 relating to fruit jams, jellies and marmalades and sweetened chestnut purée intended for human consumption (OJ L 10, 12.1.2002, p. 67).
- (6) Regulation (EU) No 1308/2013 of the European Parliament and of the Council of 17 December 2013 establishing a common organisation of the markets in agricultural products and repealing Council Regulation (EEC) No 922/72, (EEC) No 234/79, (EC) No 1037/2001 and (EC) 1234/2007 (OJ L 347, 20.12.2013, p. 671).
- ►M32 (7) Maximum use levels in the final product ready for use, marketed as such or reconstituted as instructed by the manufacturer. ◄
- ▶ M45 (8) Council Directive 2001/112/EC of 20 December 2001 relating to fruit juices and certain similar products intended for human consumption (OJ L 10, 12.1.2002, p. 58).
- ▶ M46 (°) When used in milk products xylo-oligosaccharides shall not replace, in whole or in part, any milk constituent.
- (10) Maximum levels calculated on the basis of the specifications of Powder form 1. ◀
- ► M49 (11) The minimum specification for vitamin D content in vitamin D₂ mushroom powder of 1 000 µg vitamin D₂/gram of mushroom powder is used.

 (12) Depending on the age group the food supplement is intended for.

Authorised novel food	Conditions under which t	he novel food may be us	sed	Additional specific labelling Other requirements Data protection		Data protection
Frozen, dried and powder forms of <i>Locusta migratoria</i> (migratory locust)	Specified food category	Maximum levels (g/100 g) (marketed as such or reconstituted according to the instructions)		The designation of the novel food on the labelling of the foodstuffs containing it shall be 'frozen Locusta migratoria		Authorised on 5.12.2021. This inclusion is based on proprietary scientific evidence and scientific data protected in
(mgratory toeast)	1 9 7	Frozen Dried or Powder Powder Dried or powder Locusta migratoria (migratory locust)', 'dried/ powder Locusta migratoria (migratory locust)', 'Whole Locusta migratory locust)', 'Whole Locusta migratory locust', 'Whole M		accordance with Article 26 of Regulation (EU) 2015/2283. Applicant: Fair Insects BV, Industriestraat 3, 5107 NC		
	Frozen, dried and powder forms of Locusta migratoria			locust) powder' depending on the form used. 2. The labelling of the foodstuffs		Dongen, the Netherlands. During the period of data protection, the novel food is authorised for placing on the market
	Processed potato products; legumes-based dishes and pasta-based products	15	5	containing frozen dried or powder forms of <i>Locusta migratoria</i> (migratory locust) shall bear a statement that this		within the Union only by Fair Insects BV, unless a subsequent applicant obtains authorisation
	Meat analogues	80	50	ingredient may cause allergic reactions to consumers with known allergies to crustaceans,		for that novel food without reference to the proprietary scientific evidence or scientific
	Soups and concentrated soups	15	5	molluses and products thereof, and to mites. This statement shall appear in		data protected in accordance with Article 26 of Regu- lation (EU) 2015/2283, or with
	Canned/jarred legumes and vegetables	20	15	close proximity to the list of ingredients.		the agreement of Fair Insects BV. End date of the data protection: 5.12.2026.
	Salads	15	5			3.12.2020.
	Beer-like beverages, Alcoholic drink mixes	2	2			
	Chocolate confectionery	30	10			
	Nuts, oilseeds and chickpeas		20			
	Frozen fermented milk-based products	15	5			
	Sausages	30	10			

▼<u>M71</u>

	Authorised novel food	Conditions under which t	he novel food may be u	sed		Additional specific labelling requirements	Other requirements	Data protection
▼ <u>M78</u>	Frozen, dried and powder forms of	Specified food category	according to the instructions) food on the labelling of th	the designation of the novel food on the labelling of the		Authorised on 1 March 2022. This inclusion is based on proprietary scientific evidence		
	yellow mealworm (Tenebrio molitor		Frozen	Dried or powder		foodstuffs containing it shall be 'frozen yellow mealworm		and scientific data protected in accordance with Article 26 of
	larva)	Frozen, dried and powder forms of yellow mealworm (<i>Tenebrio molitor</i> larva)				(Tenebrio molitor larva)', 'dried yellow mealworm (Tenebrio molitor larva)', or		Regulation (EU) 2015/2283. Applicant: Fair Insects BV, Industriestraat 3, 5107 NC
		Multigrain bread and rolls; crackers and breadsticks	30	10	2.	'yellow mealworm (<i>Tenebrio molitor</i> larva) powder'. The labelling of the		Dongen, the Netherlands. During the period of data protection, the novel food is auth-
		Cereal bars	30	15		foodstuffs containing frozen,		orised for placing on the market
		Dried pasta based products; pasta based dishes (excluding dried puffed pasta); pizza and pizza-like dishes	15	10		dried and powder forms of yellow mealworm (<i>Tenebrio molitor</i> larva) shall bear a statement that this ingredient may cause allergic reactions to consumers with known allergies to crustaceans and products thereof and to dust mites. This statement shall appear in close proximity to the list of ingredients.	o a t t t t t t t t t t t t t t t t t t	within the Union only by Fair Insects BV, unless a subsequent applicant obtains authorisation for that novel food without reference to the proprietary scientific evidence or scientific data protected in accordance with Article 26 of Regulation (EU) 2015/2283, or with the agreement of Fair Insects BV. End date of the date protection:
		Dried stuffed pasta based products	30	15	1			
		Pre-mixes (dry) for baked products	30	15	1			
		Sauces	30	10	1			
		Potato, legumes based dishes	15	10				
		Whey powder	40	20	1			1 March 2027.
		Meat analogues	80	50				
		Soups and salads	20	5				
		Chips/crisps	40	20				
		Beer-like beverages; mixed alcoholic drinks; alcoholic drink mixes	1	1				
		Chocolate confectionary 30 10						
		Nuts, oilseeds and chickpeas	40	30				
		Frozen fermented milk-based products	15	5				
		Meat preparations	40	16				

Authorised Novel Food	Specifications
N-Acetyl-D-neuraminic acid	Description:
·	N-Acetyl-D-neuraminic acid is a white to off-white crystalline powder
	Definition:
	Chemical name:
	IUPAC names:
	N-Acetyl-D-neuraminic acid (dihydrate)
	5-Acetamido-3,5-dideoxy-D-glycero-D-galacto-non-2-ulopyranosonic acid (dihydrate)
	Synonyms:
	Sialic acid (dihydrate)
	Chemical formula:
	$C_{11}H_{19}NO_9$ (acid)
	$C_{11}H_{23}NO_{11}$ ($C_{11}H_{19}NO_9 * 2H_2O$) (dihydrate)
	Molecular mass:
	309,3 Da (acid)
	345,3 (309,3 + 36,0) (dihydrate)
	CAS No.:
	131-48-6 (free acid)
	50795-27-2 (dihydrate)
	Specifications:
	Description: white to off-white crystalline powder
	pH (20 °C, 5 % solution): 1,7 – 2,5
	N-Acetyl-D-neuraminic acid (dihydrate): > 97,0 %
	Water (dihydrate calculates to 10,4 %): \leq 12,5 % (w/w)
	Ash, sulphated: $< 0.2 \%$ (w/w)
	Acetic acid (as free acid and/or sodium acetate): < 0,5 % (w/w)
	Heavy Metals:
	Iron: < 20,0 mg/kg
	Lead: $< 0.1 \text{ mg/kg}$
	Residual proteins: < 0,01 % (w/w)

Residual solvents: 2-Propanol: < 0,1 % (w/w) Acetone: < 0,1 % (w/w) Ethyl acetate: < 0,1 % (w/w) Microbiological criteria: Salmonella: Absence in 25 g Aerobic mesophilic total count: < 500 CFU/g Enterobacteriaceae: Absence in 10 g Cronobacter (Enterobacter) sakazakii: Absence in 10 g Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Acetone: < 0,1 % (w/w) Ethyl acetate: < 0,1 % (w/w) Microbiological criteria: Salmonella: Absence in 25 g Aerobic mesophilic total count:< 500 CFU/g Enterobacteriaceae: Absence in 10 g Cronobacter (Enterobacter) sakazakii: Absence in 10 g Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Ethyl acetate: < 0,1 % (w/w) Microbiological criteria: Salmonella: Absence in 25 g Aerobic mesophilic total count:< 500 CFU/g Enterobacteriaceae: Absence in 10 g Cronobacter (Enterobacter) sakazakii: Absence in 10 g Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Microbiological criteria: Salmonella: Absence in 25 g Aerobic mesophilic total count:< 500 CFU/g Enterobacteriaceae: Absence in 10 g Cronobacter (Enterobacter) sakazakii: Absence in 10 g Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
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Aerobic mesophilic total count:< 500 CFU/g Enterobacteriaceae: Absence in 10 g Cronobacter (Enterobacter) sakazakii: Absence in 10 g Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Enterobacteriaceae: Absence in 10 g Cronobacter (Enterobacter) sakazakii: Absence in 10 g Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Cronobacter (Enterobacter) sakazakii: Absence in 10 g Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Listeria monocytogenes: Absence in 25 g Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Bacillus cereus: < 50 CFU/g Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Yeasts: < 10 CFU/g Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Moulds: < 10 CFU/g Residual endotoxins: < 10 EU/mg	
Residual endotoxins: < 10 EU/mg	
CELL Colony Forming Huiter ELL Fordstowin Huite	
CFU: Colony Forming Units; EU: Endotoxin Units.	
Adansonia digitata (Baobab) dried Description/Definition:	
fruit pulp The Baobab (Adansonia digitata) fruits are harvested from trees. The hard shells are cracked open and the pulp is sel is milled, separated into coarse and fine lots (particle size 3 to 600 μ) and then packaged.	parated from the seeds and the shell. This
Typical nutritional components:	
Moisture (loss on drying) (g/100 g): 4,5-13,7	
Protein (g/100 g): 1,8-9,3	
Fat (g/100 g): 0-1,6	
Total carbohydrate (g/100 g): 76,3-89,5	
Total sugars (as glucose): 15,2-36,5	
Sodium (mg/100 g): 0,1-25,2	
Analytical specifications:	
Foreign matter: Not more than 0,2 %	
Moisture (loss on drying) (g/100 g): 4,5-13,7	
Ash (g/100 g): 3,8-6,6	

Specifications
Description/Definition: Hydroalcoholic extract from <i>Ajuga reptans</i> L. tissue cultures which is substantially equivalent to extracts from flowering aerial parts of <i>Ajuga reptans</i> obtain by traditional cultures.
Pasteurised Akkermansia muciniphila (strain ATCC BAA-835, CIP 107961) is produced by anaerobic growth of the bacteria followed by pasteurisatic concentration of the cells, cryopreservation, and freeze drying. Characteristics/Composition: Total A. muciniphila cell count (cells/g): 2,5 × 10 ¹⁰ to 2,5 × 10 ¹² Viable A. muciniphila cell count (CFU/g): < 10 (LoD)(*) Water activity: ≤ 0,43 Moisture (%): ≤ 12,0 Protein (%): ≤ 35,0 Fat (%): ≤ 4,0 Crude ash (%): ≤ 21,0 Carbohydrates (%): 36,0 − 86,0 Microbiological criteria:Aerobic mesophilic total count: ≤ 500 CFU(**)/g Sulphite reducing anaerobes: ≤ 50 CFU/g Coagulase* Staphylococi: ≤ 10 CFU/g Enterobacteriaceae: ≤ 10 CFU/g Yeast: ≤ 10 CFU/g Mould: ≤ 10 CFU/g Bacillus cereus: ≤ 100 CFU/g Listeria spp.: Absence in 25 g Salmonella spp.: Absence in 25 g Escherichia coli: Absence in 1 g

Authorised Novel Food	Specifications
L-Alanyl-L-Glutamine	Description/Definition:
	L-Alanyl-L-Glutamine is produced by fermentation with a genetically modified strain of <i>Escherichia coli</i> . During the fermentation process, the ingredient is secreted into the growth medium from which it is subsequently separated and purified to a concentration of > 98 %.
	Appearance: White crystalline powder
	Purity: > 98 %
	Infrared spectroscopy: Conformity with ref. standard
	Appearance of solution: Colourless and clear
	Assay (dry basis): 98-102 %
	Related substances (each): $\leq 0.2 \%$
	Residue on ignition: ≤ 0.1 %
	Loss on drying: ≤ 0,5 %
	Optical rotation: +9,0 - +11,0°
	pH (1 %; H ₂ O): 5,0-6,0
	Ammonium (NH ₄): $\leq 0,020 \%$
	Chloride (Cl): ≤ 0,020 %
	Sulphate (SO ₄): $\leq 0.020 \%$
	Microbiological criteria:
	Escherichia coli: Absence/g
Algal oil from the microalgae <i>Ulkenia</i> sp.	Description/Definition:
Sp.	Oil from the micro-algae <i>Ulkenia</i> sp.
	Acid value: ≤ 0.5 mg KOH/g
	Peroxide value (PV): ≤ 5,0 meq/kg oil
	Moisture and volatiles: $\leq 0.05 \%$
	Unsaponifiables: ≤ 4,5 %
	Trans-fatty acids: $\leq 1.0 \%$
	DHA content: ≥ 32 %

Authorised Novel Food	Specifications
5	
Allanblackia seed oil	Description/Definition:
	Allanblackia seed oil is obtained from the seeds of the allanblackia species: A. floribunda (synonymous with A. parviflora) and A. stuhlmannii.
	Composition of fatty acids (as a % of the total fatty acids):
	Lauric acid — Myristic acid — Palmitic acid (C12:0 - C14:0 - C16:0): sum of these acids < 4,0 %
	Stearic acid (C18:0): 45-58 %
	Oleic acid (C18:1): 40-51 %
	Poly unsaturated fatty acids (PUFA): < 2 %
	Characteristics:
	Free fatty acids: max 0,1 % of total fatty acids
	Trans fatty acids: max 1,0 % of total fatty acids
	Peroxide value: max 1,0 meq/kg
	Unsaponifiable matter: max 1,0 % (w/w) of the oil
	Saponification value: 185-198 mg KOH/g
Aloe macroclada Baker leaf extra	ct Description/Definition:
	Powdered gel extract derived from the leaves of <i>Aloe macroclada</i> Baker which is substantially equivalent to the same gel derived from <i>Aloe vera</i> (L.) Burn leaves.
	Ash: 25 %
	Dietary fibres: 28,6 %
	Fat: 2,7 %
	Moisture: 4,7 %
	Polysaccharides: 9,5 %
	Protein: 1,63 %
	Glucose: 8,9 %

	Authorised Novel Food	Specifications
<u>M23</u>		
	Antarctic Krill oil from Euphausia	Description/Definition:
	superba	To produce lipid extract from Antarctic Krill (<i>Euphausia superba</i>) deep-frozen crushed krill or dried krill meal is subjected to lipid extraction with a approved extraction solvent (under Directive 2009/32/EC). Proteins and krill material are removed from the lipid extract by filtration. The extraction solven and residual water are removed by evaporation.
		Saponification value: ≤ 230 mg KOH/g
		Peroxide value (PV): $\leq 3 \text{ meq } O_2/kg \text{ oil}$
		Oxidative stability: All food products containing Antarctic Krill oil from <i>Euphausia superba</i> should demonstrate oxidative stability by appropriate a recognised national/international test methodology (e.g. AOAC).
		Moisture and volatiles: ≤ 3 % or 0,6 expressed as water activity at 25 °C
		Phospholipids: ≥ 35 % to < 60 %
		Trans-fatty acids: ≤ 1 %
		EPA (eicosapentaenoic acid): ≥ 9 %
		DHA (docosahexaenoic acid): ≥ 5 %
М9		
	Antarctic Krill oil rich in phosp-	Description/Definition:
	holipids from Euphausia superba	Oil rich in phospholipids is produced from Antarctic krill (<i>Euphausia superba</i>) by repeated solvent washings with an approved solvent (under Directive 200 32/EC) to increase phospholipid content of the oil. Solvents are removed from the final product by evaporation.
		Saponification value: ≤ 230 mg KOH/g
		Peroxide value (PV): $\leq 3 \text{ meq } O_2/kg \text{ oil}$
		Moisture and volatiles: ≤ 3 % or 0,6 expressed as water activity at 25 °C
		Phospholipids: ≥ 60 %
		Trans-fatty acids: ≤ 1 %
		EPA (eicosapentaenoic acid): ≥ 9 %
		DHA (docosahexaenoic acid): ≥ 5 %

the fungus Mortierella alpina using a suitable liquid. The oil is then extracted from the biomass and purified. Arachidonic acid: ≥ 40 % by weight of the total fatty acid content Free fatty acids: ≤ 0,45 % of the total fatty acid content Trans fatty acids: ≤ 0,5 % of the total fatty acid content Unsaponifiable matter: ≤ 1,5 % Peroxide value (PV): ≤ 5 meq/kg Anisidin value: ≤ 20 Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition:	Authorised Novel Food	Specifications		
The clear yellow arachidonic acid-rich oil is obtained by fermentation of the non-genetically modified strains IS-4, I49-N18, FJRK-MA01 and CBS 210.32 of the fungus. Mortiverella alpina using a suitable liquid. The oil is then extracted from the biomass and purified. Arachidonic acid: \(\pm \) 40 by tweight of the total fatty acid content Free fatty acids: \(\pm \) 0.45 % of the total fatty acid content Trans fatty acids: \(\pm \) 0.5 % of the total fatty acid content Unsaponifiable matter: \(\pm \) 1.5 % meq/kg Anisidin value: \(\pm \) 20 Acid value: \(\pm \) 0.5 % Description/Definition: Argan oil from Argania spinosa Description/Definition: Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 43-50 % Linolcic acid (C18:1): 43-50 % Linolcic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total tocopherols: 16-90 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acid(iy: 0,2-1,5 %				
In clear yellow arachidonic acid-rich oil is obtained by termentation of the funiges modified strains IS-4, 149-N18, FJRK-MA01 and CBS 210-32 of the fungus Moriterella alpina using a suitable liquid. The oil is then extracted from the biomass and purified. Arachidonic acid: ≥ 40 % by weight of the total fatty acid content Free fatty acids: ≤ 0.45 % of the total fatty acid content Unsaponifiable matter: ≤ 1,5 % Peroxide value (PV): ≤ 5 meq/kg Anisidin value: ≤ 20 Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:1): 43-50 % Linoleic acid (C18:1): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Description/Definition:		
Free fatty acids: ≤ 0,45 % of the total fatty acid content Trans fatty acids: ≤ 0,5 % of the total fatty acid content Unsaponifiable matter: ≤ 1,5 % Peroxide value (PV): ≤ 5 meq/kg Anisidin value: ≤ 20 Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Description/Definition: Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:1): 43-50 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Oleic acidity: 0,2-1,5 %	tungus <i>Mortierella alpina</i>	The clear yellow arachidonic acid-rich oil is obtained by fermentation of the non-genetically modified strains IS-4, I49-N18, FJRK-MA01 and CBS 210.32 of the fungus <i>Mortierella alpina</i> using a suitable liquid. The oil is then extracted from the biomass and purified.		
Trans fatty acids: ≤ 0,5 % of the total fatty acid content Unsaponifiable matter: ≤ 1,5 % Peroxide value (PV): ≤ 5 meq/kg Anisidin value: ≤ 20 Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Description/Definition: Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Arachidonic acid: ≥ 40 % by weight of the total fatty acid content		
Unsaponifiable matter: ≤ 1,5 % Peroxide value (PV): ≤ 5 meq/kg Anisidin value: ≤ 20 Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Description/Definition: Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:1): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Free fatty acids: ≤ 0.45 % of the total fatty acid content		
Peroxide value (PV): ≤ 5 meq/kg Anisidin value: ≤ 20 Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Description/Definition: Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Trans fatty acids: ≤ 0,5 % of the total fatty acid content		
Anisidin value: ≤ 20 Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acid(ity: 0,2-1,5 %		Unsaponifiable matter: ≤ 1,5 %		
Acid value: ≤ 1,0 KOH/g Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Peroxide value (PV): ≤ 5 meq/kg		
Moisture: ≤ 0,5 % Description/Definition: Argan oil from Argania spinosa Description/Definition: Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Anisidin value: ≤ 20		
Argan oil from Argania spinosa Description/Definition: Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of Argania spinosa (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Acid value: ≤ 1,0 KOH/g		
Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of <i>Argania spinosa</i> (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Moisture: $\leq 0.5 \%$		
Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of <i>Argania spinosa</i> (L.) Skeels. Kernels may be roasted prior to pressing but with no direct contact with a flame. Composition: Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %				
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Palmitic acid (C16:0): 12-15 % Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Argan oil is the oil obtained by cold pressing of the almond like kernels of the fruits of <i>Argania spinosa</i> (L.) Skeels. Kernels may be roasted prior to pressing, but with no direct contact with a flame.		
Stearic acid (C18:0): 5-7 % Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Composition:		
Oleic acid (C18:1): 43-50 % Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Palmitic acid (C16:0): 12-15 %		
Linoleic acid (C18:2): 29-36 % Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Stearic acid (C18:0): 5-7 %		
Unsaponifiable matter: 0,3-2 % Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Oleic acid (C18:1): 43-50 %		
Total sterols: 100-500 mg/100 g Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Linoleic acid (C18:2): 29-36 %		
Total tocopherols: 16-90 mg/100 g Oleic acidity: 0,2-1,5 %		Unsaponifiable matter: 0,3-2 %		
Oleic acidity: 0,2-1,5 %		Total sterols: 100-500 mg/100 g		
		Total tocopherols: 16-90 mg/100 g		
Peroxide value (PV): < 10 meq O ₂ /kg		Oleic acidity: 0,2-1,5 %		
		Peroxide value (PV): < 10 meq O ₂ /kg		

Authorised Novel Food

Astaxanthin-rich oleoresin from Haematococcus pluvialis algae	Description/Definition:
Themalococcus pluvillus aigae	Astaxanthin is a carotenoid produced by <i>Haematococcus pluvialis</i> algae. Production methods for the growth of the algae are variable; using either closed systems exposed to sunlight or strictly controlled illuminated light; alternatively open ponds may be used. The algal cells are harvested and dried; the oleoresin is extracted using either super critical CO ₂ or a solvent (ethyl acetate). The Astaxanthin is diluted and standardized to 2,5 %, 5,0 %, 7,0 %, 10 %, 15 % or 20 % using olive oil, safflower oil or MCT (Medium Chain Triglycerides).
	Composition of the Oleoresin:
	Fat: 42,2- 99 %
	Protein: 0,3-4,4 %
	Carbohydrate: 0-52,8 %
	Fibre: < 1,0 %
	Ash: 0,0-4,2 %
	Specification of Carotenoids w/w%
	Total Astaxanthins: 2,9-11,1 %
	9-cis-astaxanthin: 0,3-17,3 %
	13-cis-astaxanthin: 0,2-7,0 %
	Astaxanthin monoesters: 79,8-91,5 %
	Astaxanthin diesters: 0,16-19,0 %
	B-Carotene: 0,01-0,3 %
	Lutein: 0-1,8 %
	Canthaxanthin: 0-1,30 %
	Microbiological criteria:
	Total aerobic bacteria: < 3 000 CFU/g
	Yeast and Moulds: < 100 CFU/g
	Coliforms: < 10 CFU/g
	E. coli: Negative
	Salmonella: Negative
	Staphylococcus: Negative

Specifications

Authorised Novel Food	Specifications		
Basil seeds (Ocimum basilicum)	Description/Definition:		
	Basil (Ocimum basilicum L.) belongs to the family 'Lamiaceae' within the order 'Lamiales'. Post-harvest the seeds are cleaned mechanically. Flowers, leaves and other parts of the plant are removed. Highest level of purity of Basil seeds has to be ensured by filtering (optical, mechanical). Production process of fruit juice and fruit/vegetable blend beverages containing Basil seeds (Ocimum basilicum L.) includes seed pre-hydration and pasteurisation steps. Microbiological controls and monitoring systems are in place.		
	Dry Matter: 94,1 %		
	Protein: 20,7 %		
	Fat: 24,4 %		
	Carbohydrate: 1,7 %		
	Dietary Fibre: 40,5 % (Method: AOAC 958,29)		
	Ash: 6,78 %		
Betaine	Description/Definition:		
	Betaine (N,N,N-trimethylglycine or carboxy-N,N,N-trimethylmethanaminium), in anhydrous (CH ₃) ₃ N ⁺ CH ₂ COO ⁻ (CAS No: 107-43-7) and monohydrate (CH ₃) ₃ N ⁺ CH ₂ COO ⁻ .H ₂ O (CAS No: 590-47-6) forms is obtained from processing of sugar beets (i.e. molasses, vinasses or betaine-glycerol).		
	Characteristics/Composition		
	Appearance: Free-flowing white crystals		
	Betaine: ≥ 99,0 % (w/w on dry weight basis)		
	Moisture: ≤ 2,0 % (anhydrous); ≤ 15,0 % (monohydrate)		
	Ash: $\leq 0.1 \%$		
	pH: 5,0-7,0		
	Residual protein: ≤ 1,0 mg/g		
	Heavy metals:		
	Arsenic: < 0,1 mg/kg		
	Mercury: < 0,005 mg/kg		
	Cadmium: < 0,01 mg/kg		
	Lead: < 0,05 mg/kg		

▼<u>M32</u>

	Authorised Novel Food	Specifications
		Microbiological criteria:
		Total viable count: ≤ 100 CFU/g
		Coliforms: Negative/10 g
		Salmonella sp.: Negative/25 g
		Yeast: $\leq 10 \text{ CFU/g}$
		Mould: ≤ 10 CFU/g
		CFU: Colony Forming Units.
▼ <u>M9</u>		
	Fermented black bean extract	Description/Definition:
		Fermented black bean extract (Touchi extract) is a fine light-brown protein-rich powder obtained by water extraction of small soybeans (<i>Glycine max (L.) Merr.</i>) fermented with <i>Aspergillus oryzae</i> . The extract contains an α-glucosidase inhibitor.
		Characteristics:
		Fat: ≤ 1,0 %
		Protein: ≥ 55 %
		Water: ≤ 7,0 %
		Ash: ≤ 10 %
		Carbohydrate: ≥ 20 %
		α-glucosidase inhibitory activity: IC50 min 0,025 mg/ml
		Soy isoflavone: $\leq 0.3 \text{ g/}100 \text{ g}$

Authorised Novel Food	Specifications
Bovine lactoferrin	Description/Definition:
	Bovine lactoferrin is a protein that occurs naturally in cows' milk. It is an iron-binding glycoprotein of approximately 77 kDa and consists of a single polypeptide chain of 689 amino acids.
	Production process: Bovine lactoferrin is isolated from skimmed milk or cheese whey via ion exchange and subsequent ultra-filtration steps. Finally, it is dribby freeze drying or spraying and the large particles are sieved out. It is a virtually odourless, light pinkish powder.
	Physical-Chemical properties of Bovine lactoferrin:
	Moisture: < 4,5 %
	Ash: < 1,5 %
	Arsenic: < 2,0 mg/kg
	Iron: < 350 mg/kg
	Protein: > 93 %
	of which bovine lactoferrin: > 95 %
	of which other proteins: < 5,0 %
	pH (2 % solution, 20 °C): 5,2-7,2
	Solubility (2 % solution, 20 °C): complete
Bovine milk basic whey protein	Description
isolate	Bovine milk basic whey protein isolate is a yellowish grey powder obtained from bovine skimmed milk via a series of isolation and purification ste
	Characteristics/Composition
	Total protein (w/weight of product): ≥ 90 %
	Lactoferrin (w/weight of product): 25-75 %
	Lactoperoxidase (w/weight of product): 10-40 %
	Other proteins (w/weight of product): ≤ 30 %
	TGF-β2: 12-18 mg/100 g
	Moisture: ≤ 6,0 %
	pH (5 % solution w/v): 5,5 - 7,6

▼<u>M34</u>

Authorised Novel Food	Specifications
	Lactose: ≤ 3,0 %
	Fat: ≤ 4,5 %
	Ash: $\leq 3.5 \%$
	Iron: $\leq 25 \text{ mg/}100 \text{ g}$
	Heavy Metals
	Lead: < 0,1 mg/kg
	Cadmium: < 0,2 mg/kg
	Mercury: < 0,6 mg/kg
	Arsenic: < 0,1 mg/kg
	Microbiological criteria:
	Aerobic mesophilic count: ≤ 10 000 CFU/g
	Enterobacteriaceae: ≤ 10 CFU/g
	Escherichia coli: Negative/g
	Coagulase positive Staphylococci: Negative/g
	Salmonella: Negative/25 g
	Listeria: Negative/25 g
	Cronobacter spp.: Negative/25 g
	Moulds: $\leq 50 \text{ CFU/g}$
	Yeasts: $\leq 50 \text{ CFU/g}$
	CFU: Colony Forming Units
 M9	
	
Buglossoides arvensis seed oil	Description/Definition:
	Refined Buglossoides oil is extracted from the seeds of Buglossoides arvensis (L.) I.M.Johnst
	Alpha-linolenic acid: ≥ 35 % w/w of total fatty acids
	Stearidonic acid: ≥ 15 % w/w of total fatty acids
	Linoleic acid: ≥ 8,0 % w/w of total fatty acids
	Trans fatty acids: ≤ 2,0 % w/w of total fatty acids

Authorised Novel Food	Specifications
	Acid value: ≤ 0,6 mg KOH/g
	Peroxide value (PV): ≤ 5.0 meq O_2/kg
	Unsaponifiable content: ≤ 2,0 %
	Protein content (total nitrogen): ≤ 10 μg/ml
	Pyrrolizidine alkaloids: Not detectable with a detection limit of 4,0 μg/kg
Calanus finmarchicus oil	Description/Definition:
•	The novel food is ruby coloured, slightly viscous oil with a slight shellfish odour extracted from the crustacean (marine zooplankton) <i>Calanus finmarchicus</i> . The ingredient consists primarily of wax esters (> 85 %) with minor amounts of triglycerides and other neutral lipids.
	Specifications:
	Water: < 1,0 %
	Wax esters: > 85 %
	Total fatty acids: > 46 %
	Eicosapentaenoic acid (EPA): > 3,0 %
	Docosahexaenoic acid (DHA): > 4,0 %
	Total fatty alcohols: > 28 %
	C20:1 n-9 fatty alcohol: > 9,0 %
	C22:1 n-11 fatty alcohol: > 12 %
	Trans fatty acids: < 1,0 %
	Astaxanthinesters: < 0,1 %
	Peroxide value (PV): < 3.0 meq. O_2/kg
<u>174</u>	
Calcium fructoborate	Description/Definition
	The novel food is calcium fructoborate, a calcium salt tetrahydrate of a bis(fructose) ester of boric acid in the form of a powder, represented by Ca[(C6H10O6)2B]2•4H2O, with a molecular mass of 846 Da.

	Authorised Novel Food	Specifications
		The novel food is produced by chemical synthesis whereby fructose is combined with boric acid in water to produce a bis(fructose) ester of boric acid through various heating and mixing processes. Calcium carbonate is then added to produce a solution containing the calcium salt of fructoborate (tetrahydrate). The solution is freeze-dried, ground to produce the final powdered product, and then packaged and stored under representative storage conditions (22 ± 1°C RH 55-60 %).
		Characteristics/composition
		Free moisture: < 5,0 %
		Calcium: 4,5-5 %
		Boron: 2,5-2,9 %
		Fructose: 80-85 %
		Ash: 15-16 %
		Heavy metals
		Arsenic: ≤ 1 mg/kg
		Microbiological criteria
		Total plate count: $\leq 1~000~\text{CFU/g}^{\text{(a)}}$
		Yeast and mould: < 100 CFU/g
		Coliforms: $\leq 10 \text{ CFU/g}$
		Escherichia coli: < 10 CFU/g
		Salmonella spp.: Absence in 25 g
		Coagulase-positive staphylococci: Absence in 1 g
		(a) CFU: colony forming units
▼ <u>M66</u>		
	Calcium L-Methylfolate	Description: The novel food is produced by chemical synthesis starting from folic acid. It is a white to light yellowish, almost odourless, crystalline powder, sparingly soluble in water and very slightly soluble or insoluble in most organic solvents.

Authorised Novel Food

Specifications		

Definition:

Chemical formula: C₂₀H₂₃CaN₇O₆

Systematic name: N-{4-[[((6S)-2-amino-1,4,5,6,7,8-hexahydro-5-methyl-4-oxo-6-pteridinyl)methyl]amino]benzoyl}-L-glutamic acid, calcium salt.

CAS Numbers: 129025-21-4 (Calcium salt with an unspecified ratio of L-5-MTHF/ Ca^{2+}) and 151533-22-1 (Calcium salt with specified 1:1 ratio of L-5-MTHF/ Ca^{2+}).

Molecular weight: 497,5 Daltons

Synonyms: L-methylfolate, calcium; L-5-methyltetrahydrofolic acid, calcium salt [(L-5-MTHF-Ca)]; (6S)-5-methyltetrahydrofolic acid, calcium salt [(6S)-5-MTHF-Ca)]; (6S)-5-methyl-5,6,7,8-tetrahydropteroyl-L-glutamic acid, calcium salt, and L-5-methyl-tetrahydrofolic acid (L-5-MTHF) without the cation specified.

Structural formula:

Characteristics

Purity: > 95 % (Dry basis)

Water: ≤ 17.0 %

Calcium (on anhydrous and solvent free basis): 7.0 - 8.5 %

Calcium D-methylfolate (6R, α S isomer): $\leq 1.0 \%$

▼<u>M66</u>

▼<u>M9</u>

Authorised Novel Food	Specifications
	Other folates and related substances: ≤ 2.5 %
	Ethanol: $\leq 0.5\%$
	Lead: $\leq 1 \text{ mg/kg}$
	Boron: ≤ 10 mg/kg
	Cadmium $\leq 0.5 \text{ mg/kg}$
	$Mercury \le 1.0 mg/kg$
	Arsenic ≤ 1.5 mg/kg
	Platinum ≤ 2 mg/kg
	Microbiological criteria:
	Total viable aerobic counts: ≤ 1 000 CFU/g
	Total yeast and mould count: ≤ 100 CFU/g
	CFU: Colony Forming Units
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Chewing gum base (monome- thoxypolyethylene glycol)	Description/Definition:
thorypolyculytene glycoly	The novel food ingredient is a synthetic polymer (Patent number WO2006016179). It consists of branched polymers of monomethoxypolyethylene glycol (MPEG) grafted onto polyisoprene-graft-maleic anhydride (PIP-g-MA), and unreacted MPEG (less than 35 % by weight).
	White to off-white colour.
	CAS No.: 1246080-53-4
	Characteristics:
	Moisture: < 5,0 %
	Aluminium: < 3,0 mg/kg
	Lithium: < 0,5 mg/kg
	Nickel: < 0,5 mg/kg
	Residual anhydride: < 15 μmol/g
	Polydispersity index: < 1,4
	Isoprene: < 0,05 mg/kg
	Ethylene oxide: < 0,2 mg/kg

Authorised Novel Food	Specifications
	Free maleic anhydride: < 0,1 %
	Total oligomeres (less than 1 000 Dalton): ≤ 50 mg/kg
	Ethylene glycol: < 200 mg/kg
	Diethylene glycol: < 30 mg/kg
	Monoethylene glycol methyl ether: < 3,0 mg/kg
	Diethylene glycol methyl ether: < 4,0 mg/kg
	Triethylene glycol methyl ether: < 7,0 mg/kg
	1,4-Dioxane: < 2,0 mg/kg
	Formaldehyde: < 10 mg/kg
Chewing gum base (Methyl vinyl	Description/Definition:
ether-maleic anhydride copolymer)	Methyl vinyl ether-maleic anhydride copolymer is an anhydrous copolymer of methyl vinyl ether and maleic anhydride.
	Free-flowing, white to white-off powder
	CAS No: 9011-16-9
	Purity:
	Assay value: At least 99,5 % in dry matter
	Specific viscosity (1 % MEK): 2-10
	Residual methyl vinyl ether: ≤ 150 ppm
	Residual maleic anhydride: ≤ 250 ppm
	Acetaldehyde: $\leq 500 \text{ ppm}$
	Methanol: ≤ 500 ppm
	Dilauroyl peroxide: ≤ 15 ppm
	Total heavy metals: \(\leq 10 \) ppm
	Microbiological criteria:
	Total aerobic plate count: ≤ 500 CFU/g
	Mould/yeast: ≤ 500 CFU/g
	Escherichia coli: Negative to test
	Salmonella: Negative to test
	Staphylococcus aureus: Negative to test
	Pseudomonas aeruginosa: Negative to test

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Authorised Novel Food	Specifications
Chia oil from Salvia hispanica	Description/Definition:
	Chia oil is produced from Chia (Salvia hispanica L.) seeds (99,9 % pure) by cold pressing. No solvents are used and, once pressed, the oil is held in decantation tanks and a filtration process employed to remove impurities. It can also be produced by extraction with supercritical CO ₂ .
	Production process:
	Produced by cold pressing. No solvents are used and, once pressed, the oil is held in decantation tanks and a filtration process employed to remove impurities.
	Acidity expressed as oleic acid: ≤ 2,0 %
	Peroxide value (PV): ≤ 10 meq/kg
	Insoluble impurities: ≤ 0,05 %
	Alpha linolenic acid: ≥ 60 %
	Linoleic acid: 15-20 %
Chia seeds (Salvia hispanica)	Description/Definition:
	Chia (Salvia hispanica L.) is a summer annual herbaceous plant belonging to the Labiatae family. Post-harvest the seeds are cleaned mechanically. Flowers, leaves and other parts of the plant are removed.
	Dry matter: 90-97 %
	Protein: 15-26 %
	Fat: 18-39 %
	Carbohydrate (*): 18-43 %
	Crude Fibre(**): 18-43 %
	Ash: 3-7 %
	(*) Carbohydrates include the fibre value
	(**) Crude fibre is the part of fibre made mainly of indigestible cellulose, pentosans and lignin

Authorised Novel Food

	Production process:
	Production process of fruit juices and fruit juice blends beverages, containing Chia seeds, includes seed pre-hydration and pasteurisation steps. Microbiological controls and monitoring systems are in place.
Chitin-glucan from Aspergillus	Description/Definition:
niger	Chitin-glucan is obtained from the mycelium of <i>Aspergillus niger</i> ; it is a slightly yellow, odourless, free-flowing powder. It has a dry matter content of 90 % or more.
	Chitin-glucan is composed largely of two polysaccharides:
	— chitin, composed of repeating units of N-acetyl-D-glucosamine (CAS No: 1398-61-4),
	— beta (1, 3)-glucan, composed of repeating units of D-glucose (CAS No: 9041-22-9).
	Loss on drying: ≤ 10 %
	Chitin-glucan: ≥ 90 %
	Ratio of chitin to glucan: 30:70 to 60:40
	Ash: $\leq 3.0 \%$
	Lipids: ≤ 1,0 %
	Proteins: ≤ 6,0 %
Chitin-glucan complex from Fomes	Description/Definition:
fomentarius	Chitin-glucan complex is obtained from the cell walls of the fruit bodies of the fungus Fomes fomentarius. It consists primarily of two polysaccharides:
	— Chitin, composed of repeating units of N-acetyl-D-glucosamine (CAS No: 1398-61-4);
	— Beta-(1,3)(1,6)-D-glucan, composed of repeating units of D-glucose (CAS No: 9041-22-9).
	The production process consists of several steps, including: cleaning, reduction in size and grinding, softening in water and heating in an alkaline solution, washing, drying. No hydrolysis is applied during the production process.
	Appearance: Powder, odourless, flavourless, brown
	Purity:
	Moisture: ≤ 15 %
	Ash: $\leq 3.0 \%$
	Chitin-glucan: ≥ 90 %
	Ratio of chitin to glucan: 70:20
	Total carbohydrates, excluding glucans: ≤ 0,1 %

Specifications

Authorised Novel Food	Specifications
	Proteins: ≤ 2,0 %
	Lipids: ≤ 1,0 %
	Melanins: $\leq 8,3\%$
	Additives: None
	pH: 6,7-7,5
	Heavy metals:
	Lead (ppm): ≤ 1,00
	Cadmium (ppm): $\leq 1,00$
	Mercury (ppm): ≤ 0.03
	Arsenic (ppm): ≤ 0.20
	Microbiological criteria:
	Total mesophilic bacteria: $\leq 10^3/g$
	Yeast and moulds: $\leq 10^3/g$
	Coliforms at 30 °C: $\leq 10^3/g$
	$E. \ coli: \leq 10/g$
	Salmonella and other pathogenic bacteria: Absence/25 g
Chitosan extract from fungi	Description/Definition:
(Agaricus bisporus; Aspergillus	The chitosan extract (containing mainly poly(D-glucosamine)) is obtained from stems of Agaricus bisporus or from the mycelium of Aspergillus niger.
niger)	The patented production process consists of several steps, including: extraction and deacetylation (hydrolysis) in alkaline medium, solubilisation in acidic medium, precipitation in alkaline medium, washing and drying.
	Synonym: Poly(D-glucosamine)
	Chitosan CAS number: 9012-76-4
	Chitosan formula: (C ₆ H ₁₁ NO ₄) _n
	Appearance: fine free-flowing powder
	Aspect: Off –white to slightly brownish
	Odour: Odourless
	Purity:
	Chitosan content (% w/w dry weight):≥ 85
	Glucan content (% w/w dry weight): ≤ 15
	Loss on drying (% w/w dry weight): ≤ 10
	Viscosity (1 % in 1 % acetic acid): 1-15
	viscosity (1 /0 iii 1 /0 accite acita). 1-13

Authorised Novel Food	Specifications
	Degree of acetylation (in % mol/wet weight): 0-30
	Viscosity (1 % in 1 % acetic acid) (mPa.s): 1-14 for chitosan from Aspergillus niger; 12-25 for chitin from Agaricus bisporus
	Ash (% w/w dry weight): ≤ 3.0
	Proteins (% w/w dry weight): ≤ 2.0
	Particle size: > 100 nm
	Tapped density (g/cm ³): 0,7-1,0
	Fat binding capacity 800 × (w/w wet weight): pass
	Heavy metals:
	Mercury (ppm): ≤ 0.1
	Lead (ppm): ≤ 1,0
	Arsenic (ppm): ≤ 1.0
	Cadmium (ppm): ≤ 0.5
	Microbiological criteria:
	Aerobic count (CFU/g): $\leq 10^3$
	Yeast and mould count (CFU/g): $\leq 10^3$
	Escherichia coli (CFU/g): ≤ 10
	Enterobacteriaceae (CFU/g): ≤ 10
	Salmonella: Absence/25g
	Listeria monocytogenes: Absence/25g
Chondroitin sulphate	Description/Definition:
	Chondroitin sulphate (sodium salt) is a biosynthetic product. It is obtained by chemical sulphation of chondroitin derived from fermentation by the bacterium <i>Escherichia coli</i> O5:K4:H4 strain U1-41 (ATCC 23502).
	Chondroitin sulphate (sodium salt) (% dry basis): 95-105
	MWw (weight avg.) (kDa): 5-12
	MWn (number avg.) (kDa): 4-11
	Dispersity $(w_h/w_{0,05})$: ≤ 0.7
	Sulphation pattern (ΔDi -6S) (%): ≤ 85
	Loss on drying (%) (105 °C to constant weight): ≤ 10,0
	Residue on ignition (% dry basis): 20-30
	Protein (% dry basis): ≤ 0.5
	Endotoxins (EU/mg): ≤ 100
	Total organic impurities (mg/kg): ≤ 50

▼<u>M54</u>

Authorised Novel Food	Specifications
Chromium Picolinate	Description/Definition:
	Chromium picolinate is a reddish free-flowing powder, slightly soluble in water at pH 7. The salt is also soluble in polar organic solvents.
	Chemical name: tris(2pyridinecarboxylato-N,O)chromium(III) or 2-pyridinecarboxylic acid chromium(III) salt
	CAS No.: 14639-25-9Chemical formula: Cr(C ₆ H ₄ NO ₂) ₃
	Chemical characteristics:
	Chromium Picolinate: ≥ 95 %
	Chromium (III): 12-13 %
	Chromium (VI): not detected
	Water: ≤ 4,0 %
Chromium-containing yeast	Description/Definition:
(Yarrowia lipolytica) biomass	The novel food is the dried and heat-killed chromium-containing biomass of the yeast <i>Yarrowia lipolytica</i> .
	The novel food is produced by fermentation in the presence of chromium chloride followed by a number of purification steps and a heat-killing step of the yeast to ensure the absence of viable <i>Yarrowia lipolytica</i> cells in the novel food.

Characteristics/Composition:

Total chromium: 18-23 µg/g

Chromium (VI): < 10 µg/kg (i.e. limit of detection)

Protein: 40–50 g/100 g

Dietary fibre: 24-32 g/100 g

Sugars: < 2 g/100 g

Fat: 6–12 g/100 g

Total ash: \leq 15 %

Water: ≤ 5 %

Dry matter: ≥ 95 %

Heavy metals:

Lead: $\leq 3.0 \text{ mg/kg}$ Cadmium: $\leq 1.0 \text{ mg/kg}$

Mercury: ≤ 0.1 mg/kg

▼<u>M54</u>

▼<u>M9</u>

Authorised Novel Food	Specifications
	Microbiological criteria:
	Total aerobic microbial count: $\leq 5 \times 10^3 \text{ CFU/g}$
	Total yeast and mould count: $\leq 10^2 \text{ CFU/g}$
	Viable Yarrowia lipolytica cells (14): < 10 CFU/g (i.e. limit of detection)
	Coliforms: ≤ 10 CFU/g
	Salmonella spp.: Absence in 25 g
	CFU: colony forming units
s incanus L. Pandalis herb	Description:

Cistus incanus L. Pandalis herb; species belonging to the Cistaceae family and native to the Mediterranean region, Chalkidiki Peninsula.

Composition:

Moisture: 9-10 g/100 g herbs Protein: 6,1 g/100 g herbs Fat: 1,6 g/100 g herbs

Carbohydrates: 50,1 g/100 g herbs

Fiber: 27,1 g/100 g herbs Minerals: 4,4 g/100 g herbs

Sodium: 0,18 g Potassium: 0,75 g Magnesium: 0,24 g Calcium: 1,0 g Iron: 65 mg Vitamin B₁: 3,0 μg Vitamin B₂: 30 μg Vitamin B₆: 54 μg Vitamin C: 28 mg

Vitamin A: less than 0,1 mg Vitamin E: 40-50 mg

Authorised Novel Food	Specifications
	Alpha-Tocopherol: 20–50 mg
	Beta and Gamma-Tocopherols: 2–15 mg
	Delta-Tocopherol: 0,1–2 mg
Citicoline	Description/Definition:
Citiconne	Citicoline is produced by a microbial process.
	Citicoline is composed of cytosine, ribose, pyrophosphate and choline.
	White crystalline powder
	Chemical name: Choline cytidine 5'-pyrophosphate, Cytidine 5'-(trihydrogen diphosphate) P'-[2-(trimethylammonio)ethyl]ester inner salt
	Chemical formula: C ₁₄ H ₂₆ N ₄ O ₁₁ P ₂
	Molecular weight: 488,32 g/mol
	CAS No.: 987-78-0
	pH (sample solution of 1 %): 2,5-3,5
	Purity:
	Assay value: ≥ 98 % of dry matter
	Loss on drying (100 °C for 4 hours): $\leq 5,0$ %
	Ammonium: ≤ 0,05 %
	Arsenic: Not more than 2 ppm
	Free phosphoric acids: ≤ 0,1 %
	5'-Cytidylic acid: ≤ 1,0 %
	Microbiological criteria:
	Total plate count: $\leq 10^3$ CFU/g
	Yeast and moulds: $\leq 10^2 \text{ CFU/g}$
	Escherichia coli: Absence in 1 g
Clostridium butyricum	Description/Definition:
	Clostridium butyricum (CBM-588) is a Gram-positive, spore-forming, obligate anaerobic, non-pathogenic, non-genetically modified bacterium. Depository number FERM BP-2789

▼M76

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Coffea arabica L. and/or Coffea
canephora Pierre ex A.Froehner
dried cherry pulp and its infusion
(Traditional food from a third

country)

Authorised Novel Food

Description/Definition:

Microbiological criteria:

Total viable aerobic count: ≤ 10³ CFU/g

Escherichia coli: Not detected in 1 g

Staphylococcus aureus: Not detected in 1 g

Pseudomonas aeruginosa: Not detected in 1 g

Yeast and moulds: $\leq 10^2$ CFU/g

The traditional food consists of the dried unroasted coffee cherry pulp of Coffea arabica L. and/or Coffea canephora Pierre ex A.Froehner (genus: Coffea family: Rubiaceae) and its infusion. The infusion can be used as such or concentrated or dried.

Specifications

Ripe coffee cherries are collected, and then the coffee beans are mechanically removed, prior or after a drying process, leaving the dried coffee cherry pulp, which can be milled to a powder.

The separated coffee cherry pulp is also known as 'cascara', from the Spanish 'cáscara', meaning 'husk'.

Typically, the infusion is prepared by mixing up to 6 g of cascara pulp or husk in 100 ml of hot water (> 75 °C) for a few minutes and then pouring through a strainer, or using corresponding amounts in dried or instant infusions.

Composition of the dried coffee cherry pulp:

Water: < 18 %

Water activity (a_w) : ≤ 0.65

Ash: < 10,4 % DM Protein: < 15 % DM Fat: < 5 % DM

Carbohydrates: < 85 % DM

Microbiological criteria:

Aerobic Plate Count: $< 10^4$ CFU/g Total yeasts and moulds: < 100 CFU/g

Enterobacteriaceae: < 50 CFU/g Salmonella: Absence in 25 g Bacillus cereus: < 100 CFU/g

▼<u>M76</u>

Authorised Novel Food	Specifications
	Mycotoxins:
	Ochratoxin A: < 5,0 µg/kg
	Aflatoxin B1: < 2,0 μg/kg
	Aflatoxin B1, B2, G1, G2 (as sum): < 4,0 μg/kg
	Heavy metals:
	Cadmium (Cd): < 0,05 mg/kg
	Lead (Pb): < 1,0 mg/kg
	Copper: ≤ 50 mg/kg
	Mercury: ≤ 0.02 mg/kg
	Arsenic: ≤ 0,2 mg/kg
	Impurities:
	Benzo(a)pyrene: < 10,0 μg/kg
	Sum of benzo(a)pyrene, benz(a)anthracene, benzo(b)fluoranthene and chrysene: < 50,0 µg/kg
	Pesticides:
	Pesticide levels in the traditional food shall comply with levels set by Regulation (EC) No 396/2005 for '0639000' for 'Herbal infusions from any other parts of the plant'.
	CFU: Colony Forming Units
	DM: Dry Matter
▼ <u>M29</u>	
D-ribose	Description
	D-ribose is an aldopentose monosaccharide which is produced by fermentation using a transketolase-deficient strain of Bacillus subtilis.
	Chemical formula: C ₅ H ₁₀ O ₅
	CAS No: 50-69-1
	Molecular mass: 150,13 Da

Authorised Novel Food	Specifications
	Characteristics/Composition
	Appearance: Dry with powdery texture, white to slightly yellow in colour
	Specific rotation $[\alpha]_D^{25}$: - 19,0° to - 21,0°
	D-ribose purity (% dry basis):
	-HPLC/RI (8) Method 98,0–102,0 %
	Ash: < 0,2 %
	Loss on drying (moisture): < 0,5 %
	Clarity on solution: ≥ 95 % transmittance
	Heavy metals
	Lead: $\leq 0.1 \text{ mg/kg}$
	Arsenic: ≤ 0,1 mg/kg
	Cadmium: ≤ 0,1 mg/kg
	Mercury: ≤ 0,1 mg/kg
	Microbiological criteria
	Total plate count: ≤ 100 CFU (9)/g
	Yeast: ≤ 100 CFU/g
	Moulds: ≤ 100 CFU/g
	Coliforms: ≤ 10 CFU/g
	Salmonella sp: Negative/25 g

▼<u>M52</u>

Authorised Novel Food

Transcribed Trover Food	Specifications
2	
Dried Euglena gracilis	Description/Definition:
	The novel food is dried whole cell Euglena, which is the dried biomass of the microalga Euglena gracilis.
	The novel food is produced by fermentation followed by filtration and a heat-killing step of the microalga to ensure the absence of viable Euglena gracilis cells in the novel food.
	Characteristics/Composition:
	Total carbohydrates: ≤ 75 %
	β-glucan: > 50 %
	Protein: ≥ 15 %
	Fat: ≤ 15 %
	Ash: ≤ 10 %
	Moisture: ≤ 6 %
	Heavy metals:
	Lead: $\leq 0.5 \text{ mg/kg}$
	Cadmium: ≤ 0,5 mg/kg
	Mercury: ≤ 0,05 mg/kg
	Arsenic: ≤ 0,02 mg/kg
	Microbiological criteria:
	Aerobic plate count: ≤ 10 000 CFU/g
	Coliforms: ≤ 100 MPN/g
	Yeast and mould: ≤ 500 CFU/g
	Escherichia coli: Absence in 10 g
	Staphylococcus aureus: Absence in 10 g
	Salmonella: Absence in 25 g
	Listeria monocytogenes: Absence in 25 g
	CFU: colony forming units.
	MPN: most probable number

Specifications

Authorised Novel Food	Specifications
Extract of defatted cocoa powder	Cocoa (<i>Theobroma cacao</i> L.) Extract
-	Appearance: Dark brown powder free of visible impurities
	Physical and chemical properties:
	Polyphenol content: Min 55,0 % GAE
	Theobromine content: Max 10,0 %
	Ash content: Max 5,0 %
	Moisture content: Max 8,0 %
	Bulk density: 0,40-0,55 g/cm ³
	pH: 5,0-6,5
	Residual solvent: Max 500 ppm
Low fat cocoa extract	Low fat Cocoa (<i>Theobroma cacao</i> L.) extract
	Appearance: Dark red to purple powder
	Cocoa extract, concentrate: Min 99 %
	Silicon dioxide (technological aid): Max 1,0 %
	Cocoa flavanols: Min. 300 mg/g
	— Epicatechin: Min. 45 mg/g
	Loss on drying: Max. 5,0 %
Coriander seed oil from <i>Coriandrum</i>	Description/Definition:
sativum	Coriander seed oil is an oil containing glycerides of fatty acids that is produced from the seeds of the coriander plant <i>Coriandrum sativum</i> L.
	Yellowish to brown colour, bland taste
	CAS No.: 8008-52-4
	Composition of fatty acids:
	composition of any without

▼<u>M67</u>

Authorised Novel Food	Specifications
	Stearic acid (C18:0): < 1,5 %
	Petroselinic acid (cis-C18:1(n-12)): 60-75 %
	Oleic acid (cis-C18:1 (n-9)): 7-15 %
	Linoleic acid (C18:2): 12-19 %
	α-Linolenic acid (C18:3): < 1,0 %
	Trans fatty acids: ≤ 1,0 %
	Purity:
	Refractive index (20 °C): 1.466-1.474
	Acid value: ≤ 4 mg KOH/g
	Peroxide value (PV): ≤ 5,0 meq/kg
	Iodine value: 88-110 units
	Saponification value: 179-200 mg KOH/g
	Unsaponifiable matter: ≤ 15 g/kg
▼ <u>M15</u>	
Cranberry extract powder	Description/Definition:
	Cranberry extract powder is a water-soluble phenolic-rich powder extract prepared through an ethanolic extraction from the juice concentrate of sound, mature berries of the cranberry cultivar <i>Vaccinium macrocarpon</i> .
	Characteristics/Composition
	Moisture (% w/w): ≤ 4
	Proanthocyanidins — PACs (% w/w dry weight)
	— OSC-DMAC method (³) (⁵): 55.0-60.0 or
	— BL-DMAC method (4) (5): 15.0-18.0
	Total phenolics (GAE (6), % w/w dry weight) (5)

— Folin-Ciocalteau method: > 46.2

Solubility (water): 100 %, with no visible insoluble particles

▼<u>M15</u>

▼<u>M9</u>

 $\alpha\text{-cyclodextrin}$

Description/Definition:

Authorised Novel Food	Specifications
	Ethanol Content (mg/kg): ≤ 100
	Screen Analysis: 100 % through 30 mesh screen
	Appearance and aroma, as powder: Free-flowing, deep red colour. Earthy aroma with no burnt character.
	Heavy metals:
	Arsenic (ppm): < 3
	Microbiological criteria:
	Yeast: $< 100 \text{ CFU } (^7)/\text{g}$
	Mould: < 100 CFU/g
	Aerobic plate count: < 1 000 CFU/g
	Coliforms: < 10 CFU/g
	Escherichia coli: < 10 CFU/g
	Salmonella: Absent in 375 g
Crataegus pinnatifida dried fruit	Description/Definition:
G I	Dried fruits of <i>Crataegus pinnatifida</i> species belonging to the <i>Rosaceae</i> family and native to north China and Korea.
	Composition:
	Dry matter: 80 %
	Carbohydrates: 55 g/kg fresh weight
	Fructose: 26,5–29,3 g/100 g
	Glucose: 25,5–28,1 g/100 g
	Vitamin C: 29,1 mg/100 g fresh weight
	Sodium: 2,9 g/100 g fresh weight
	Compotes are products obtained by thermal processing of the edible part of one or several species of fruits, whole or in pieces, sieved or not, without significant concentration. Sugars, water, cider, spices and lemon juice may be used.

A non-reducing cyclic saccharide consisting of six α -1,4-linked D-glucopyranosyl units produced by the action of cyclodextrin glucosyltransferase (CGTase, EC 2.4.1.19) on hydrolyzed starch. Recovery and purification of α -cyclodextrin may be carried out using one of the following procedures: precipitation of a complex of α -cyclodextrin with 1-decanol, dissolution in water at elevated temperature and re-precipitation, steam-stripping of the

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Authorised Novel Food	Specifications
	complexant, and crystallisation of α -cyclodextrin from the solution; or chromatography with ion-exchange or gel filtration followed by crystallisation of α -cyclodextrin from the purified mother liquor; or membrane separation methods such as ultra-filtration and reverse osmosis: Description: Virtually odourless, white or almost white crystalline solid.
	Synonyms: α-cyclodextrin, α-dextrin, cyclohexaamylose, cyclomaltohexaose, α-cycloamylase
	Chemical name: Cyclohexaamylose
	CAS No.: 10016-20-3
	Chemical formula: (C ₆ H ₁₀ O ₅) ₆
	Formula weight: 972,85
	Assay: $\geq 98\%$ (dry basis)
	Identification:
	Melting range: Decomposes above 278 °C
	Solubility: Freely soluble in water; very slightly soluble in ethanol
	Specific rotation: [α] _D ²⁵ : Between +145° and +151° (1 % solution)
	Chromatography: The retention time for the major peak in a liquid chromatogram of the sample corresponds to that for α-cyclodextrin in a chromatogram of reference α-cyclodextrin (available from <i>Consortium für Elektrochemische Industrie GmbH, München, Germany or Wacker Biochem Group, Adrian, MI, USA</i>) using the conditions described in the METHOD OF ASSAY
	Purity:
	Water: ≤ 11 % (Karl Fischer Method)
	Residual complexant: ≤ 20 mg/kg
	(1-decanol)
	Reducing substances: ≤ 0,5 % (as glucose)
	Sulphated ash: $\leq 0.1 \%$
	Lead: $\leq 0.5 \text{ mg/kg}$
	Method of assay:
	Determine by liquid chromatography using the following conditions:
	Sample solution: Weigh accurately about 100 mg of test sample into a 10 ml volumetric flask and add 8 ml of deionised water. Dissolve the sample completely using an ultra-sonification bath (10-15 min) and dilute to the mark with purified deionised water. Filter through a 0,45-micrometer filter

Authorised Novel Food	Specifications
	Reference solution: Weigh accurately about 100 mg of α-cyclodextrin into a 10 ml volumetric flask and add 8 ml of deionised water. Dissolve the sample completely using an ultra-sonification bath and dilute to the mark with purified deionised water.
	Chromatography: Liquid chromatograph equipped with a refractive index detector and an integrating recorder.
	Column and packing: Nucleosil-100-NH2 (10 µm) (Macherey & Nagel Co. Düren, Germany) or similar
	Length: 250 mm
	Diameter: 4 mm
	Temperature: 40 °C
	Mobile phase: acetonitrile/water (67/33, v/v)
	Flow rate: 2,0 ml/min
	Injection volume: 10 μlProcedure: Inject the sample solution into the chromatograph, record the chromatogram, and measure the area of the α-CD peak. Calculate the percentage of α-cyclodextrin in the test sample as follows:
	$\%$ α-cyclodextrin (dry basis) = $100 \times (A_S/A_R)$ (W_R/W_S)
	where
	A_S and A_R are the areas of the peaks due to α -cyclodextrin for the sample solution and reference solution, respectively.
	W _S and W _R are the weights (mg) of the test sample and reference α-cyclodextrin, respectively, after correcting for water content.
γ-cyclodextrin	Description/Definition:
-cyclodext iii	A non-reducing cyclic saccharide consisting of eight α -1,4-linked D-glucopyranosyl units produced by the action of cyclodextrin glucosyltransferase (CGTase, EC 2.4.1.19) on hydrolysed starch. Recovery and purification of γ -cyclodextrin may be carried out by precipitation of a complex of γ -cyclodextrin with 8-cyclohexadecen-1-one, dissolution of the complex with water and n-decane, steam-stripping of the aqueous phase and recovery of gamma-CD from the solution by crystallisation.
	Virtually odourless, white or almost white crystalline solid
	Synonyms: γ-cyclodextrin, γ-dextrin, cyclooctaamylose, cyclomaltooctaose, γ-cycloamylase
	Chemical name: Cyclooctaamylose
	CAS number: 17465-86-0
	Chemical formula: (C ₆ H ₁₀ O ₅) ₈
	Assay: ≥ 98 % (dry basis)

	Authorised Novel Food	Specifications
		Identification: Melting range: Decomposes above 285 °C Solubility: Freely soluble in water; very slightly soluble in ethanol Specific rotation: [α] _D ²⁵ : between + 174° and + 180° (1 % solution) Purity: Water: ≤ 11 % Residual complexant (8-cyclohexadecen-1-one (CHDC)): ≤ 4 mg/kg Residual solvent (n-decane): ≤ 6mg/kg
		Reducing substances: ≤ 0.5 % (as glucose) Sulphated ash: ≤ 0.1 %
▼ M21		Supraced usin 2 0,1 70
▼ <u>M21</u>	Decorticated grains of <i>Digitaria</i> exilis (Kippist) Stapf (fonio) (Traditional food from a third country)	Description/Definition The traditional food is the decorticated grain (bran removed) of Digitaria exilis (Kippist) Stapf. Digitaria exilis (Kippist) Stapf) is an annual herbaceous plant belonging to the Poaceae family. Typical nutritional components of decorticated grain of fonio Carbohydrates: 76,1 g/100 g of fonio Water: 12,4 g/100 g of fonio Protein: 6,9 g/100 g of fonio Fat: 1,2 g/100 g of fonio Fibre: 2,2 g/100 g of fonio Ash: 1,2 g/100 g of fonio Phytate content: ≤ 2,1 mg/g
▼ <u>M9</u>	Dextran preparation produced by Leuconostoc mesenteroides	1. Powdered form: Carbohydrates: 60 % with: (Dextran: 50 %, Mannitol: 0,5 %, Fructose: 0,3 %, Leucrose: 9,2 %) Protein: 6,5 %

Authorised Novel Food	Specifications
	Lipid: 0,5 %
	Lactic acid: 10 %
	Ethanol: traces
	Ash: 13 %
	Moisture: 10 %
	2. Liquid form:
	Carbohydrates: 12 % with: (Dextran: 6,9 %, Mannitol: 1,1 %, Fructose: 1,9 %, Leucrose: 2,2 %)
	Protein: 2,0 %
	Lipid: 0,1 %
	Lactic acid: 2,0 %
	Ethanol: 0,5 %
	Ash: 3,4 % Moisture: 80 %
	Workline. 60 70
Diacylglycerol oil of plant origin	Description/Definition:
	Manufactured from glycerol and fatty acids derived from edible vegetable oils, in particular from soybean oil (Glycine max) or rapeseed oil (Brassica campestris, Brassica napus) using a specific enzyme.
	Acylglycerol Distribution:
	Diacylglycerols (DAG): ≥ 80 %
	1,3-Diacylglycerols (1,3-DAG): ≥ 50 %
	Triacylglycerols (TAG): ≤ 20 %
	Monoacylglycerols (MAG): ≤ 5,0 %
	Fatty Acid Composition (MAG, DAG, TAG):
	Oleic acid (C18:1): 20-65 %
	Linoleic acid (C18:2): 15-65 %
	Linolenic acid (C18:3): ≤ 15 %
	Saturated fatty acids: ≤ 10 %

Authorised Novel Food	Specifications
	Others:
	Acid value: ≤ 0,5 mg KOH/g
	Moisture and volatile: ≤ 0,1 %
	Peroxide value (PV): ≤ 1,0 meq/kg
	Unsaponifiables: ≤ 2,0 %
	Trans fatty acids≤ 1,0 %
	MAG = monoacylglycerols, DAG = diacylglycerols, TAG = triacylglycerols
Dihydrocapsiate (DHC)	Description/Definition:
	Dihydrocapsiate is synthesised by enzyme-catalysed esterification of vanillyl alcohol and 8-methylnonanoic acid. Following the esterification dihydrocapsia is extracted with n-hexane.
	Viscous to colourless to yellow liquid
	Chemical formula: C ₁₈ H ₂₈ O ₄
	CAS No: 205687-03-2
	Physical-chemical properties:
	Dihydrocapsiate: > 94 %
	8-Methylnonanoic acid: < 6,0 %
	Vanillyl acohol: < 1,0 %
	Other synthesis related substances: < 2,0 %
Dried aerial parts of Hoodia	Description/Definition:
parviflora	It is the whole dried aerial parts of <i>Hoodia parviflora</i> N.E.Br., (family <i>Apocynaceae</i>)
	Characteristics/Composition
	Plant material: Aerial parts of at least 3-year-old plants
	Appearance: Light green to tan fine powder
	Solubility (water): > 25 mg/mL
	Moisture: < 5,5 %
	A_{w} : < 0,3

Authorised Novel Food	Specifications
	pH: < 5,0
	Protein: < 4,5 g/100 g
	Fat: $< 3 \text{ g}/100 \text{ g}$
	Carbohydrate (including dietary fibre): < 80 g/100 g
	Dietary fibre: < 55 g/100 g
	Total sugars: < 10,5 g/100 g
	Ash: < 20 %
	Hoodigosides
	P57: 5–50 mg/kg
	L: 1 000–6 000 mg/kg
	O: 500–5 000 mg/kg
	Total: 1 500–11 000 mg/kg
	Heavy metals:
	Arsenic: < 1,00 mg/kg
	Mercury: < 0,1 mg/kg
	Cadmium: < 0,1 mg/kg
	Lead: < 0,5 mg/kg
	Microbiological criteria:
	Aerobic plate count: $< 10^5 \text{ CFU/g}$
	Escherichia coli: < 10 CFU/g
	Staphylococcus aureus: < 50 CFU/g
	Total coliforms: < 10 CFU/g
	Yeast: $\leq 100 \text{ CFU/g}$
	Mould: $\leq 100 \text{ CFU/g}$
	Salmonella species: Negative/25 g
	Listeria monocytogenes: Negative/25 g
	CFU: Colony Forming Units

	Authorised Novel Food	Specifications
	Dried extract of Lippia citriodora	Description/Definition:
	from cell cultures	Dried extract of Lippia citriodora (Palau) Kunth from cell cultures HTN®Vb.
	Echinacea angustifolia extract from cell cultures	Description/Definition:
		Extract of the roots of Echinacea angustifolia obtained from plant tissue culture which is substantially equivalent to a root extract from Echinacea angustifolia obtained in ethanol-water titrated to 4 % echinacoside.
▼ <u>M31</u>		
	Echinacea purpurea extract from cell cultures	Description/Definition:
		Dried extract of <i>Echinacea purpurea</i> from cell cultures EchiPure-PC TM
▼ <u>M9</u>		
	Echium plantagineum oil	Description/Definition:
		Echium oil is the pale yellow product obtained by refining oil extracted from the seeds of <i>Echium plantagineum</i> L. Stearidonic acid: ≥ 10 % w/w of total fatty acids
		Trans fatty acids: ≤ 2,0 % (w/w of total fatty acids)
		Acid value: ≤ 0,6 mg KOH/g
		Peroxide value (PV): ≤ 5.0 meq O_2/kg
		Unsaponifiable content: ≤ 2,0 %
		Protein content (total nitrogen): ≤ 20 μg/ml
		Pyrrolizidine alkaloids: Not detectable with a detection limit 4,0 μg/kg

	Authorised Novel Food	Specifications
▼ <u>M50</u>		
	Ecklonia cava phlorotannins	Description/Definition
		Ecklonia cava phlorotannins are obtained via alcohol extraction from the edible marine alga Ecklonia cava. The extract is a dark brown powder, rich in phlorotannins, polyphenolic compounds found as secondary metabolites in certain brown algae species.
		Characteristics/Composition
		Phlorotannin content: 90 ± 5 %
		Antioxidant activity: > 85 %
		Moisture: < 5 %
		Ash: < 5 %
		Microbiological criteria
		Total viable cell count: < 3 000 CFU/g
		Mould/yeast: < 300 CFU/g
		Coliforms: Negative to test
		Salmonella spp.: Negative to test
		Staphylococcus aureus: Negative to test
		Heavy metals and Halogens
		Lead: < 3,0 mg/kg
		Mercury: < 0,1 mg/kg
		Cadmium: < 3,0 mg/kg
		Arsenic: < 25,0 mg/kg
		Inorganic Arsenic: < 0,5 mg/kg
		Iodine: 150,0 – 650,0 mg/kg
		CFU: Colony Forming Units
		Iodine: 150,0 - 650,0 mg/kg

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Authorised	Novel	Food

Specifications

▼<u>M18</u>

Egg membrane hydrolysate

Description

The egg membrane hydrolysate is derived from the eggshell membranes of chicken eggs. The eggshells undergo hydro-mechanical separation in order to obtain the egg membranes, which are then further processed using a patented solubilisation method. Following the solubilisation process, the solution is filtered, concentrated, spray-dried and packaged.

Characteristics/Composition

Chemical parameters

Total nitrogen-containing compounds (% w/w): ≥ 88

Collagen (% w/w): ≥ 15 Elastin (% w/w): ≥ 20

Total glycosaminoglycans (% w/w): ≥ 5

Calcium: ≤ 1 %

Physical parameters

pH: 6,5 - 7,6

Ash (% w/w): ≤ 8

Moisture (% w/w): ≤ 9

Water activity: ≤ 0.3

Solubility (in water): soluble

Bulk density: ≥ 0,6 g/cc

Heavy metals

Arsenic ≤ 0.5 mg/kg

Microbiological criteria

Aerobic plate count: ≤ 2 500 CFU/g

Escherichia coli: ≤ 5 MPN/g

Salmonella: Negative (in 25 g)

Coliforms: $\leq 10 \text{ MPN/g}$

Staphylococcus aureus: ≤ 10 CFU/g

Mesophilic spore count: ≤ 25 CFU/g

Thermophilic spore count: ≤ 10 CFU/10 g

Methods

Combustion according to AOAC 990.03 and AOAC 992.15

SircolTM Soluble Collagen Assay

FastinTM Elastin Assay

USP26 (chondroitin sulphate K0032 method)

▼<u>M18</u>

Authorised Novel Food	Specifications					
	Yeast: ≤ 10 CFU/g					
	Mould: ≤ 200 CFU/g					
	CFU: Colony Forming Units; MPN = Most Probable Number; USP: United States Pharmacopeia.					
			•			
Epigallocatechin gallate as a	Description/Definition:					
purified extract from green tea leaves (Camellia sinensis)	A highly purified extract from the leaves of green tea (Camellia sinensis (L.) Kuntze) in the form of a fine, off-white to pale pink powder. It is composed of minimum of 90 % epigallo-catechin gallate (EGCG), and has a melting point between approx. 210 and 215 °C					
	Appearance: off-white to pale pink powder					
	Chemical name: polyphenol (-) epigallocatechin-3-gallate					
	Synonyms: epigallocatechin gallate (EGCG)					
	CAS No.: 989-51-5					
	INCI name: epigallocatechin gallate					
	Molecular mass: 458,4 g/mol					
	Loss on drying: max 5,0 %					
	Heavy metals:					
	Arsenic: max 3,0 ppm					
	Lead: max 5,0 ppm					
	Assay:	Assay:				
	Min. 94 % EGCG (on dry material)					
	max. 0,1 % caffeine					
Solubility: EGCG is fairly soluble in water, ethanol, methanol and acetone						
L-ergothioneine	Definition Chemical name (IUPAC): (2S)-3-(2-thioxo-2,3-dihydro-1 <i>H</i> -imidazol-4-yl)-2-(trimethylammonio)-Propanoate Chemical formula: C ₉ H ₁₅ N ₃ O ₂ S					
	Molecular mass: 229,3 Da					
	CAS No.: 497-30-3					
	Parameter	Specification	Method			
	Appearance	White powder	Visual			
	Optical rotation	$[\alpha]_D \ge (+) \ 122^\circ \ (c = 1, H_2O)^{a)}$	Polarimetry			

Authorised Novel Food	Specifications		
	Chemical purity	≥ 99,5 %	HPLC [Eur. Ph. 2,2.29]
		≥ 99,0 %	1H-NMR
	Identification	Compliant with the structure	1H-NMR
		C: $47,14 \pm 0,4 \%$	Elemental analysis
		H: $6,59 \pm 0,4 \%$	
		N: $18,32 \pm 0,4 \%$	
	Total residual solvents	[Eur. Ph. 01/2008:50400]	Gas chromatography
	(methanol, ethyl acetate, isopropanol, ethanol)	< 1 000 ppm	[Eur. Ph. 01/2008:20424]
	Loss on drying	Internal standard < 0,5 %	[Eur. Ph. 01/2008:20232]
	Impurities	< 0,8 %	HPLC/GPC or 1H-NMR
	Heavy metals ^{b) c)}		
	Lead	< 3,0 ppm	ICP/AES
	Cadmium	< 1,0 ppm	(Pb, Cd)
	Mercury	< 0,1 ppm	Atomic fluorescence (Hg)
	Microbiological specifications ^{b)}		
	Total viable aerobic count (TVAC)	$\leq 1 \times 10^3 \text{ CFU/g}$	[Eur. Ph. 01/2011:50104]
	Total yeast and mould count (TYMC)	$\leq 1 \times 10^2 \text{ CFU/g}$	
	Escherichia coli	Absence in 1 g	

▼ <u>M9</u>		
	Authorised Novel Food	Specifications
Eur. Ph.: European Pharmacopoeia; 1H-NMR: proton nuclear magnetic resonance; HPLC: high-performance liquid chromatography; ICP/AES: Inductively coupled plasma atomic emission spectroscopy;		Eur. Ph.: European Pharmacopoeia; 1H-NMR: proton nuclear magnetic resonance; HPLC: high-performance liquid chromatography; GPC: gel permeation chromatography; ICP/AES: Inductively coupled plasma atomic emission spectroscopy;
		CFU: colony-forming units.
		a) Lit. $[\alpha]_D = (+) 126.6^\circ (c = 1, H_2O)$
		b) Analyses conducted on each batch
		c) Maximum levels in accordance with Regulation (EC) No 1881/2006
▼ <u>M50</u>		
	Extract of three herbal roots (Cynanchum wilfordii Hemsley, Phlomis umbrosa Turcz. and Angelica gigas Nakai)	Description/Definition The mixture of the three herbal roots is yellowish brown fine powder produced by hot-water extraction, concentration by evaporation, and spray drying Composition of the extract of mixture of the 3 herbal roots Cynanchum wilfordii root: 32,5 % (w/w) Phlomis umbrosa root: 32,5 % (w/w) Angelica gigas root: 35,0 % (w/w) Specifications Loss on drying: NMT 100 mg/g Assay Cinnamic acid: 0,012 – 0,039 mg/g Shanzhiside methyl ester: 0,20 – 1,55 mg/g Nodakenin: 3,35 – 10,61 mg/g Methoxsalen: < 3 mg/g Phenols: 13,0 – 40,0 mg/g

Coumarins: 13,0 - 40,0 mg/g Iridoids: 13,0 - 39,0 mg/g Saponins: 5,0 - 15,5 mg/g **Nutritive components**

Carbohydrates: 600 - 880 mg/g Proteins: 70 - 170 mg/g

Microbiological parameters

Total viable plate count: < 5000 CFU/g Total mold and yeast: < 100 CFU/g Coliform bacteria: < 10 CFU/g Salmonella: Negative/25 g

Fats: < 4 mg/g

▼<u>M50</u>

▼<u>M9</u>

Authorised Novel Food	Specifications
	Escherichia coli: Negative/25 g Staphylococcus aureus: Negative/25 g Heavy metals Lead: < 0,65 mg/kg Arsenic: < 3,0 mg/kg Mercury: < 0,1 mg/kg Cadmium: < 1,0 mg/kg CFU: Colony Forming Units
Ferric Sodium EDTA	Description/Definition: Ferric Sodium EDTA (ethylenediaminetetraacetic acid) is an odourless free-flowing, yellow to brown powder with a chemical purity of more than 99 % (w/w). It is freely soluble in water. Chemical formula: $C_{10}H_{12}FeN_2NaO_8*3H_2O$ Chemical characteristics: pH of 1 % solution: 3,5-5,5 Iron: 12,5-13,5 % Sodium: 5,5 % Water: 12,8 % Organic matter (CHNO): 68,4 % EDTA: 65,5-70,5 % Water insoluble matter: ≤ 0,1 % Nitrilo-triacetic acid: ≤ 0,1 %
Ferrous ammonium phosphate	Description/Definition: Ferrous ammonium phosphate is a grey/green fine powder, practically insoluble in water and soluble in dilute mineral acids. CAS No.: 10101-60-7 Chemical formula: FeNH₄PO₄ Chemical characteristics: pH of 5 % suspension in water: 6,8-7,8 Iron (total): ≥ 28 % Iron (II): 22-30 % (w/w) Iron (III): ≤ 7,0 % (w/w) Ammonia: 5-9 % (w/w) Water: ≤ 3,0 %

Authorised Novel Food	Specifications		
Fish peptides from Sardinops sagax	Description/Definition: The novel food ingredient is a peptide mixture, which is obtained by an alkaline protease-catalysed hydrolysis of fish (Sardinops sagax) muscle, subsequent isolation of the peptide fraction by column chromatography, concentration under vacuum and spray drying. Yellowish white powderPeptides (¹) (short chain peptides, dipeptides and tripeptides with a molecular weight of less than 2 kDa): ≥ 85 g/100 g Val-Tyr (dipeptide): 0,1-0,16 g/100 g Ash: ≤ 10 g/100 g Moisture: ≤ 8 g/100 g (¹) Kjeldahl method		
Flavonoids from Glycyrrhiza glabra	Description/Definition: Flavonoids derived from the roots or rootstock of <i>Glycyrrhiza glabra</i> L. are extracted with ethanol followed by further extraction of this ethanolic extract with medium-chain triglycerides. It is a dark-brown coloured liquid, containing 2,5 % to 3,5 % of glabridin. Moisture: < 0,5 % Ash: < 0,1 % Peroxide value (PV): < 0,5 meq/kg Glabridin: 2,5-3,5 % of fat Glycyrrhizinic acid: < 0,005 % Fat including polyphenol-type substances: ≥ 99 % Protein: < 0,1 % Carbohydrates: not detectable		
Fruit pulp, pulp juice, concentrated pulp juice from <i>Theobroma cacao</i> L. (Traditional food from a third country)	Description/Definition The traditional food is the fruit pulp from the cocoa (<i>Theobroma cacao</i> L) plant, which is the 'aqueous, mucilaginous and acidic substance in which the seed are embedded'. Cocoa fruit pulp is obtained by splitting cocoa pods followed by separation from husks and beans; the pulp is then subject to pasteurisation and freezing Cocoa pulp juice and/or cocoa concentrated pulp juice are produced following processing (enzymatic treatment, pasteurization, filtration, and concentration Typical compositional data of cocoa fruit pulp, pulp juice, concentrated pulp juice Protein (g/100 g): 0,0 to 2,0 Total fat (g/100 g): 0,0 to 0,2 Total sugars (g/100 g): > 11,0 Brix level (° Brix): ≥ 14 pH: 3,3 to 4,0 Microbiological criteria Total Plate Count (aerobic): < 10 000 cfu (°)/g Enterobacteriaceae: ≤ 10 cfu/g Salmonella: Absence in 25 g		

Authorised Novel Food	Specifications

▼<u>M71</u>

Frozen, dried and powder forms of Locusta migratoria (migratory locust)

Description/Definition:

The novel food consists of the frozen, dried and powder forms of migratory locust. The term 'migratory locust' refers to the adult of *Locusta migratoria*, an insect species that belongs to the Acrididae family (subfamily Locustinae).

The novel food is intended to be marketed in three different forms, namely: (i) thermally processed and frozen *L. migratoria* (LM frozen); (ii) thermally processed and freeze-dried *L. migratoria* (LM dried), and (iii) thermally processed freeze-dried and ground whole *L. migratoria* (whole LM powder). The LM dried may be marketed as such or in powder.

For LM frozen and LM dried, legs and wings must be removed to reduce the risk of intestinal constipation that could be possibly caused by ingestion of the large spines on the insect tibia. The whole LM powder is obtained via mechanical grinding of the insect with legs and wings, and sieving to reduce particle size below 1 mm.

A minimum 24 hours fasting period is required before killing the insects by freezing, to allow the adults to discard their bowel content.

Parameters	LM frozen	LM dried	Whole LM powder
Characteristics/Composition			
Ash (% w/w)	0,6-1,0	2,0-3,1	1,8-1,9
Moisture (% w/w)	67-73	≤ 5	≤ 5
Crude protein (N × 6,25) (% w/w)	11-21	43-53	50 - 60
Fat (% w/w)	7-13	31-41	31-41
Saturated fatty acids (% fat)	35-43	35-43	35-43
Digestible carbohydrates (% w/w)	0,1-2,0	0,1-2,0	1,0-3,5
(18) Dietary fibre (% w/w)	1,5-3,5	5,5-9,0	5,5-9,0
Chitin (% w/w)	1,7-2,4	6,4-10,4	10,5-13,9
Peroxide value (Meq O ₂ /kg fat)	≤ 5	≤ 5	≤ 5

Authorised Novel Food	Specifications				
	Contaminants				
	Lead (mg/kg)	≤ 0,07	≤ 0,07	≤ 0,07	
	Cadmium (mg/kg)	≤ 0,05	≤ 0,05	≤ 0,05	
	Aflatoxins (Sum of B1, B2, G1, G2) (µg/kg)	≤ 4	≤ 4	≤ 4	
	Aflatoxin B1 (μg/kg)	≤ 2	≤ 2	≤ 2	
	Deoxynivalenol (μg/kg)	≤ 200	≤ 200	≤ 200	
	Ochratoxin A (µg/kg)	≤ 1	≤ 1	≤ 1	
	Sum of dioxins and dioxins-like PCBs UB ((19) WHO ₂₀₀₅ PCDD/ F-PCB-TEQ) (pg/g fat)	≤ 1,2	≤ 1,2	≤ 1,2	
	Microbiological criteria				
	Total aerobic colony count ((7) CFU/g)	≤ 10 ⁵	≤ 10 ⁵	≤ 10 ⁵	
	Enterobacteriaceae (presumptive) (CFU/g)	≤ 100	≤ 100	≤ 100	
	Escherichia coli (CFU/g)	≤ 50	≤ 50	≤ 50	
	Listeria monocytogenes	Not detected in 25g	Not detected in 25g	Not detected in 25g	
	Salmonella spp.	Not detected in 25g	Not detected in 25g	Not detected in 25g	
	Bacillus cereus (presumptive) (CFU/g)	≤ 100	≤ 100	≤ 100	
	Coagulase positive Staphylococci (CFU/g	≤ 100	≤ 100	≤ 100	
	Sulfite-reducing Anaerobes (CFU/g)	≤ 30	≤ 30	≤ 30	
	Yeasts and moulds (CFU/g)	≤ 100	≤ 100	≤ 100	

Authorised Novel Food

Fucoidan: 60-65 %

Fucus vesiculosus	Description/Definition: Fucoidan from the seaweed <i>Fucus vesiculosus</i> is extracted using aqueous extraction in acidic solution and filtration processes without the use of organic solvents. The resulting extract is concentrated and dried to yield the fucoidan extract with the following specifications:		
	Off-white to brown powder Odour and Taste: Bland odour and taste Moisture: < 10 % (105 °C for 2 hours)		
	pH value: 4,0-7,0 (1 % suspension at 25 °C) Heavy metals: Arsenic (inorganic): < 1,0 ppm Cadmium: < 3,0 ppm		
	Lead: < 2,0 ppm		
	Mercury: < 1,0 ppm		
	Microbiological criteria:		
	Total aerobic microbial count: < 10 000 CFU/g		
	Yeast and mould count: < 100 CFU/g		
	Total enterobacteria count: Absence/g		
	Escherichia coli: Absence/g		
	Salmonella: Absence/10 g		
	Staphylococcus aureus: Absence/g		
	Composition of the two permitted types of extracts, based on the level of fucoidan:		
	Extract 1:		
	Fucoidan: 75-95 %		
	Alginate: 2,0-5,5 %		
	Polyphloroglucinol: 0,5-15 %		
	Mannitol: 1-5 %		
	Natural salts/Free Minerals: 0,5-2,5 %		
	Other carbohydrates: 0,5-1,0 %		
	Protein: 2,0-2,5 %		
	Extract 2:		

Specifications

Authorised Novel Food	Specifications	
	Alginate: 3,0-6,0 %	
	Polyphloroglucinol: 20-30 %	
	Mannitol: < 1,0 %	
	Natural salts/Free Minerals: 0,5-2,0 %	
	Other carbohydrates: 0,5-2,0 %	
	Protein: 2,0-2,5 %	
coidan extract from the seaweed	Description/Definition:	
ndaria pinnatifida		
	Fucoidan from seaweed <i>Undaria pinnatifida</i> is extracted using aqueous extraction in acidic solution and filtration processes without the use of organ solvents. The resulting extract is concentrated and dried to yield the fucoidan extract with the following specifications:	
	Off-white to brown powder	
	Odour and Taste: Bland odour and tasteMoisture: < 10 % (105 °C for 2 hours)	
	pH value: 4,0-7,0 (1 % suspension at 25 °C)	
	Heavy metals:	
	Arsenic (inorganic): < 1,0 ppm	
	Cadmium: < 3,0 ppm	
	Lead: < 2,0 ppm	
	Mercury: < 1,0 ppm	
	Microbiology:	
	Total aerobic microbial count: < 10 000 CFU/g	
	Yeast and mould count: < 100 CFU/g	
	Total enterobacteria count: Absence/g	
	Escherichia coli: Absence/g	
	Salmonella: Absence/10 g	
	Staphylococcus aureus: Absence/g	
	Composition of the two permitted types of extracts, based on the level of fucoidan:	
	Extract 1:	
	Fucoidan: 75-95 %	
	Alginate: 2,0-6,5 %	

Authorised Novel Food	Specifications
	Polyphloroglucinol: 0,5-3,0 %
	Mannitol: 1-10 %
	Natural salts/Free Minerals: 0,5-1,0 %
	Other carbohydrates: 0,5-2,0 %
	Protein: 2,0-2,5 %
	Extract 2:
	Fucoidan: 50-55 %
	Alginate: 2,0-4,0 %
	Polyphloroglucinol: 1,0-3,0 %
	Mannitol: 25-35 %
	Natural salts/Free Minerals: 8-10 %
	Other carbohydrates: 0,5-2,0 %
	Protein: 1,0-1,5 %
2'-Fucosyllactose	Definition:
synthetic)	Chemical name: α -L-Fucopyranosyl- $(1\rightarrow 2)$ - β -D-galactopyranosyl- $(1\rightarrow 4)$ - D-glucopyranose
	Chemical formula: C ₁₈ H ₃₂ O ₁₅
	CAS No: 41263-94-9
	Molecular weight: 488,44 g/mol
	Description:
	2'-fucosyllactose is a white to off-white powder that is produced by a chemical synthesis process.
	Purity:
	2'-Fucosyllactose: ≥ 95 %
	D-Lactose: $\leq 1.0 \text{ w/w } \%$
	L-Fucose: $\leq 1,0$ w/w %
	Difucosyl- D-lactose isomers: ≤ 1,0 w/w %
	2'-Fucosyl- D-lactulose: ≤ 0,6 w/w %
	pH (20 °C, 5 % solution): 3,2-7,0
	Water (%): ≤ 9,0 %
	Ash, sulphated: $\leq 0.2\%$

Authorised Novel Food	Acetic acid: ≤ 0,3 % Residual solvents (methanol, 2-propanol, methyl acetate, acetone): ≤ 50,0 mg/kg singly, ≤ 200,0 mg/kg in combination Residual proteins: ≤ 0,01 % Heavy Metals: Palladium: ≤ 0,1 mg/kg Nickel: ≤ 3,0 mg/kg Microbiological criteria: Aerobic mesophilic bacteria total count: ≤ 500 CFU/g Yeasts and Moulds: ≤ 10 CFU/g		
2'-Fucosyllactose (microbial source)	Residual endotoxins: ≤ 10 EU/mg M27 Definition: Chemical name: α-L-Fucopyranosyl-(1→2)-β-D-galactopyranosyl-(1→4)-D-glucopyranose Chemical formula: C ₁₈ H ₃₂ O ₁₅ CAS No: 41263-94-9 Molecular weight: 488,44 g/mol Source: Source:		
	Genetically modified strain of <i>Escherichia coli</i> K-12 Description: 2'-Fucosyllactose is a white to off-white powder that is produced by a microbial process. Purity: 2'-Fucosyllactose: ≥ 83 % D-Lactose: ≤ 10,0 % L-Fucose: ≤ 2,0 % Difucosyl-D-lactose: ≤ 5,0 % 2'-Fucosyl-D-lactulose: ≤ 1,5 % Sum of saccharides (2'-Fucosyllactose, D-Lactose, L-Fucose, Difucosyl-D-lactose, 2'-Fucosyl-D-lactulose): ≥ 90 % pH (20 C, 5 % solution): 3,0-7,5 Water: ≤ 9,0 %	Genetically modified strain of <i>Escherichia coli</i> BL21 Description: 2'-Fucosyllactose is a white to off white powder and the liquid concentrate (45 % ± 5 % w/v) aqueous solution is a colourless to slight yellow clear aqueous solution. 2'-Fucosyllactose is produced by a microbiological process. Purity: 2'-Fucosyllactose: ≥ 90 % Lactose: ≤ 5,0 % Fucose: ≤ 3,0 % 3-Fucosyllactose: ≤ 5,0 % Fucosylgalactose: ≤ 5,0 % Difucosyllactose: ≤ 5,0 %	

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Authorised Novel Food		pecifications
	Sulphated ash: ≤ 2,0 % Acetic acid: ≤ 1,0 % Residual proteins: ≤ 0,01 % Microbiological criteria: Aerobic mesophilic bacteria total count: ≤ 3 000 CFU/g Yeasts: ≤ 100 CFU/g Moulds: ≤ 100 CFU/g Endotoxins: ≤ 10 EU/mg	Glucose: ≤ 3,0 % Galactose: ≤ 3,0 % Water: ≤ 9,0 % (powder) Ash, sulphated: ≤ 0,5 % (powder and liquid) Residual proteins: ≤ 0,01 % (powder and liquid) Heavy Metals: Lead: ≤ 0,02 mg/kg (powder and liquid) Arsenic: ≤ 0,2 mg/kg (powder and liquid) Cadmium: ≤ 0,1 mg/kg (powder and liquid) Microbiological criteria: Total plate count: ≤ 10 ⁴ CFU/g (powder), ≤ 5 000 CFU/g (liquid Yeasts and Moulds: ≤ 100 CFU/g (powder); ≤ 50 CFU/g (liquid Enterobacteriaceae/Coliforms: absence in 11 g (powder and liquid) Salmonella: negative/100 g (powder), negative/200 ml (liquid) Cronobacter: negative/100 g (powder), negative/200 ml (liquid) Cronobacter: 100 EU/g (powder), ≤ 100 EU/ml (liquid) Aflatoxin M1: ≤ 0,025 μg/kg (powder and liquid) ◀
2'-Fucosyllactose/Difucosyllactose mixture ('2'-FL/DFL') (microbial source)		te powder or agglomerates thereof that is produced by a microbial process.
(microbiai source)	Source: Genetically modified <i>Escherichia coli</i> strain K-12 DH1 Characteristics/Composition: Appearance: White to off white powder or agglomerates Sum of 2'-Fucosyllactose, Difucosyllactose, D-Lactose, L-Fucose, and 3-	Fucosyllactose (% of dry matter): $> 92.0\%$ (w/w)

Sum of 2'-Fucosyllactose and Difucosyllactose (% of dry matter): \geq 85,0 % (w/w)

▼<u>M56</u>

Authorised Novel Food	Specifications
	2'-Fucosyllactose (% of dry matter): ≥ 75,0 % (w/w) Difucosyllactose (% of dry matter): ≥ 5,0 % (w/w) D-Lactose: ≤ 10,0 % (w/w) L-Fucose: ≤ 1,0 % (w/w) 2'-Fucosyl-D-lactulose: ≤ 2,0 (w/w) Sum of other carbohydrates (¹¹): ≤ 6,0 % (w/w) Moisture: ≤ 6,0 % (w/w) Ash, sulfated: ≤ 0,8 % (w/w) pH (20 °C, 5 % solution): 4,0 -6,0 Residual protein: ≤ 0,01 % (w/w) Microbiological criteria: Aerobic mesophilic total plate count: ≤ 1000 CFU/g Enterobacteriaceae: ≤ 10 CFU/g Salmonella sp.: Negative/25 g Yeast: ≤ 100 CFU/g Mould: ≤ 100 CFU/g
	Residual endotoxins: ≤ 10 EU/mg
	CFU: Colony Forming Units; EU: Endotoxin Units
▼ <u>M72</u>	
3-Fucosyllactose ('3-FL') (microbial source)	Description: 3-Fucosyllactose (3-FL) is a purified, white to off-white powder that is produced by microbial fermentation and contains limited levels of D-Lactose, L-Fucose, D-Galactose, and D-Glucose. Source: Genetically modified strain of Escherichia coli K-12. Definition: Chemical formula: C ₁₈ H ₃₂ O ₁₅ Chemical name: β-D-galactopyranosyl-(1→4)[-α-L-fucopyranosyl-(1→3)]-D-glucopyranose Molecular mass: 488,44 Da CAS No 41312-47-4 Characteristics/Composition: 3-Fucosyllactose (% of dry matter): ≤ 90,0 % (w/w) D-Lactose (% of dry matter): ≤ 5,0 % (w/w) L-Fucose (% of dry matter): ≤ 3,0 % (w/w)

	Authorised Novel Food	Specifications
		Sum of D-Galactose/D-Glucose (% of dry matter): ≤ 3.0 % (w/w) Sum of other carbohydrates ^a (% of dry matter): ≤ 3.0 % (w/w) Moisture: ≤ 5.0 % (w/w) pH (20 °C, 5 % solution): 3,0-7,5 Residual protein: ≤ 0.01 % (w/w) Ash (%): ≤ 0.5
		Heavy metals/Contaminants:
		Arsenic: $\le 0.2 \text{ mg/kg}$ Cadmium: $\le 0.05 \text{ mg/kg}$ Lead: $\le 0.05 \text{ mg/kg}$ Mercury: $\le 0.1 \text{ mg/kg}$ Aflatoxin M1: $\le 0.025 \text{ µg/kg}$ Aflatoxin B1: $\le 0.1 \text{ µg/kg}$ Residual endotoxins: $\le 0.3 \text{ EU/mg}$
		Microbiological criteria:
		Total plate count: ≤ 1 000 CFU/g
		Enterobacteriaceae: Absence in 10 g
		Salmonella sp.: Absence in 25 g
		Cronobacter (Enterobacter) sakazakii: Absence in 10 g
		Listeria monocytogenes: Absence in 25 g
		Bacillus cereus: ≤ 10 CFU/g
		Yeast: $\leq 100 \text{ CFU/g}$
		Mould: $\leq 100 \text{ CFU/g}$
		CFU: Colony Forming Units; EU: Endotoxin Units; ^a Sum of other carbohydrates: 3-Fucosyllactose isomer, difucosyllactose isomer, and oligomers.
▼ <u>M9</u>		
	Galacto-oligosaccharide	Description/Definition:
		Galacto-oligosaccharide is produced from milk lactose by an enzymatic process using β-galactosidases from Aspergillus oryzae, Bifidobacterium bifidum, Pichia pastoris, Sporobolomyces singularis, Kluyveromyces lactis, Bacillus circulans, and Papiliotrema terrestris.
		GOS: min 46 % Dry Matter (DM)
		Lactose: max 40 % DM
		Glucose: max 27 % DM
		1

Authorised Novel Food	Specifications
	Galactose: min 0,8 % DM Ash: max 4,0 % DM Protein: max 4,5 % DM Nitrite: max. 2 mg/kg
Glucosamine HCl from Aspergillus niger and genetically modified strain of E. coli K-12	White crystalline odourless powder Molecular formula: C ₆ H ₁₃ NO ₅ · HCl Relative molecular mass: 215,63 g/mol D-Glucosamine HCl 98,0-102,0 % of reference standard (HPLC) Specific rotation + 70,0° - + 73,0°
Glucosamine sulphate KCl from Aspergillus niger and genetically modified strain of E. coli K-12	White crystalline odourless powder Molecular formula: (C ₆ H ₁₄ NO ₅) ₂ SO ₄ · 2KCl Relative molecular mass: 605,52 g/mol D-Glucosamine Sulphate 2KCl 98,0-102,0 % of reference standard (HPLC) Specific Rotation +50,0° to +52,0°
Glucosamine sulphate NaCl from Aspergillus niger and genetically modified strain of E. coli K-12	White crystalline odourless powder Molecular formula: (C ₆ H ₁₄ NO ₅) ₂ SO ₄ · 2NaCl Relative molecular mass: 573,31 g/mol D-Glucosamine HCl: 98-102 % of reference standard (HPLC) Specific Optical Rotation: +52° - +54°
Guar Gum	Description/Definition: Native guar gum is the ground endosperm of seeds from natural strains of guar Cyamopsis tetragonolobus L. Taub. (Leguminosae family). It consists of a high molecular weight polysaccharide, primarily composed of galactopyranose and mannopyranose units combined through glycosidic linkages, which may be described chemically as a galactomannan (galactomannan content not less than 75 %). Appearance: White to yellowish powder Molecular weight: Between 50 000 – 8 000 000 Daltons CAS number: 9000-30-0 Einecs Number: 232-536-8 Purity: As specified by Commission Regulation (EU) No 231/2012 laying down specifications for food additives listed in Annexes II and III to Regulation (EC) No 1333/2008 of the European Parliament and of the Council (¹) & by Commission Implementing Regulation (EU) 2015/175 of 5 February 2015 laying down special conditions applicable to the import of guar gum originating in or consigned from India due to contamination risks by pentachlorophenol and dioxins (²).

Authorised Novel Food	Specifications
	Physico-chemical properties:
	Powder
	Shelf-life: 2 years
	Colour: White
	Odour: Light
	Average diameter of particles: 60-70µm
	Moisture: Max 15 %
	Viscosity * at 1 hour —
	Viscosity * at 2 hours: Min 3 600 mPa.s
	Viscosity * at 24 hours: Min 4 000 mPa.s
	Solubility: Soluble in hot and cold water
	pH for 10g/L, at 25 °C - 6-7,5
	Flakes
	Useful life: 1 year
	Colour: White/off white with absence or minimal presence of black spots
	Odour: Light
	Average diameter of particles: 1-10 mm
	Moisture: Max 15 %
	Viscosity * at 1 hour: Min 3 000 mPa.s
	Viscosity * at 2 hours —
	Viscosity * at 24 hours —
	Solubility — Soluble in hot and cold water
	pH for 10g/L, at 25 °C - 5-7,5
	(*) The measurements of viscosity are carried out under the following conditions: 1 %, 25 °C, 20 rpm
Heat-treated milk products	Description/Definition:
fermented with Bacteroides xylani- solvens	Heat-treated fermented milk products are produced with <i>Bacteroides xylanisolvens</i> (DSM 23964) as starter culture.

Authorised Novel Food	Specifications
	Semi-skimmed milk (between 1,5 % and 1,8 % fat) or skimmed milk (0,5 % fat or less) is pasteurised or ultra-heat-treated before starting the fermentation wi <i>Bacteroides xylanisolvens</i> (DSM 23964). The resulting fermented milk product is homogenised and then heat-treated to inactivate <i>Bacteroides xylanisolvens</i> (DSI 23964). The final product does not contain viable cells of <i>Bacteroides xylanisolvens</i> (DSM 23964)(¹). (¹) Modified DIN EN ISO 21528-2.
Hydroxytyrosol	Description/Definition:
Hydroxytyrosor	Hydroxytyrosol is a pale yellow viscous liquid obtained by chemical synthesis
	Molecular formula: C ₈ H ₁₀ O ₃
	Molecular weight: 154,6 g/mol
	CAS No: 10597-60-1
	Moisture $\leq 0.4 \%$
	Odour: CharacteristicTaste: Slightly bitter
	Solubility (water): Miscible with water
	pH: 3,5-4,5
	Refractive Index: 1,571-1,575
	Purity:
	Hydroxytyrosol: ≥ 99 %
	Acetic acid: $\leq 0.4\%$
	Hydroxytyrosol acetate: ≤ 0,3 %
	Sum of homovanillic acid, iso-homovanilic acid, and 3-methoxy-4hydroxyphenylglycol: $\leq 0.3\%$
	Heavy Metals
	Lead: $\leq 0.03 \text{ mg/kg}$
	Cadmium: $\leq 0.01 \text{ mg/kg}$
	Mercury: $\leq 0.01 \text{ mg/kg}$
	Residual Solvents
	Ethyl acetate: $\leq 25,0$ mg/kg
	Isopropanol: ≤ 2,50 mg/kg
	Methanol: $\leq 2,00 \text{ mg/kg}$
	Tetrahydrofuran: $\leq 0.01 \text{ mg/kg}$

Authorised Novel Food

	Ice Structuring Protein type III	Description/Definition:
	HPLC 12	The Ice Structuring Protein (ISP) preparation is a light-brown liquid produced by submerged fermentation of a genetically-modified strain of food-grade baker's yeast (<i>Saccharomyces cerevisiae</i>) in which a synthetic gene for the ISP has been inserted into the yeast's genome. The protein is expressed and secreted into the growth medium where it is separated from the yeast cells by micro-filtration and concentrated by ultra-filtration. As a result, the yeast cells are not transferred into the ISP preparation as such or under an altered form. The ISP preparation consists of native ISP, glycosylated ISP and proteins and peptides from the yeast and sugars as well as acids and salts commonly found in food. The concentrate is stabilised with 10 mM citric acid buffer.
		Assay: ≥ 5 g/l active ISP
		pH: 2,5-3,5
		Ash: $\leq 2,0 \%$
		DNA: Not detectable
	Aqueous extract of dried leaves of	Description/Definition:
	Ilex guayusa	Dark brown liquid. Aqueous extracts of dried leaves of <i>Ilex guayusa</i> .
		Composition:
		Protein: < 0,1 g/100 ml
		Fat: $< 0.1 \text{ g/}100 \text{ ml}$
		Carbohydrate: 0,2–0,3 g/100 ml
		Total sugars: < 0,2 g/100 ml
		Caffeine: 19,8–57,7 mg/100 ml
		Theobromine: 0,14–2,0 mg/100 ml
		Chlorogenic acids: 9,9–72,4 mg/100ml
▼ <u>M47</u>		
	Infusion from coffee leaves of	Description/Definition:
	Coffee arabica L. and/or Coffee	The traditional food consists of an infusion of leaves from Coffea arabica L. and/or Coffea canephora Pierre ex A.Froehner (family: Rubiaceae).
	canephora Pierre ex A. Froehner	The traditional food is prepared by mixing a maximum of 20 g of dried leaves from Coffea arabica L. and/or Coffea canephora Pierre ex A.Froehner with 1 L
	(Traditional food from a third country)	of hot water. Leaves are removed and the infusion is then subjected to pasteurization (at least 71 °C for 15 seconds).

Specifications

▼<u>M47</u>

Authorised Novel Food	Specifications
	Composition:
	Visual: Brown green liquid
	Odour and taste: Characteristic
	Chlorogenic acid (5-CQA): < 100 mg/L
	Caffeine: < 80 mg/L
	Epigallocatechin gallate (EGCG): < 700 mg/L
	Microbiological criteria:
	Total plate count: < 500 CFU/g
	Total yeast and mould count: < 100 CFU/g
	Total coliforms: < 100 CFU/g
	Escherichia coli: Absence in 1 g
	Salmonella: Absence in 25 g
	Heavy metals:
	Lead (Pb): < 3,0 mg/L
	Arsenic (As): < 2,0 mg/L
	Cadmium (Cd): < 1,0 mg/L
	CFU: Colony Forming Units
Isomalto-oligosaccharide	Powder:
	Solubility (water) (%): > 99
	Glucose (% dry basis): ≤ 5.0
	Isomaltose + DP3 to DP9 (% dry basis): ≥ 90
	Moisture (%): ≤ 4.0
	Sulphated $ash(g/100 g)$: ≤ 0.3
	Heavy metals:
	Lead (mg/kg): ≤ 0.5
	Arsenic (mg/kg): ≤ 0.5

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Authorised Novel Food	Specifications
	Syrup:
	Dried solids (g/100 g): > 75
	Glucose (% dry basis): ≤ 5.0
	Isomaltose + DP3 to DP9 (% dry basis): ≥ 90
	pH: 4 - 6
	Sulphated $ash(g/100 g)$: ≤ 0.3
	Heavy metals:
	Lead (mg/kg): ≤ 0.5
	Arsenic (mg/kg): ≤ 0.5

Isomaltulose

Description/Definition:

A reducing disaccharide that consists of one glucose and one fructose moiety linked by an alpha-1,6-glycosidic bond. It is obtained from sucrose by an enzymatic process. The commercial product is the monohydrate. Appearance: Virtually odourless, white or almost white crystals with a sweet taste

Chemical name: 6-O-α-D-glucopyranosyl-D-fructofuranose, monohydrate

CAS No.: 13718-94-0

Chemical formula: C₁₂H₂₂O₁₁ · H₂O

Structural formula

Formula weight: 360,3 (monohydrate)

Authorised Novel Food	Specifications
	Purity:
	Assay: $\geq 98\%$ on the dry basis
	Loss on drying: $\leq 6.5 \%$ (60 °C, 5 hours)
	Heavy metals:
	Lead: ≤ 0,1 mg/kg Determine using an atomic absorption technique appropriate to the specified level. The selection of sample size and method of sample preparation may be based on the principles of the method described in FNP 5(¹), 'Instrumental methods'
	(1) Food and Nutrition Paper 5 Rev. 2 — Guide to specifications for general notices, general analytical techniques, identification tests, test solutions and other reference materials (JECFA), 1991, 322 pp., English, ISBN 92-5-102991-1.
Lactitol	Description/Definition:
	Crystalline powder or colourless solution manufactured via catalytic hydrogenation of lactose. Crystalline products occur in anhydrous, monohydrate and dihydrate forms. Nickel is used as a catalyst.
	Chemical name: 4-O-β-D-Galactopyranosyl-D-glucitol
	Chemical formula: C ₁₂ H ₂₄ O ₁₁
	Molecular weight: 344,31 g/mol
	CAS No: 585-86-4
	Purity:
	Solubility (in water): Very soluble in water
	Specific rotation $\left[\alpha\right]_{D}^{20} = +13^{\circ} \text{ to } +16^{\circ}$
	Assay: ≥ 95 % d.b (d.b — expressed on the dry weight basis)
	Water: ≤ 10,5 %
	Other polyols: ≤ 2,5 % d.b
	Reducing sugars: ≤ 0,2 % d.b
	Chlorides: ≤ 100 mg/kg d.b
	Sulphates: ≤ 200 mg/kg d.b
	Sulphated ash: ≤ 0,1 % d.b
	Nickel: ≤ 2.0 mg/kg d.b
	Arsenic: ≤ 3,0 mg/kg d.b
	Lead: ≤ 1,0 mg/kg d.b

Authorised Novel Food	Specifications
Lacto-N-neotetraose	Definition:
(synthetic)	Chemical name: β-D-Galactopyranosyl-(1→4)-2-acetamido-2-deoxy-β-D-glucopyranosyl-(1→3)-β-D-galactopyranosyl-(1→4)- D-glucopyranose
	Chemical formula: C ₂₆ H ₄₅ NO ₂₁
	CAS No: 13007-32-4
	Molecular weight: 707,63 g/mol
	Description:
	Lacto-N-neotetraose is a white to off-white powder. Produced by a chemical synthesis process and is isolated by crystallisation.
	Purity:
	Assay (water free): ≥ 96 %
	D-Lactose: ≤ 1,0 %
	Lacto-N-triose II: ≤ 0,3 %
	Lacto-N-neotetraose fructose isomer: ≤ 0,6 %
	pH (20 °C, 5 % solution): 5,0-7,0
	Water: ≤ 9,0 %
	Ash, sulphated: ≤ 0,4 %
	Acetic acid: ≤ 0,3 %Residual solvents (methanol, 2-propanol, methyl acetate, acetone): ≤ 50 mg/kg singly, ≤ 200 mg/kg in combination
	Residual proteins: ≤ 0,01 %
	Palladium: ≤ 0,1 mg/kg
	Nickel: ≤ 3,0 mg/kg
	Microbiological criteria:
	Aerobic mesophilic bacteria total count: ≤ 500 CFU/g
	Yeasts: ≤ 10 CFU/g
	Moulds: $\leq 10 \text{ CFU/g}$
	Residual endotoxins: ≤ 10 EU/mg
Lacto-N-neotetraose	Definition:
(microbial source)	Chemical name: β-D-Galactopyranosyl-(1→4)-2-acetamido-2-deoxy-β-D-glucopyranosyl-(1→3)-β-D-galactopyranosyl-(1→4)-D-glucopyranose
	Chemical formula: C ₂₆ H ₄₅ NO ₂₁
	CAS No: 13007-32-4
	Molecular weight: 707,63 g/mol

Authorised Novel Food	Specifications
	Source:
	— Genetically modified strain of Escherichia coli K-12, or
	— a combination of the genetically modified strains PS-LNnT-JBT and DS-LNnT-JBT of Escherichia coli BL21(DE3)
	Description:
	Lacto-N-neotetraose is a white to off-white powder that is produced by a microbiological process.
	Purity:
	Assay (water free): $\geq 80 \%$
	D-Lactose: ≤ 10,0 %
	Lacto- <i>N</i> -triose II: ≤ 3,0 %
	para-Lacto-N-neohexaose: ≤ 5,0 %
	Lacto- <i>N</i> -neotetraose fructose isomer: ≤ 1,0 %
	Sum of saccharides (Lacto-N-neotetraose, D-Lactose, Lacto-N-triose II, para-Lacto-N-neohexaose, Lacto-N-neotetraose fructose isomer): ≥ 92 % (% w/w dr matter)
	pH (20 °C, 5 % solution): 4,0-7,0
	Water: ≤ 9,0 %
	Ash, sulphated: ≤ 1,0 %
	Residual solvents (methanol): ≤ 100 mg/kg
	Residual proteins: $\leq 0.01 \%$
	Microbiological criteria:
	Aerobic mesophilic bacteria total count: ≤ 500 CFU/g
	Yeasts and moulds: ≤ 50 CFU/g
	Residual endotoxins: ≤ 10 EU/mg
	CFU: Colony Forming Units; EU: Endotoxin Units
<u>M43</u>	
<u>M44</u>	
Lacto-N-tetraose ('LNT')	Definition:
(microbial source)	Chemical formula: C ₂₆ H ₄₅ NO ₂₁ Chemical name: β-D-Galactopyranosyl-(1→3)-2-acetamido-2-deoxy-β-D-glucopyranosyl-(1→3)-β-D-galactopyranosyl-(1→4)-D-glucopyranose Molecular mass: 707.63 Da CAS No 14116-68-8
	Description:
	Lacto- <i>N</i> -tetraose is a purified, white to off-white amorphous powder or agglomerates that is produced by a microbial process.
	Lacto-11-teraose is a purifica, white to off-white amorphous powder of agglomerates that is produced by a inicioulal process.

Source:
Genetically modified strain of Escherichia coli strain K-12 DH1
Characteristics/Composition:
Appearance: White to off white powder or agglomerates
Sum of lacto-N-tetraose, D-Lactose and lacto-N-triose II (% of dry matter): ≥ 90.0 % (w/w)
Lacto-N-tetraose (% of dry matter): ≥ 70.0 % (w/w)
D-Lactose: $\leq 12.0 \% (w/w)$
Lacto- N -triose II: $\leq 10.0 \%$ (w/w)
$Para$ -lacto- N -hexaose- $2: \le 3.5 \% (w/w)$
Lacto- <i>N</i> -tetraose fructose isomer: ≤ 1.0 % (w/w)
Sum of other carbohydrates: $\leq 5.0 \%$ (w/w)
Moisture: $\leq 6.0 \%$ (w/w)
Ash, sulfated: $\leq 0.5 \%$ (w/w)
pH (20 °C, 5 % solution): 4.0 -6.0
Residual protein: $\leq 0.01 \% \text{ (w/w)}$
Microbiological criteria:
Aerobic mesophilic bacteria total plate count: ≤ 1 000 CFU/g
Enterobacteriaceae: ≤ 10 CFU/g
Salmonella spp.: Negative/25 g
$Yeast: \leq 100 \text{ CFU/g}$
Mould: \leq 100 CFU/g
Residual endotoxins: ≤ 10 EU/mg
CFU: Colony Forming Units
Description/Definition:
The traditional food are fresh and frozen berries from <i>Lonicera caerulea</i> var. edulis.
Lonicera caerulea L. is a deciduous shrub belonging to the Caprifoliaceae family.
Typical nutritional components of haskap berries (given in fresh berries):
Carbohydrates: 12,8 %
Fibre: 2,1 %
Lipids: 0,6 %
Proteins: 0,7 %

▼<u>M9</u>

Authorised Novel Food	Specifications
	Ash: 0,4 % Water: 85,5 %
Lucerne leaf extract from	Description/Definition:
Medicago sativa	The Lucerne (<i>Medicago sativa</i> L.) is processed within 2 hours after harvest. It is chopped and crushed. By passing through an oleaginous-type press, the Lucerne provides a fibrous residue and press juice (10 % of dry matter). The dry matter of this juice contains about 35 % of crude protein. The press juice (pH 5,8-6,2) is neutralised. Preheating and vapour injection allows coagulation of proteins associated with carotenoid and chlorophyll pigments. The protein precipitate is separated by centrifugation and thereafter dried. After adding ascorbic acid the Lucerne protein concentrate is granulated and stored in inert gas or in cold storage.
	Composition:
	Protein: 45-60 %
	Fat: 9-11 %
	Free carbohydrates (soluble fibre): 1-2 %
	Polysaccharides (insoluble fibre): 11-15 %
	including cellulose: 2-3 %
	Minerals: 8-13 %
	Saponins: ≤ 1,4 %
	Isoflavones: ≤ 350 mg/kg
	Coumestrol: ≤ 100 mg/kg
	Phytates: ≤ 200 mg/kg
	L-canavanine: ≤ 4,5 mg/kg
Lycopene	Description/Definition:
	Synthetic lycopene is produced by the Wittig condensation of synthetic intermediates commonly used in the production of other carotenoids used in food. Synthetic lycopene consists of ≥ 96 % lycopene and minor quantities of other related carotenoid components. Lycopene is presented either as a powder in a suitable matrix or an oily dispersion. The colour is dark red or red-violet. Antioxidative protection has to be assured.
	Chemical name: Lycopene
	CAS No.: 502-65-8 (all-trans lycopene)
	Chemical formula: C ₄₀ H ₅₆
	Formula weight: 536,85 Da

Authorised Novel Food	Specifications
Lycopene from Blakeslea trispora	Description/Definition:
	The purified lycopene from <i>Blakeslea trispora</i> consists of ≥ 95 % lycopene and ≤ 5 % other carotenoids. It is presented either as a powder in a suitable matrix or an oily dispersion. The colour is dark red or red-violet. Anti-oxidative protection has to be assured.
	Chemical name: Lycopene
	CAS No.: 502-65-8 (all trans lycopene)
	Chemical formula: C ₄₀ H ₅₆
	Formula weight: 536,85 Da
Lycopene from tomatoes	Description/Definition:
Lycopolic from tollinoos	The purified lycopene from tomatoes (<i>Lycopersicon esculantum</i> L.) consists of ≥ 95 % lycopene and ≤ 5 % other carotenoids. It is presented either as a powder in a suitable matrix or an oily dispersion. The colour is dark red or red-violet. Anti-oxidative protection has to be assured.
	Chemical name: Lycopene
	CAS No.: 502-65-8 (all trans lycopene)
	Chemical formula: C ₄₀ H ₅₆
	Formula weight: 536,85 Da
Lycopene oleoresin from tomatoes	Description/Definition:
	Lycopene oleoresin from tomatoes is obtained by solvent extraction of ripe tomatoes (<i>Lycopersicon esculentum Mill.</i>) with subsequent removal of the solvent. It is a red to dark brown viscous, clear liquid.
	Total lycopene: 5-15 %
	Thereof trans-lycopene: 90-95 %
	Total carotenoids (calculated as lycopene): 6,5-16,5 %
	Other carotenoids: 1,75 %
	(Phytoene/phytofluene/β-carotene): (0,5-0,75/0,4-0,65/0,2-0,35 %)
	Total tocopherols: 1,5-3,0 %
	Unsaponifiable matter: 13-20 %
	Total fatty acids: 60-75 %
	Water (Karl Fischer): ≤ 0,5 %

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Hen egg white lysozyme hydrolysate

Authorised Novel Food

Description/Definition

Hen egg white lysozyme hydrolysate is obtained from hen egg white lysozyme by an enzymatic process, using subtilisin from *Bacillus licheniformis* The product is a white to light yellow powder.

Specifications

Specification

Protein (TN(*) x 5,30): 80-90 %

Tryptophan: 5-7 %

Ratio Tryptophan/LNAA(**): 0,18-0.25

Degree of hydrolysis: 19-25 %

Moisture: < 5 %
Ash: < 10 %
Sodium: < 6 %

Heavy metals

Arsenic: < 1 ppm Lead: < 1 ppm Cadmium: < 0,5 ppm

Mercury: < 0,1 ppm

Microbiological criteriaTotal aerobic count: $< 10^3$ CFU/g

Total combined yeasts/moulds count: < 10² CFU/g

Enterobacteria: < 10 CFU/g

Salmonella spp: Absence in 25 g Escherichia coli: Absence in 10 g

Staphylococcus aureus: Absence in 10 g Pseudomonas aeruginosa: Absence in 10 g

* TN: total nitrogen

** LNAA: large neutral amino acids

Authorised Novel Food

Magnesium citrate malate	Description/Definition:
	Magnesium citrate malate is a white to yellowish-white, amorphous powder. Chemical formula: Mg ₅ (C ₆ H ₅ O ₇) ₂ (C ₄ H ₄ O ₅) ₂
	Chemical name: Pentamagnesium di-(2-hydroxybutanedioate)-di-(2- hydroxypropane-1,2,3-tricarboxylate)
	CAS No.: 1259381-40-2
	Molecular weight: 763,99 Daltons (anhydrous)
	Solubility: Freely soluble in water (about 20 g in 100 ml)
	Description of the physical state: Amorphous powder
	Assay magnesium: 12,0-15,0 %
	Loss on drying (120 °C/4 hours): ≤ 15 %
	Colour (solid): White to yellowish-white
	Colour (20 % aqueous solution): Colourless to yellowish
	Appearance (20 % aqueous solution): Clear solution
	pH (20 % aqueous solution): Approx. 6,0
	Impurities:
	Chloride: ≤ 0,05 %
	Sulphate: ≤ 0,05 %
	Arsenic: ≤ 3.0 ppm
	Lead: ≤ 2.0 ppm
	Cadmium: ≤ 1 ppm
	Mercury: ≤ 0.1 ppm
Magnolia Bark Extract	Description/Definition:
	Magnolia bark extract is obtained from the bark of the plant <i>Magnolia officinalis</i> L. and produced with supercritical carbon dioxide. The bark is washed and oven dried to reduce moisture content before being crushed and extracted with supercritical carbon dioxide. The extract is dissolved in medical-grade ethanol and re-crystallised to yield magnolia bark extract.
	Magnolia bark extract is mainly composed of two phenolic compounds, magnolol and honokiol.
	Appearance: Light brownish powder
	Purity:
	Magnolol: ≥ 85,2 %
	Honokiol: $\geq 0.5\%$

Specifications

Authorised Novel Food	Specifications
	Magnolol & Honokiol: ≥ 94 %
	Total Eudesmol: $\leq 2\%$
	Moisture: 0,50 %
	Heavy metals:
	Arsenic (ppm): ≤ 0.5
	Lead (ppm): ≤ 0.5
	Methyl eugenol (ppm): ≤ 10
	Tubocurarine (ppm): ≤ 2.0
	Total Alkaloid (ppm): ≤ 2.00
	Total Alkalold (ppill). \$\log 100
Taize-germ oil high in unsapo-	Description/Definition:
ifiable matter	Maize-germ oil high in unsaponifiable matter is produced by vacuum distillation and it is different from refined maize-germ oil in the concentration of the unsaponifiable fraction (1,2 g in refined maize-germ oil and 10 g in 'maize-germ oil high in unsaponifiable matter').
	Purity:
	Unsaponifiable matter: > 9,0 g/100 g
	Tocopherols: $\geq 1.3 \text{ g/}100 \text{ g}$
	α-tocopherol (%): 10-25 %
	β-tocopherol (%): < 3,0 %
	γ-tocopherol (%): 68-89 %
	δ-tocopherol (%): < 7,0 %
	Sterols, triterpenic alcohols, methylsterols: > 6,5 g/100 g
	Fatty acids in triglycerides:
	palmitic acid: 10,0-20,0 %
	stearic acid: < 3,3 %
	oleic acid: 20,0-42,2 %
	linoleic acid: 34,0-65,6 %
	linolenic acid: < 2,0 %
	Acid value: ≤ 6,0 mg KOH/g
	Peroxide value (PV): $\leq 10 \text{ mEq } O_2/\text{kg}$

Authorised Novel Food	Specifications
	Heavy metals:
	Iron (Fe): < 1 500 μg/kg
	Copper (Cu): < 100 µg/kg
	Impurities:
	Polycyclic aromatic hydrocarbons (PAH) Benzo(a)pyrene: < 2 μg/kg
	Treatment with active carbon is required to ensure that polycyclic aromatic hydrocarbons (PAH) are not enriched in the production of 'maize-germ oil high unsaponifiable matter'
lethylcellulose	Description/Definition:
tettiyicenulose	Methyl cellulose is cellulose obtained directly from natural strains of fibrous plant material and partially etherified with methyl groups.
	Chemical name: Methyl ether of cellulose
	Chemical formula: The polymers contain substituted anhydroglucose units with the following general formula:
	C6H7O2(OR1)(OR2)(OR3) where R1, R2, R3 each may be one of the following:
	— H
	— CH ₃ or
	— CH ₂ CH ₃
	Molecular weight: Macromolecules: from about 20 000 (n about 100) up to about 380 000 g/mol (n about 2 000)
	Assay: Content not less than 25 % and not more than 33 % of methoxyl groups (-OCH ₃) and not more than 5 % of hydroxyethoxyl groups (-OCH ₂ CH ₂ Ol
	Slightly hygroscopic white or slightly yellowish or greyish odourless and tasteless, granular or fibrous powder.
	Solubility: Swelling in water, producing a clear to opalescent, viscous, colloidal solution. Insoluble in ethanol, ether and chloroform. Soluble in glacial acet acid.
	Purity:
	Loss on drying: $\leq 10 \%$ (105 °C, 3 hours)
	Sulphated Ash: ≤ 1.5 % determined at 800 ± 25 °C
	pH: ≥ 5.0 and ≤ 8.0 (1 % colloidal solution)
	Heavy metals:
	Arsenic: $\leq 3.0 \text{ mg/kg}$
	Lead: $\leq 2.0 \text{ mg/kg}$
	Mercury: $\leq 1.0 \text{ mg/kg}$
	Cadmium: ≤ 1,0 mg/kg

Authorised Novel Food	Specifications
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1-Methylnicotinamide chloride	Definition:
	Chemical name: 3-carbamoyl-1-methyl-pyridinium chloride
	Chemical formula: C ₇ H ₉ N ₂ OCl
	CAS No: 1005-24-9
	Molecular weight: 172,61 Da
	Description
	1-Methylnicotinamide chloride is white or off-white, crystalline solid produced by a chemical synthesis process.
	Characteristics/Composition
	Appearance: White – off-white, crystalline solid
	Purity: ≥ 98,5 %
	Trigonelline: ≤ 0,05 %
	Nicotinic Acid: ≤ 0,10 %
	Nicotinamide: ≤ 0,10 %
	Largest unknown impurity: ≤ 0,05 %
	Sum of unknown impurities: ≤ 0,20 %
	Sum of all impurities: $\leq 0.50 \%$
	Solubility: soluble in water and methanol. Practically insoluble in 2-propanol and dichloromethane
	Moisture: $\leq 0.3\%$
	Loss on drying: $\leq 1.0 \%$
	Residue on ignition: $\leq 0.1 \%$
	Residual Solvents and Heavy Metals
	Methanol: $\leq 0.3 \%$
	Heavy metals: $\leq 0,002 \%$
	Microbiological criteria:
	Total aerobic microbial count: ≤ 100 CFU/g
	Mould/yeast: ≤ 10 CFU/g
	Enterobacteriaceae: absence in 1 g
	Pseudomonas aeruginosa: absence in 1 g
	Staphylococcus aureus: absent in 1 g
	CFU: Colony Forming Units

Authorised Novel Food	Specifications
6S)-5-methyltetrahydrofolic acid, glucosamine salt	Description/Definition: Chemical name: N-[4-[[[(6S)-2-amino-1,4,5,6,7,8-hexahydro-5-methyl-4-oxo-6-pteridinyl]methyl]amino]benzoyl]-L-glutamic acid, glucosamine salt Chemical formula: C ₃₂ H ₅₁ N ₉ O ₁₆ Molecular weight: 817,80 g/mol (anhydrous) CAS No.: 1181972-37-1 Appearance: Creamy to light-brown powder Purity: Diastereoisomeric purity: At least 99 % of (6S)-5-methyltetrahydrofolic acid Glucosamine assay: 34-46 % in dry basis 5-Methyltetrahydrofolic acid assay: 54-59 % in dry basis Water: ≤ 8,0 % Heavy metals: Lead: ≤ 2,0 ppm Cadmium: ≤ 1,0 ppm Mercury: ≤ 0,1 ppm Arsenic: ≤ 2,0 ppm Boron: ≤ 10 ppm Microbiological criteria: Total aerobic microbial count: ≤ 100 CFU/g Yeasts and moulds: ≤ 100 CFU/g
Aonomethylsilanetriol (Organic ilicon)	Description/Definition: Chemical name: Silanetriol, 1-methyl- Chemical formula: CH ₆ O ₃ Si Molecular weight: 94,14 g/mol CAS No: 2445-53-6

Authorised Novel Food	Specifications
	Purity:
	Organic Silicon (monomethylsilanetriol) preparation (aqueous solution):
	Acidity (pH): 6,4-6,8
	Silicon: 100-150 mg Si/l
	Heavy metals:
	Lead: $\leq 1.0 \mu \text{g/l}$
	Mercury: $\leq 1.0 \mu \text{g/l}$
	Cadmium: $\leq 1,0 \mu g/l$
	Arsenic: $\leq 3.0 \mu g/l$
	Solvents:
	Methanol: ≤ 5,0 mg/kg (residual presence)
Mycelial extract from Shiitake	Description/Definition:
mushroom (Lentinula edodes)	The novel food ingredient is a sterile aqueous extract obtained from the mycelium of <i>Lentinula edodes</i> cultivated in a submerged fermentation. It is a light brown, slightly turbid liquid.
	Lentinan is a β -(1-3) β -(1-6)-D-glucan which has a molecular weight of approximately 5 \times 10 ⁵ Daltons, a degree of branching of 2/5 and a triple helical tertiary structure.
	Purity/Composition of the mycelial extract from Lentinula edodes:
	Moisture: 98 %
	Dry matter: 2 %
	Free glucose: < 20 mg/ml
	Total protein(1): $< 0.1 \text{ mg/ml}$
	N-containing constituents(²): < 10 mg/ml
	Lentinan: $0.8 - 1.2 \text{ mg/ml}$
	(¹) Bradford method
	(²) Kjeldahl method
M38	
	
Nicotinamide riboside chloride	Description/Definition:
	The novel food is a synthetic form of nicotinamide riboside.
	The novel food contains ≥ 90 % nicotinamide riboside chloride, predominantly in its β form, the remaining components being residual solvents, reaction by-products and degradation products.

▼<u>M38</u>

Authorised Novel Food	Specifications
	Nicotinamide riboside chloride:
	CAS number: 23111-00-4
	EC number: 807-820-5
	IUPAC name: 1-[(2R,3R,4S,5R)-3,4-dihydroxy-5-(hydroxymethyl)oxolan-2-yl]pyridin-1-ium-3-carboxamide;chloride
	Chemical formula: C ₁₁ H ₁₅ N ₂ O ₅ Cl
	Molecular weight: 290,7 g/mol
	Characteristics/Composition:
	Colour: White to light brown
	Form: Powder
	Identification: Conforms by NMR (nuclear magnetic resonance)
	Nicotinamide riboside chloride: ≥ 90 %
	Water content: ≤ 2 %
	Residual solvents:
	Acetone: $\leq 5000\text{mg/kg}$
	Methanol: ≤ 1 000 mg/kg
	Acetonitrile: ≤ 50 mg/kg
	Methyl tert-butyl ether: ≤ 500 mg/kg
	Reaction by-products:
	Methyl acetate: ≤ 1 000 mg/kg
	Acetamide: ≤ 27 mg/kg
	Acetic acid: ≤ 5 000 mg/kg
	Heavy metals:
	Arsenic: ≤ 1 mg/kg
	Microbiological criteria:
	Total Plate Count: ≤ 1 000 CFU/g
	Yeast and Mould: ≤ 100 CFU/g
	Escherichia coli: Absence in 10 g
▼ <u>M9</u>	
Noni fruit juice (Morinda citrifoli	a) Description/Definition:
	Noni fruits (fruits of Morinda citrifolia L.) are pressed. The obtained juice is pasteurised. An optional fermentation step before or after the pressing may occur.
	Rubiadin: $\leq 10 \mu g/kg$
	Lucidin: ≤ 10 μg/kg

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ed to or to	

Noni fruit puree and concentrate

Authorised Novel Food

Noni fruit juice powder (Morinda

citrifolia)

(Morinda citrifolia)

Description/Definition:

amount as used in atomisation).

Description/Definition:

produced juice occurs in one or two ways:

The fruits of *Morinda citrifolia* are harvested by hand. Seeds and skin may be separated mechanically from the pureed fruits. After pasteurisation, the puree is packaged in aseptic containers and stored under cold conditions.

Specifications

Seeds and skin of the sun-dried fruits of Morinda citrifolia are separated. The obtained pulp is filtered to separate juice from the flesh. Desiccation of the

Or by zeodratation or drying and then mixing with an excipient, this process allows the juice to be dried initially and then mixed with maltodextrins (same

Either by atomisation using maize maltodextrins, this mixture is obtained by keeping the rates of inflow of the juice and maltodextrins constant

Morinda citrifolia concentrate is prepared from M. citrifolia puree by treatment with pectinolytic enzymes (50–60 °C for 1-2 h). Then the puree is heated to inactivate the pectinases and then immediately cooled. The juice is separated in a decanter centrifuge. Afterwards the juice is collected and pasteurised, prior to being concentrated in a vacuum evaporator from a brix of 6 to 8 to a brix of 49 to 51 in the final concentrate.

Composition:

Puree:

Moisture: 89-93 % Protein: < 0,6 g/100 g Fat: ≤ 0,4 g/100 g Ash: < 1,0 g/100 g

Total carbohydrates: 5-10 g/100 g

Fructose: 0,5-3,82 g/100 g Glucose: 0,5-3,14 g/100 g Dietary fibre: < 0,5-3 g/100 g

5,15-dimethylmorindol (1): $\leq 0,254 \mu \text{g/ml}$

Lucidin (1): Not detectable Alizarin (1): Not detectable Rubiadin (1): Not detectable

Concentrate:
Moisture: 48-53 %

Authorised Novel Food	Specifications
	Protein: 3-3,5 g/100 g
	Fat: < 0,04 g/100 g
	Ash: 4,5-5,0 g/100 g
	Total carbohydrates: 37-45 g/100 g
	Fructose: 9-11 g/100 g
	Glucose: 9-11 g/100 g
	Dietary fibre: 1,5-5,0 g/100 g
	5,15-dimethylmorindol (1): $\leq 0,254 \mu \text{g/ml}$
	(¹) By an HPLC-UV method developed and validated for the analysis of anthraquinones in Morinda citrifolia puree and concentrate. Limits of detection: 2,5 ng/ml (5,15 dimethylmorindol); 50,0 ng/ml (lucidin); 6,3 ng/ml (alizarin) and 62,5 ng/ml (rubiadin).
Noni leaves (Morinda citrifolia)	Description/Definition:
	After cutting, the leaves of <i>Morinda citrifolia</i> are subject to drying and roasting steps. The product has a particle size ranging from broken leaves to coarse powder with fines. It is of greenish brown to brown colour.
	Purity/Composition:
	Moisture: < 5,2 %
	Protein: 17- 20 %
	Carbohydrate: 55-65 %
	Ash: 10-13 %
	Fat: 4-9 %
	Oxalic acid: < 0,14 %
	Tannic acid: < 2,7 %
	5,15-dimethylmorindol: < 47 mg/kg
	Rubiadin: non detectable, ≤ 10 µg/kg
	Lucidin: non detectable, ≤ 10 μg/kg
Noni fruit powder (<i>Morinda</i> citrifolia)	Description/Definition: Noni fruit powder is made from pulped noni (Morinda citrifolia L.) fruits by freeze-drying. Fruits are pulped and seeds are removed. After freeze-drying
	during which water is removed from noni fruits, the remaining noni pulp is milled to a powder and encapsulated.

Authorised Novel Food	Specifications
	Purity/Composition
	Moisture: 5,3-9 %
	Protein: 3,8-4,8 g/100 g
	Fat: 1-2 g/100 g
	Ash: 4,6-5,7 g/100 g
	Total carbohydrates: 80-85 g/100 g
	Fructose: 20,4-22,5 g/100 g
	Glucose: 22-25 g/100 g
	Dietary fibre: 15,4-24,5 g/100 g
	5,15-dimethylmorindol (1): $\leq 2,0 \mu g/ml$
	(1) By an HPLC-UV method developed and validated for the analysis of anthraquinones in Morinda citrifolia fruit powder. Limits of detection: 2,5 ng/ml (5,15 dimethylmorindol)
Odontella aurita microalgae	Silicon: 3,3 %
8	Crystalline silica: max 0,1-0,3 % as impurity
Oil enriched with phytosterols/phytostanols	Description/Definition:
	Oil enriched with phytosterols/phytostanols is composed of an oil fraction and a phytosterol fraction.
	Acylglycerol Distribution:
	Free fatty acids (expressed as oleic acid): ≤ 2.0 %
	Monoacylglycerols (MAG): ≤ 10 %
	Diacylglycerols (DAG): ≤ 25 %
	Triacylglycerols (TAG): Making up the balance
	Phytosterol fraction:
	$β$ -sitosterol: $\le 80 \%$
	$β$ -sitostanol: $\le 15\%$
	campesterol: ≤ 40 %
	campestanol: $\leq 5.0 \%$
	stigmasterol: ≤ 30 %
	stigmasterol: $\leq 30 \%$ brassicasterol $\leq 3.0 \%$

Authorised Novel Food		Specifications	
	Others: Moisture and volatile: $\leq 0.5 \%$		
	Peroxide value (PV): < 5,0 meq/kg		
	Trans fatty acids: ≤ 1 %		
	- · ·	ivalent method) of phytosterols/phytostanols:	
	Phytosterols and phytostanols extracte than 99 %.	d from sources other than vegetable oil suitable for for	od have to be free of contaminants, best ensured by a purity of more
Oil extracted from squids	Acid value: ≤ 0.5 KOH/g oil Peroxide value (PV): ≤ 5 meq O_2/kg p-Anisidine value: ≤ 20 Cold test at 0 °C: ≤ 3 hours Moisture: ≤ 0.1 % (w/w) Unsaponifiable matter: ≤ 5.0 %Trans Docosahexaeonic acid: ≥ 20 % Eicosapentaenoic acid: ≥ 10 %		
Partially defatted chia seed (Salvia hispanica) powders	Description/Definition: The novel foods are partially defatted Physical-sensorial: Foreign matter: 0,1 %	d chia seed (Salvia hispanica) powders obtained by	pressing and grinding of the whole seeds of Salvia hispanica L.
		Powder with high protein content	Powder with high fibre content
	Particle size	≤ 130 μm	≤ 400 μm
	Chemical composition:		
		Salvia hispanica powder with high protein content	Salvia hispanica powder with high fibre content
	Moisture	≤ 9,0 %	≤ 9,0 %
	Protein	≥ 40,0 %	≥ 24,0 %
	Fat	≤ 17 %	≤ 12 %
	Fibre	≤ 30 %	≥ 50 %

	Authorised Novel Food	Specifications	
		Microbiological criteria:	
		Total plate count: ≤ 10 000 CFU/g	
		Yeasts: $\leq 500 \text{ CFU/g}$	
		Moulds: ≤ 500 CFU/g	
		Staphylococcus aureus: ≤ 10 CFU/g	
		Coliforms: < 100 MPN/g	
		Enterobacteriaceae: ≤ 100 CFU/g	
		Bacillus cereus: ≤ 50 CFU/g	
		Escherichia coli: < 10 MPN/g	
		Listeria monocytogenes: Absence/g	
		Salmonella spp.: Absence in 25 g	
		Contaminants:	
		Arsenic: ≤ 0,1 ppm	
		Cadmium: ≤ 0,1 ppm	
		Lead: ≤ 0,1 ppm	
		Mercury: ≤ 0.1 ppm	
		Total aflatoxins: ≤ 4 ppb	
		Ochratoxin A: ≤ 1 ppb	
▼ <u>M60</u>			
	Partially defatted rapeseed powder from <i>Brassica rapa</i> L. and	Definition: The powder is produced from the partially defatted seeds of non-genetically modified <i>Brassica rapa</i> L. and <i>Brassica napus</i> L. double low (00) cultivars through a series of processing steps to reduce glucosinolates and phytates.	
	Brassica napus L.	Source: Brassica rapa L. and Brassica napus L. seeds	
		Characteristics/Composition:	
		Protein (N × 6,25): 33,0-43,0 %	
		Lipids: 14,0 – 22,0 %	
		Total Carbohydrates(*): 33,0 – 40,0 %	
		Total Fibre(**): 33,0 – 43,0 %	

Authorised Novel Food	Specifications
	Moisture: < 7,0 %
	Ash: 2,0–5,0 %
	Total Glucosinolates: < 0,3 mmol/kg (≤ 120 mg/kg)
	Phytate: < 1,5 %
	Peroxide value (in novel food weight): ≤ 3,0 mEq O ₂ /kg
	Heavy Metals:
	Lead: < 0,2 mg/kg
	Arsenic (inorganic): < 0,2 mg/kg
	Cadmium: < 0,2 mg/kg
	Mercury: < 0,1 mg/kg
	Aluminium: < 35,0 mg/kg
	Microbiological criteria:
	Total plate count (30 °C): < 5 000 CFU/g
	Enterobacteriaceae: < 10 CFU/g
	Salmonella sp.: Negative/25 g
	Yeast and mould: < 100 CFU/g
	Bacillus cereus: < 100 CFU/g
	(*) By difference: 100 % – [protein % + moisture % + fat % + ash %]
	(**) AOAC 2011.25 (Enzymatic gravimetry)
	CFU: Colony Forming Units, AOAC: Association of Official Agricultural Chemists
et from Panax notoginseng stragalus membranaceus	Description/Definition:

▼<u>M53</u>

The novel food contains two extracts. One is an ethanol extract of the roots of Astragalus membranaceus (Fisch.) Bunge. The other is a hot water extract of the roots of Panax notoginseng (Burkill) F.H. Chen that is further concentrated using absorption on a resin and subsequent elution with 60 % ethanol. At the end of the manufacturing process both extracts are mixed (45-47,5 % of each extract) with maltodextrin (5-10 %).

Characteristics/Composition:

Total saponins: 1,5-5 % Ginsenoside Rb1: 0,1-0,5 % Astragaloside I: 0,01-0,1 %

▼<u>M53</u>

treatment agricultural and manufacturing practices Fruit added 40 % to 60 % of thawed fruit Fruit homogenised and added to other ingredients pH 3,2 to 4,2 Brix 7 to 42 Assured by added sugars a _w < 0,95 Assured by added sugars Final storage 60 days maximum at + 5 °C maximum Equivalent to storage regimen for conventionally process.		Authorised Novel Food		Specifications	
Ash: ≤ 1 % Moisture: ≤ 5 % Fat: ≤ 1.5 % Heavy metals: Arsenic: 5 0.3 mg/kg Microbiological criteria: Total plate count: ≤ 5000 CFU/g Total yeast and mould count: ≤ 500 CFU/g Enterobacteriaccae: < 10 CFU/g Enterobacteriaccae: < 10 CFU/g Enterobacteriaccae: < 10 CFU/g Escherichia coli: Absence in 375 g Samphylococcus aureus: Absence in 25 g CFU: colony forming units Parameter Parameter Fruit storage before high-pressure treatment Fruit added 40 % to 60 % of thawed fruit Fruit homogenised and added to other ingredients pl1 3.2 to 4.2 "Brix 7 to 42 Assured by added sugars Final storage 60 days maximum at + 5 °C maximum Equivalent to storage regimen for conventionally process." Full valued to storage regimen for conventionally process.					
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Fat: ≤ 1,5 % Heavy metals: Arsenic: ≤ 0,3 mg/kg Microbiological criteria: Total plate count: ≤ 5000 CFU/g Total yeast and mould count: ≤ 500 CFU/g Enterobacteriaceae: < 10 CFU/g Enterobacteriaceae: < 10 CFU/g Escherichia coit. Absence in 25 g Salmonella: Absence in 375 g Staphylococcus aureus: Absence in 25 g CFU: colony forming units Parameter Target Target Target Fruit storage before high-pressure treatment Fruit storage before high-pressure Fruit added 40 % to 60 % of thawed fruit Fruit homogenised and added to other ingredients pH 3,2 to 4,2 Brix 7 to 42 Assured by added sugars Final storage Final storage 60 days maximum at +5 °C maximum Equivalent to storage regimen for conventionally processing to the processing of the conventionally processing to the processing of the conventionally processing to the processing of the conventionally processing the processing of the processing of the processing the processing of the			Ash: ≤ 1 %		
Heavy metals: Arsenic: ≤ 0,3 mg/kg Microbiological criteria: Total plate count: ≤ 5000 CFU/g Total yeast and mould count: ≤ 5000 CFU/g Enterobacteriaccae: < 10 CFU/g Escherichia coli: Absence in 25 g Salmonella: Absence in 375 g Slaphylococcus aureus: Absence in 25 g CFU: colony forming units Parameter Parameter Fruit storage before high-pressure treatment Fruit storage before high-pressure treatment Absured truit-based preparations produced using high-pressure treatment Fruit added 40 % to 60 % of thawed fruit Fruit homogenised and added to other ingredients PH 3,2 to 4,2 Brix 7 to 42 Assured by added sugars Assured by added sugars Final storage Final storage 60 days maximum at +5 °C maximum Equivalent to storage regimen for conventionally processors.			Moisture: ≤ 5 %		
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Enterobacteriaceae: < 10 CFU/g Escherichia coli: Absence in 25 g Salmonella: Absence in 25 g Staphylococcus aureus: Absence in 25 g CFU: colony forming units Pasteurised fruit-based preparations produced using high-pressure treatment Parameter Parameter Target Comments Fruit storage before high-pressure treatment Fruit storage before high-pressure treatment Fruit added 40 % to 60 % of thawed fruit Fruit homogenised and added to other ingredients pH 3,2 to 4,2 Brix 7 to 42 Assured by added sugars a _w < 0,95 Assured by added sugars Final storage Final storage Final storage Final storage 60 days maximum at + 5 °C maximum Equivalent to storage regimen for conventionally process			Total plate count: ≤ 5 000 CFU/g		
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Pasteurised fruit-based preparations produced using high-pressure treatment Parameter Fruit storage treatment Fruit storage treatment Fruit added Fruit homogenised and added to other ingredients Parameter Fruit storage treatment Fruit added Fruit homogenised and added to other ingredients Fruit homogenised and added to other ingredients Parameter Fruit storage before high-pressure treatment Fruit added Fruit homogenised and added to other ingredients Assured by added sugars Final storage Final storage Final storage 60 days maximum at + 5 °C maximum Equivalent to storage regimen for conventionally process			CFU: colony forming units		
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Fruit storage before high-pressure treatment Fruit added 40 % to 60 % of thawed fruit pH 3,2 to 4,2 Brix 7 to 42 Assured by added sugars a _w <0,95 Assured by added sugars Final storage 60 days maximum at +5 °C maximum Equivalent to storage regimen for conventionally process.					
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° Brix 7 to 42 Assured by added sugars a _w < 0,95 Assured by added sugars Final storage 60 days maximum at +5 °C maximum Equivalent to storage regimen for conventionally process			Fruit added	40 % to 60 % of thawed fruit	Fruit homogenised and added to other ingredients
a _w < 0,95 Assured by added sugars Final storage 60 days maximum at + 5 °C maximum Equivalent to storage regimen for conventionally process			рН	3,2 to 4,2	
Final storage 60 days maximum at + 5 °C maximum Equivalent to storage regimen for conventionally processing to the final storage of the final storage for the final storage at the final storage of the final storage for the final storage for the final storage at the final storage for the			° Brix	7 to 42	Assured by added sugars
			$a_{ m w}$	< 0,95	Assured by added sugars
			Final storage	60 days maximum at + 5 °C maximum	Equivalent to storage regimen for conventionally processed product

	Authorised Novel Food	Specifications	
▼ <u>M35</u>			
	Phenylcapsaicin	Description/Definition:	
		Phenylcapsaicin (<i>N</i> -[(4-hydroxy-3-methoxyphenyl)methyl]-7-phenylhept-6-ynamide, C ₂₁ H ₂₃ NO ₃ , CAS no: 848127-67-3), is synthesized chemically via a two step synthesis process involving in a first step the production of the acetylenic acid intermediate through a reaction of phenyl acetylene with a carboxylic acid derivative, and in a second step a series of reactions of the acetylenic acid intermediate with vanillylamine derivative to produce phenylcapsaicin.	
		Characteristics/Composition:	
		Purity (% of dry matter): ≥ 98 %	
		Moisture: ≤ 0,5 %	
		Total synthesis related production by-products: $\leq 1.0 \%$	
		N,N-dimethyl formamide: ≤ 880 mg/kg	
		Dichloromethane: ≤ 600 mg/kg	
		Dimethoxyethane: ≤ 100 mg/kg	
		Ethyl acetate: $\leq 0.5 \%$	
		Other solvents: $\leq 0.5 \%$	
		Heavy metals:	
		Lead: ≤ 1,0 mg/kg	
		Cadmium: ≤ 1,0 mg/kg	
		Mercury: ≤ 0,1 mg/kg	
		Arsenic: ≤ 1,0 mg/kg	
		Microbiological criteria:	
		Total plate count: ≤ 10 CFU/g	
		Coliforms: ≤ 10 CFU/g	
		Escherichia coli: Negative/10 g	
		Salmonella sp.: Negative/10 g	
		Yeast and mould: ≤ 10 CFU/g	
		CFU: Colony Forming Units	

Authorised Novel Food	Specifications		
Phosphated maize starch	Description/Definition:		
	Phosphated maize starch (phosphated distarch phosphate) is a chemically modified resistant starch derived from high amylose starch by combining chemical treatments to create phosphate cross-links between carbohydrate residues and esterified hydroxyl groups.		
	The novel food ingredient is a white or nearly white powder.		
	CAS No: 11120-02-8		
	Chemical formula: $(C_6H_{10}O_5)_n$ $[(C_6H_9O_5)_2PO_2H]x$ $[(C_6H_9O_5)PO_3H_2]y$		
	n = number of glucose units; x, y = degrees of substitution		
	The chemical characteristics of phosphated distarch phosphate:		
	Loss on drying: 10-14 %		
	pH: 4,5-7,5		
	Dietary fibre: ≥ 70 %		
	Starch: 7-14 %		
	Protein: ≤ 0,8 %		
	Lipids: $\leq 0.8 \%$		
	Residual bound phosphorus: ≤ 0,4 % (as phosphorus) 'high amylose maize' as source		
Phosphatidylserine from fish	Description/Definition:		
phospholipids	The novel food ingredient is yellow to brown powder. Phosphatidylserine is obtained from fish phospholipids by an enzymatic transphosphorylation with the amino acid L-serine.		
	Specification of the phosphatidylserine product manufactured from fish phospholipids:		
	Moisture: < 5,0 %		
	Phospholipids: ≥ 75 %		
	Phosphatidylserine: ≥ 35 %		
	Glycerides: < 4,0 %		
	Free L-serine: < 1,0 %		
	Tocopherols: < 0,5 % (1)		
	Peroxide value (PV): < 5,0 meq O ₂ /kg		
	(1) Tocopherols may be added as antioxidants according to Commission Regulation (EU) No 1129/2011		

Authorised Novel Food	Specifications		
Phosphatidylserine from soya	Description/Definition:		
phospholipids	The novel food ingredient is off-white to light yellow powder. It is also available in liquid form with a clear brown to orange colour. The liquid form contains medium chain triacylglycerides (MCT) as a carrier. It contains lower levels of Phosphatidylserine due to the fact that it includes significant amounts of oil (MCT).		
	Phosphatidylserine from soya phospholipids is obtained through enzymatic transphosphatidylation of high-phosphatidylcholine soybean lecithin with the amino acid L-serine. Phosphatidylserine consists of a glycerophosphate skeleton conjugated with two fatty acids and L-serine via a phosphodiester linkage.		
	Characteristics of Phosphatidylserine from soya phospholipids:		
	Powder form:		
	Moisture: < 2,0 %		
	Phospholipids: ≥ 85 %		
	Phosphatidylserine: ≥ 61 %		
	Glycerides: < 2,0 %		
	free L-serine: < 1,0 %		
	Tocopherols: < 0,3 %		
	Phytosterols: < 0,2 %		
	Liquid form:		
	Moisture: < 2,0 %		
	Phospholipids: ≥ 25 %		
	Phosphatidylserine: ≥ 20 %		
	Glycerides: not applicable		
	free L-serine: < 1,0 %		
	Tocopherols: < 0,3 %		
	Phytosterols: < 0,2 %		
Phospholipid product containing	Description/Definition:		
equal amounts of phosphati- dylserine and phosphatidic acid	The product is manufactured through enzymatic conversion of soy lecithin. The phospholipid product is a highly concentrated, yellow-brown powder form of phosphatidylserine and phosphatidic acid at an equal level.		
	Specification of the product:		
	Moisture: ≤ 2,0 %		

Authorised Novel Food	Specifications
	Total phospholipids: ≥ 70 %
	Phosphatidylserine: ≥ 20 %
	Phosphatidic acid: ≥ 20 %
	Glycerides: ≤ 1,0 %
	Free L-serine: ≤ 1,0 %
	Tocopherols: $\leq 0.3\%$
	Phytosterols: $\leq 2.0 \%$
	Silicon dioxide is used with a maximum content of 1,0 %
Phospholipides from egg yolk	85 % and 100 % pure Phospholipides from egg yolk
Phytoglycogen	Description: White to off-white powder which is an odourless, colourless, flavourless polysaccharide derived from non-GM sweet corn using conventional food processing techniques
	Definition: Glucose polymer ($C_6H_{12}O_6$)n with linear linkages of $\alpha(1-4)$ glycosidic bonds branched every 8 to 12 glucose units by $\alpha(1-6)$ glycosidic bonds
	Specifications:
	Carbohydrates: 97 %
	Sugars: 0,5 %
	Fibre: 0,8 %
	Fat: 0,2 %
	Protein: 0,6 %
Phytosterols/phytostanols	Description/Definition:
	Phytosterols and phytostanols are sterols and stanols that are extracted from plants and may be presented as free sterols and stanols or esterified with food grade fatty acids.
	Composition (with GC-FID or equivalent method):
	β-sitosterol: < 81 %
	β -sitostanol: $< 35 \%$
	campesterol: < 40 %
	campestanol: < 15 %

Authorised Novel Food	Specifications
	stigmasterol: < 30 %
	brassicasterol: < 3,0 %
	other sterols/stanols: < 3,0 %
	Contamination/Purity (GC-FID or equivalent method):
	Phytosterols and phytostanols extracted from sources other than vegetable oil suitable for food have to be free of contaminants, best ensured by a purity of more than 99 % of the phytosterol/phytostanol ingredient.
Plum kernel oil	Description/Definition:
	Plum kernel oil is a vegetable oil obtained by cold pressing of plum (Prunus domestica) kernels.
	Composition:
	Oleic acid (C18:1): 68 %
	Linoleic acid (C18:2): 23 %
	γ-Tocopherol:80 % of total tocopherols
	β-Sitosterol: 80-90 % of total sterols
	Triolein: 40-55 % of triglycerides
	Cyanhydric acid: maximum 5 mg/kg oil
Potato proteins (coagulated) and	Dry substance: ≥ 800 mg/g
hydrolysates thereof	Protein (N * 6,25): \geq 600 mg/g (dry substance)
	Ash: $\leq 400 \text{ mg/g (dry substance)}$
	Glycoalkaloid (total): ≤ 150 mg/kg
	Lysinoalanine (total): ≤ 500 mg/kg
	Lysinoalanine (free): ≤ 10 mg/kg
Prolyl oligopeptidase (enzyme	Specification of the enzyme:
preparation)	Systematic name: Prolyl oligopeptidase
	Synonyms: Prolyl endopeptidase, proline-specific endopeptidase, endoprolylpeptidase
	Molecular weight: 66 kDa
	Enzyme Commission number: EC 3.4.21.26
	CAS number: 72162-84-6

Authorised Novel Food	Specifications
	Source: A genetically modified strain of Aspergillus niger (GEP-44)
	Description: Prolyl oligopeptidase is available as an enzyme preparation containing approximately 30 % maltodextrin.
	Specifications of the enzyme preparation of prolyl oligopeptidase:
	Activity: $> 580\ 000\ PPI(^1)/g\ (> 34,8\ PPU(^2)/g)$
	Appearance: Microgranulate
	Colour: Off-white to orange yellowish. The colour may change from batch to batch
	Dry Matter: > 94 %
	Gluten: < 20 ppm
	Heavy metals:
	Lead: ≤ 1,0 mg/kg
	Arsenic: ≤ 1,0 mg/kg
	Cadmium: ≤ 0,5 mg/kg
	Mercury: ≤ 0.1 mg/kg
	Microbiological criteria:
	Total aerobic plate count: $\leq 10^3$ CFU/g
	Total yeasts and moulds: $\leq 10^2 \text{ CFU/g}$
	Sulphite reducing anaerobes: ≤ 30 CFU/g
	Enterobacteriaceae: < 10 CFU/g Salmonella: Absence in 25 g
	Escherichia coli: Absence in 25 g
	Staphylococcus aureus: Absence in 10 g
	Pseudomonas aeruginosa: Absence in 10 g
	Listeria monocytogenes: Absence in 25 g
	Antimicrobial activity: Absent Mycotoxins: Below limits of detection: Aflatoxin B1, B2, G1, G2 (< 0,25 μg/kg), total Aflatoxins (< 2,0 μg/kg), Ochratoxin A (< 0,20 μg/kg), T-2 Toxin (< 5 μg/kg), Zearalenone (< 2,5 μg/kg), Fumonisin B1 and B2 (< 2,5 μg/kg)
	(1) PPI – Protease Picomole International
	(²) PPU – Prolyl Peptidase Units or Proline Protease Units

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Authorised Novel Food	Specifications

▼ M48

Protein extract from pig kidneys

Description/Definition:The protein extract is obtained from homogenised pig kidneys through a combination of salt precipitation and high speed centrifugation. The obtained precipitate contains essentially proteins with 7 % of the enzyme diamine oxidase (enzyme nomenclature E.C. 1.4.3.22) and is resuspended in a physiologic buffer system. The obtained pig kidney extract is formulated as encapsulated enteric coated pellets or enteric coated tablets to reach the active sites of digestion.

Basic Product:

Specification: pig kidney protein excerpt with natural content of Diamine oxidase (DAO):

Physical condition: liquid

Colour: brownish

Appearance: slightly turbid solution

pH value: 6,4-6,8

Enzymatic activity: > 2 677 kHDU DAO/ml (DAO REA (DAO Radioextractionassay))

Microbiological criteria:

Brachyspira spp.: negative (Real Time PCR)

Listeria monocytogenes: negative (Real Time PCR)

Staphylococcus aureus: < 100 CFU/g

Influenza A: negative (Reverse Transcription Real Time PCR)

Escherichia coli: < 10 CFU/g

Total aerobic microbiological count: < 10⁵ CFU/g

Yeasts/moulds count: < 10⁵ CFU/g

Salmonella: Absence/10g

Bile salt resistant enterobacteriaceae: < 10⁴ CFU/g

Final product:

Specification pig kidney protein excerpt with natural content of DAO (E.C. 1.4.3.22) in an enteric coated formulation:

Physical condition: solid

Colour: yellow grey

Appearance: micropellets or tablets

Enzymatic activity: 110-220 kHDU DAO/g pellet or g tablet (DAO REA (DAO Radioextractionassay))

▼<u>M48</u>

	Authorised Novel Food	Specifications
		Acid stability 15 min 0,1M HCl followed by 60 min Borat pH = 9,0: > 68 kHDU DAO/g pellet or g tablet (DAO REA (DAO Radioextractionassay)
		Humidity: < 10 %
		Staphylococcus aureus: < 100 CFU/g
		Escherichia coli: < 10 CFU/g
		Total aerobic microbiological count: < 10 ⁴ CFU/g
		Total combined yeasts/moulds count: < 10 ³ CFU/g
		Salmonella: Absence/10g
		Bile salt resistant enterobacteriaceae: < 10 ² CFU/g
7 M10		
	Pyrroloquinoline quinone disodium	Definition:
	salt	Chemical name: disodium 9-carboxy-4,5-dioxo-1 <i>H</i> -pyrrolo[5,4-f]quinoline-2,7-dicarboxylate
		Chemical formula: C ₁₄ H ₄ N ₂ Na ₂ O ₈
		CAS No: 122628-50-6
		Molecular weight: 374,17 Da
		Description
		Pyrroloquinoline quinone disodium salt is a reddish-brown powder produced by the non-genetically modified bacterium <i>Hyphomicrobium denitrificans</i> strain CK-275.
		Characteristics/Composition
		Appearance: Reddish-brown powder
		Purity: ≥ 99,0 % (dry weight)
		UV absorbance (A322/A259): 0.56 ± 0.03
		UV absorbance (A233/A259): 0,90 ± 0,09
		Moisture: ≤ 12,0 %
		Residual Solvent
		Ethanol: $\leq 0.05 \%$
		Heavy metals
		Lead: < 3 mg/kg
		Arsenic: < 2 mg/kg

▼M10

Authorised Novel Food	Specifications
	Microbiological criteria:
	Total viable cell count: ≤ 300 CFU/g
	Mould/yeast: ≤ 12 CFU/g
	Coliforms: absent in 1 g
	Hyphomicrobium denitrificans: ≤ 25 CFU/g
	CFU: Colony Forming Units

▼<u>M9</u>

Rapeseed oil high in unsaponifiable matter

Description/Definition:

Rapeseed oil high in unsaponifiable matter' is produced by vacuum distillation and it is different from refined rapeseed oil in the concentration of the unsaponifiable fraction (1 g in refined rapeseed oil and 9 g in 'rapeseed oil high in unsaponifiable matter'). There is a minor reduction of triglycerides containing monounsaturated and polyunsaturated fatty acids.

Purity:

Unsaponifiable matter: > 7,0 g/100 g

Tocopherols: > 0,8 g/100 g α-tocopherol (%): 30-50 % γ-tocopherol (%): 50-70 % δ-tocopherol (%): < 6,0 %

Sterols, triterpenic alcohols, methylsterols: > 5,0 g/100 g

Fatty acids in triglycerides:

palmitic acid: 3-8 %
stearic acid: 0,8-2,5 %
oleic acid: 50-70 %
linoleic acid: 15-28 %
linolenic acid: 6-14 %
erucic acid: < 2,0 %
Acid value: ≤ 6,0 mg KOH/g

Peroxide value (PV): ≤ 10 mEq O_2/kg

Authorised Novel Food	Specifications
	Heavy metals:
	Iron (Fe): < 1 000 µg/kg
	Copper (Cu): < 100 µg/kg
	Impurities:
	Polycyclic aromatic hydrocarbons (PAH) Benzo(a)pyrene: < 2 μg/kg
	Treatment with active carbon is required to ensure that polycyclic aromatic hydrocarbons (PAH) are not enriched in the production of 'rapeseed oil high in unsaponifiable matter.
Rapeseed Protein	Definition:
•	Rapeseed protein is an aqueous protein-rich extract from rapeseed press cake originating from non-genetically modified Brassica napus L. and Brassica rapa L
	Description:
	White to off-white, spray dried powder
	Total protein: ≥ 90 %
	Soluble protein: ≥ 85 %
	Moisture: ≤ 7,0 %
	Carbohydrates: ≤ 7,0 %
	Fat: $\leq 2,0 \%$
	Ash: ≤ 4,0 %
	Fibre: ≤ 0,5 %
	Total glucosinolates: ≤ 1 mmol/kg
	Purity:
	Total phytate: ≤ 1,5 %
	Lead: $\leq 0.5 \text{ mg/kg}$
	Microbiological criteria:
	Yeast and mould count: ≤ 100 CFU/g
	Aerobic bacteria count: ≤ 10 000 CFU/g
	Total coliform count: ≤ 10 CFU/g
	Escherichia coli: Absence in 10 g
	Salmonella: Absence in 25 g

	Authorised Novel Food	Specifications
▼ <u>M17</u>		
	Refined shrimp peptide	Description
C	concentrate	Refined shrimp peptide concentrate is a peptide mixture obtained from northern shrimp (<i>Pandalus borealis</i>) shells and heads via a series of purification steps following enzymatic proteolysis using a protease from <i>Bacillus licheniformis</i> and/or <i>Bacillus amyloliquefaciens</i> .
		Characteristics/Composition
		Total Dry matter (%): \geq 95,0 %
		Peptides (w/weight dry matter): ≥ 87,0 % of which peptides with molecular weight < 2 kDa: ≥ 99,9 %
		Fat (w/w) : $\leq 1.0 \%$
		Carbohydrates (w/w): ≤ 1.0 %
		Ash (w/w) : $\leq 15.0 \%$
		Calcium: ≤ 2,0 %
		Potassium: ≤ 0,15 %
		Sodium: ≤ 3,5 %
		Heavy Metals
		Arsenic (inorganic): ≤ 0,22 mg/kg
		Arsenic (organic): ≤ 51,0 mg/kg
		Cadmium: ≤ 0,09 mg/kg
		Lead: $\leq 0.18 \text{ mg/kg}$
		Total mercury: ≤ 0.03 mg/kg
		Microbiological criteria:
		Total viable cell count: ≤ 20 000 CFU/g
		Salmonella: ND/25g
		Listeria monocytogenes: ND/25g
		Escherichia coli: ≤ 20 CFU/g
		Coagulase positive <i>Staphylococcus aureus</i> : ≤ 200 CFU/g
		Pseudomonas aeruginosa: ND/25g
		Mould/yeast: ≤ 20 CFU/g
		CFU: Colony Forming Units
		ND: Not Detectable

Authorised Novel Food	Specifications
Trans-resveratrol	Description/Definition:
	Synthetic <i>Trans</i> -resveratrol is off-white to beige crystals.
	Chemical name: 5-[(E)-2-(4-hydroxyphenyl)ethenyl]benzene-1,3-diol
	Chemical formula: C ₁₄ H ₁₂ O ₃
	Molecular weight: 228,25 Da
	CAS No: 501-36-0
	Purity:
	Trans-resveratrol: $\geq 98 \%-99 \%$
	Total by-products (related substances): $\leq 0.5 \%$
	Any single related substance: ≤ 0,1 %
	Sulphated ash: ≤ 0,1 %
	Loss on drying: ≤ 0,5 %
	Heavy metals:
	Lead: ≤ 1,0 ppm
	Mercury: ≤ 0.1 ppm
	Arsenic: ≤ 1,0 ppm
	Impurities:
	Diisopropylamine: ≤ 50 mg/kg
	Microbial source: A genetically modified strain of Saccharomyces cerevisiae
	Appearance: Off-white to slight yellow powder
	Particle size: 100 % less than 62,23 μm
	Trans-resveratrol content: Min. 98 % w/w (dry weight basis)
	Ash: Max. 0,5 % w/w
	Moisture: Max. 3 % w/w
Rooster comb extract	Description/Definition: Rooster comb extract is obtained from Gallus gallus by enzymatic hydrolysis of rooster comb and by subsequent filtration, concentration and precipitation
	steps. The principal constituents of rooster comb extract are the glycosaminoglycans hyaluronic acid, chondroitin sulphate A and dermatan sulphate (chondroitin sulphate B). White or almost white hygroscopic powder.

Authorised Novel Food	Specifications
	Hyaluronic acid: 60-80 %
	Chondroitin sulphate A: ≤ 5,0 %
	Dermatan sulphate (chondroitin sulphate B): $\leq 25 \%$
	pH: 5,0-8,5
	Purity:
	Chlorides: ≤ 1,0 %
	Nitrogen: ≤ 8,0 %
	Loss on drying: (105 °C for 6 hours): \leq 10 %
	Heavy metals:
	Mercury: $\leq 0.1 \text{ mg/kg}$
	Arsenic: ≤ 1,0 mg/kg
	Cadmium: ≤ 1,0 mg/kg
	Chromium: ≤ 10 mg/kg
	Lead: $\leq 0.5 \text{ mg/kg}$
	Microbiological criteria:
	Total viable aerobic count: $\leq 10^2$ CFU/g
	Escherichia coli: Absence in 1 g
	Salmonella: Absence in 1 g
	Staphylococcus aureus: Absence in 1 g
	Pseudomonas aeruginosa: Absence in 1g
Sacha Inchi oil from <i>Plukenetia</i>	Description/Definition:
volubilis	Sacha inchi oil is a 100 % cold pressed vegetable oil obtained from the seeds of <i>Plukenetia volubiis</i> L. It is a transparent, fluid (liquid) and shiny oil at room temperature. It has a fruity, light, green vegetable taste without undesirable flavours.
	Aspect, limpidity, shine, colour: Fluid at room temperature, clean, shiny yellow gold
	Odour and taste: Fruity, vegetable without non acceptable taste or odour

	Purity:
	Water and Volatiles: < 0,2 g/100 g
	Impurities insoluble in hexane: < 0,05 g/100 g
	Oleic acidity: < 2,0 g/100 g
	Peroxide value (PV): < 15 meq O ₂ /kg
	Trans fatty acids: < 1,0 g/100 g
	Total unsaturated fatty acids: > 90 %Omega 3 alpha linolenic acid (ALA): > 45 %
	Saturated fatty acids: < 10 %
	No trans fatty acids (< 0,5 %)
	No erucic acid (< 0,2 %)
	More than 50 % of tri-linolenin and di-linolenin-triglycerides
	Phytosterols composition and level
	No cholesterol (< 5,0 mg/100 g)
Salatrims	Description/Definition:
5 4.1.1. 5	Salatrim is the internationally recognised acronym for (short and long chain acyl triglyceride molecules). Salatrim is prepared by non-enzymatic inter-esterification of triacetin, tripropionin, tributyrin, or their mixtures with hydrogenated canola, soybean, cottonseed, or sunflower oil. Description: Clear, slightly amber liquid to a light coloured waxy solid at room temperature. Free of particulate matter and of foreign or rancid odour.
	Glycerol ester disribution:
	Triacylglycerols: > 87 %
	Diacylglycerols: ≤ 10 %
	Monoacylglycerols: ≤ 2,0 %
	Fatty acid composition:
	MOLE % LCFA (long chain fatty acids): 33-70 %

Authorised Novel Food	Specifications
	MOLE % SCFA (short chain fatty acids): 30-67 %
	Saturated long chain fatty acids: < 70 % by weight
	Trans fatty acids: ≤ 1,0 %
	Free fatty acids as oleic acid: $\leq 0.5\%$
	Triacylglycerol profile:
	Triesters (short/long of 0,5 to 2,0): \geq 90 %
	Triesters (short/long = 0): $\leq 10\%$
	Unsaponifiable material: $\leq 1,0\%$
	Moisture: $\leq 0.3\%$
	Ash: $\leq 0.1 \%$
	Colour: ≤ 3,5 Red (Lovibond)
	Peroxide value (PV): ≤ 2.0 Meq/Kg
	Teloxide value (1 v). \(\leq 2.0\) Med/Kg
Schizochytrium sp. oil rich in DHA	A Acid value: $\leq 0.5 \text{ mg KOH/g}$
and EPA	Peroxide value (PV): ≤ 5,0 meq/kg oil
	Oxidative stability: All food products containing <i>Schizochytrium sp.</i> oil rich in DHA and EPA should demonstrate oxidative stability by appropriate recognised national/international test methodology (e.g. AOAC)
	Moisture and volatiles: ≤ 0,05 %
	Unsaponifiables: ≤ 4,5 %
	Trans-fatty acids: ≤ 1 %
	DHA content: $\geq 22.5\%$
	EPA content: ≥ 10 %
Schizochytrium sp. (ATCC	The novel food is obtained from the strain ATCC PTA-9695 of the microalgae Schizochytrium sp.
PTA-9695) oil	Peroxide value (PV): ≤ 5.0 meq/kg oil
	Unsaponifiables: ≤ 3,5 %
	Trans-fatty acids: ≤ 2,0 %
	Free fatty acids: $\leq 0.4 \%$
	Docosapentaenoic acid (DPA) n-6: ≤ 7,5 %
	DHA content: ≥ 35 %

Authorised Novel Food	Specifications
Schizochytrium sp. (FCC-3204) oil	Description/Definition:
	The novel food is an oil produced from the strain FCC-3204 of the microalgae <i>Schizochytrium</i> sp.
	Composition:
	Acid value: ≤ 0,5 mg KOH/g
	Peroxide value (PV): ≤ 5,0 meq/kg oil
	Moisture and volatiles: ≤ 0,05 %
	Unsaponifiables: ≤ 4,5 %
	Trans-fatty acids: ≤ 1,0 %
	Docosahexaenoic acid (DHA): ≥ 32,0 %
	p-anisidine value: ≤ 10
Schizochytrium sp. oil	Acid value: ≤ 0,5 mg KOH/g
	Peroxide value (PV): ≤ 5,0 meq/kg oil
	Moisture and volatiles: $\leq 0.05 \%$
	Unsaponifiables: ≤ 4,5 %
	Trans-fatty acids: ≤ 1,0 %
	DHA content: ≥ 32,0 %
Schizochytrium sp. (118) oil	Acid value: ≤ 0,8 mg KOH/g Peroxide value (PV): ≤ 5,0 meq/kg oil
	Moisture and volatiles: ≤ 0,05 %
	Unsaponifiables: $\leq 3.5\%$
	Trans-fatty acids: $\leq 2.0 \%$ Free fatty acids: $\leq 0.4 \%$
	DHA content: ≥ 35 %
	Schizochytrium sp. (FCC-3204) oil Schizochytrium sp. oil Schizochytrium sp. (T18) oil

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	Authorised Novel Food	Specifications
▼ <u>M62</u>		
	Schizochytrium sp. (WZU477) oil	Description/Definition:
		The novel food is an oil produced from the strain WZU477 of the microalgae <i>Schizochytrium</i> sp. Composition:
		Acid value: ≤ 0,5 mg KOH/g
		Peroxide value (PV): ≤ 5,0 meq/kg oil
		Moisture and volatiles: ≤ 0,05 %
		Unsaponifiables: ≤ 4,5 %
		Trans-fatty acids: ≤ 1,0 %
		Docosahexaenoic acid (DHA): ≥ 32,0 %
		p-anisidine value: ≤ 10
▼ M22		
	Syrup from Sorghum bicolor (L.)	Description/Definition
	Moench.	The traditional food is syrup from Sorghum bicolor (L.) Moench (genus, Sorghum; family, Poaceae (alt. Gramineae)).
	(Traditional food from a third country)	The syrup is obtained from stalks of <i>S. bicolor</i> , after applying production processes such as crushing, extraction, and evaporation including a heat treatment in order to obtain a minimum of 74 °Brix syrup
		Compositional data of syrup from Sorghum bicolor (L.) Moench
		Water: 22,7 g/100 g
		Ash: 2,4
		Sugars, total: > 74,0 g/100 g
▼ <u>M9</u>		
	Fermented soybean extract	Description/Definition: Fermented soybean extract is an odourless milk-white coloured powder. It is comprised of 30 % fermented soybean extract powder and 70 % resistant dextrin (as carrier) from corn-starch, which is added during the processing. Vitamin K ₂ is removed during the manufacturing process. Fermented soybean extract contains nattokinase isolated from natto, a foodstuff produced by the fermentation of non-genetically modified soybeans (Glycine max (L.)) with a selected strain of Bacillus subtilis var. natto. Nattokinase activity: 20 000 -28 000 Fibrin degradation unit/g(¹) Identity: Confirmable

Authorised Novel Food	Specifications
	Condition: No offensive taste or smell Loss on drying: ≤ 10 % Vitamin K ₂ : ≤ 0,1 mg/kg Heavy metals:
	Lead: $\leq 5.0 \text{ mg/kg}$
	Arsenic: ≤ 3,0 mg/kg
	Microbiological criteria:
	Total viable aerobic count: $\leq 10^3 \text{ CFU}(^3)/\text{g}$
	Yeast and mould: $\leq 10^2$ CFU/g
	Coliforms: ≤ 30 CFU/g
	Spore-forming bacteria: ≤ 10 CFU/g
	Escherichia coli: Absence/25 g
	Salmonella: Absence/25 g
	Listeria: Absence/25 g
	(1) Assay method as described by Takaoka et al. (2010).
V <u>M55</u>	
Selenium-containing yeast (Yarrowia lipolytica) biomass	Description/Definition: The novel food is the dried and heat-killed selenium-containing biomass of the yeast <i>Yarrowia lipolytica</i> . The novel food is produced by fermentation in the presence of sodium selenite followed by a number of purification steps including a heat-killing step of the yeast to ensure the absence of viable <i>Yarrowia lipolytica</i> cells in the novel food. Characteristics/Composition: Total selenium: 165–200 μg/g Se-methionine (¹³): 100–140 μg/g Protein: 40–50 g/100 g Dietary fibre: 24–32 g/100 g Sugars: < 1 g/100 g Fat: 6–12 g/100 g Total ash: ≤ 15 % Water: ≤ 5 % Dry matter: ≥ 95 %

▼<u>M55</u>

	Authorised Novel Food	Specifications
		Heavy metals:
		Lead: $\leq 3.0 \text{ mg/kg}$
		Cadmium: ≤ 1,0 mg/kg
		Mercury: $\leq 0.1 \text{ mg/kg}$
		Microbiological criteria:
		Total aerobic microbial count: $\leq 5 \times 10^3$ CFU/g
		Total yeast and mould count: $\leq 10^2$ CFU/g
		Viable Yarrowia lipolytica cells (14): < 10 CFU/g (i.e. limit of detection)
		Coliforms: ≤ 10 CFU/g
		Salmonella spp.: Absence in 25 g
		CFU: colony forming units
▼ <u>M59</u>		
	3'-Sialyllactose (3'-SL) sodium salt	Description:
	(microbial source)	3'-Sialyllactose (3'-SL) sodium salt is a purified, white to off-white powder or agglomerate that is produced by a microbial process and contains limited levels
	(microsian source)	of lactose, 3'-sialyl-lactulose, and sialic acid
		Source: Genetically modified strain of Escherichia coli K-12 DH1
		Definition:
		Chemical formula: C ₂₃ H ₃₈ NO ₁₉ Na
		Chemical name: N -Acetyl- α -D-neuraminyl- $(2\rightarrow 3)$ - β -D-galactopyranosyl- $(1\rightarrow 4)$ -D-glucose, sodium salt
		Molecular mass: 655,53 Da
		CAS No 128596-80-5
		Characteristics/Composition:
		Appearance: White to off-white powder or agglomerate
		Sum of 3'-Sialyllactose sodium salt, D-Lactose, and Sialic acid (% of dry matter): ≥ 90,0 % (w/w)
		3'-Sialyllactose sodium salt (% of dry matter): $\geq 88,0$ % (w/w)
		D-Lactose: $\leq 5.0 \%$ (w/w)
		Sialic acid: $\leq 1.5\%$ (w/w)
		3'-Sialyl-lactulose: ≤ 5,0 % (w/w) Sum of other carbohydrates: ≤ 3,0 % (w/w)
		Sum of other carbonydrates. $\leq 5.0\%$ (w/w) Moisture: $\leq 8.0\%$ (w/w)
		Sodium: $2.5 - 4.5\%$ (w/w)
		Solution: $2.5 = 4.5 \%$ (w/w) Chloride: $\leq 1.0 \%$ (w/w)
		pH (20 °C, 5 % solution): 4,5 -6,0
		Residual protein: $\leq 0.01 \%$ (w/w)

▼<u>M59</u>

Authorised Novel Food	Specifications
	Microbiological criteria:
	Aerobic mesophilic bacteria total plate count: ≤ 1000 CFU/g
	Enterobacteriaceae: ≤ 10 CFU/g
	Salmonella sp.: Absence in 25 g
	Yeast: $\leq 100 \text{ CFU/g}$
	Mould: $\leq 100 \text{ CFU/g}$
	Residual endotoxins: ≤ 10 EU/mg
	CFU: Colony Forming Units; EU: Endotoxin Units.
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	Description:
	6'-Sialyllactose (6'-SL) sodium salt is a purified, white to off-white powder or agglomerate that is produced by a microbial process and contains limited level
(microbial source)	of lactose, 6'-sialyl-lactulose, and sialic acid.
	Source: Genetically modified strain of <i>Escherichia coli</i> K-12 DH1 Definition:
	Chemical formula: C ₂₃ H ₃₈ NO ₁₉ Na Chemical name: N-Acetyl-α-D-neuraminyl-(2→6)-β-D-galactopyranosyl-(1→4)-D-glucose, sodium salt
	Molecular mass: 655,53 Da
	CAS No 157574-76-0
	CAS No 15/5/4-70-0 Characteristics/Composition:
	Appearance: White to off-white powder or agglomerate
	Sum of 6'-Sialyllactose sodium salt, D-Lactose and Sialic acid (% of dry matter): ≥ 94.0 % (w/w)
	6'-Sialyllactose sodium salt (% of dry matter): ≥ 90.0 % (w/w)
	D-Lactose: ≤ 5,0 % (w/w)
	Sialic acid: $\leq 2.0 \%$ (w/w)
	6'-Sialyl-lactulose: $\leq 3.0\%$ (w/w)
	Sum of other carbohydrates: $\leq 3.0 \%$ (w/w)
	Moisture: $\leq 6.0 \%$ (w/w)
	Sodium: 2,5-4,5 % (w/w)
	Chloride: $\leq 1.0 \%$ (w/w)
	pH (20 °C, 5 % solution): 4,5-6,0
	Residual protein: $\leq 0.01 \%$ (w/w)
	6'-Sialyllactose ('6'-SL') sodium salt (microbial source)

▼<u>M58</u>

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	Authorised Novel Food	Specifications
		Microbiological criteria:
		Aerobic mesophilic bacteria total plate count: ≤ 1 000 CFU/g
		Enterobacteriaceae: ≤ 10 CFU/g
		Salmonella sp.: Absence in 25 g
		Yeast: $\leq 100 \text{ CFU/g}$
		Mould: $\leq 100 \text{ CFU/g}$
		Residual endotoxins: ≤ 10 EU/mg
		CFU: Colony Forming Units; EU: Endotoxin Units.
▼ <u>M41</u>	:	
	Spermidine-rich wheat germ extract (Triticum aestivum)	Description/Definition: Spermidine-rich wheat germ extract is obtained from non-fermented, non-sprouting wheat germs (<i>Triticum aestivum</i>) by the process of solid-liquid extraction targeting specifically, but not exclusively polyamines. Spermidine:(N-(3-aminopropyl)butane-1,4-diamine):0,8-2,4 mg/g Spermine: 0,4-1,2 mg/g Spermidine trichloride < 0,1 μg/g Putrescine: < 0,3 mg/g Cadaverine: ≤ 16,0 μg/g Mycotoxins: Aflatoxins (total): < 0,4 μg/kg Microbiological criteria: Total aerobic bacteria: < 10 000 CFU/g Yeast and moulds: < 100 CFU/g Escherichia coli: < 10 CFU/g Salmonella: Absence/25g Listeria monocytogenes: Absence/25g
▼ <u>M9</u>		
	Sucromalt	Description/Definition: Sucromalt is a complex mixture of saccharides which is produced from sucrose and a starch hydrolysate by means of an enzymatic reaction. In this process, glucose units are attached to saccharides from the starch hydrolysate by means of an enzyme produced by the bacterium <i>Leuconostoc citreum</i> or by means of a recombinant strain of the production organism <i>Bacillus licheniformis</i> . The resulting oligosaccharides are characterised by the presence of α -(1 \rightarrow 6) and α -(1 \rightarrow 3) glycosidic compounds. The overall product is syrup, in addition to these oligosaccharides, contains mainly fructose but also the disaccharide leucrose and other disaccharides. Total solids: 75-80 %

Authorised Novel Food	Specifications		
	Moisture: 20-25 % Sulphatase: Max 0,05 % pH: 3,5-6,0 Conductivity < 200 (30 %) Nitrogen < 10 ppm Fructose: 35-45 % d.w. Leucrose: 7-15 % d.w. Other disaccharides: Max 3 % Higher saccharides: 40-60 % d.w		
Sugar cane fibre	Description/Definition: Sugar Cane Fibre is derived from the dry cell wall or fibrous residue remaining after expression or extraction of sugar juice from sugar cane, of the Saccharum genotype. It consists primarily of cellulose and hemicellulose. The production process consists of several steps, including: chipping, alkaline digestion, removal of lignins and other non-cellulosic components, bleaching of purified fibres, acid washing and neutralization. Moisture: ≤ 7,0 % Ash: ≤ 0,3 % Total Dietary Fibre (AOAC) dry basis (all insoluble): ≥ 95 % of which: Hemicellulose (20-25 %) and cellulose (70-75 %) Silica (ppm): ≤ 200 Protein: 0,0 % Fat: Trace pH: 4-7 Heavy metals: Mercury (ppm): ≤ 0,1 Lead (ppm): ≤ 1,0 Cadmium (ppm): ≤ 0,1 Microbiological criteria: Yeast and moulds (CFU/g): ≤ 1 000 Salmonella: Absence		

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▼ M5	1

Sugars	obtained	from	cocoa
(Theobi	roma cac	ao L.)	pulp

Authorised Novel Food

Description/Definition:

Sugars are obtained from the concentrated cocoa pulp (*Theobroma cacao* L.) juice either via a drying process or via a purification process to produce high purity glucose or fructose.

Specifications

Sugars produced by a drying process

Nutritional composition:

Total sugars (g/100g): > 80

Moisture (%): < 5

Microbiological criteria:

Total Plate Count (aerobic) (cfu/g): < 10⁴

Moulds and Yeasts (cfu/g): < 50

Enterobacteriaceae (cfu/g): < 10

Salmonella spp.: Absence in 25 g

Alicyclobacillus: Absence in 50 g

Thermo-acidophilic bacteria: Absence in 50 g

Sugars produced by a purification process

Nutritional composition of Glucose obtained from cocoa (Theobroma cacao L.) pulp:

Glucose content (%): > 93

Ash (%): < 0.2

Moisture (%): < 1,0

Nutritional composition of Fructose obtained from cocoa (Theobroma cacao L.) pulp:

Fructose content (%): > 98

Glucose content (%): < 0.5 %

Ash (%): < 0.2

Moisture (%):< 0,5

Microbiological criteria for glucose and fructose obtained from cocoa (Theobroma cacao L.) pulp:

Total Plate Count (aerobic) (cfu/g): $< 10^4$

Salmonella spp.: Absence in 25 g

Authorised Novel Food	Specifications
Sunflower oil extract	Description/Definition:
	The sunflower extract is obtained by a concentration factor of 10 of the unsaponifiable fraction of refined sunflower oil extracted from the seeds of the sunflower, <i>Helianthus Annuus</i> L.
	Composition:
	Oleic acid (C18:1): 20 %
	Linoleic acid (C18:2): 70 %
	Unsaponifiable matter: 8,0 %
	Phytosterols: 5,5 %
	Tocopherols: 1,1 %
70	
Synsepalum dulcificum dried fruits	Description/Definition:
	The novel food is lyophilised pulp and skin of pitted fruits of <i>Synsepalum dulcificum</i> (Schumach. & Thonn.) Daniell that belongs to the Sapotaceae famil The resulting dried cake is milled into a powder.
	Characteristics/Composition:
	Moisture $(g/100 \text{ g})$: < 6
	Ash (g/100 g): 3,5-8,5
	Total carbohydrates (g/100 g): 70-87
	Sugars (g/100 g): 50-75
	Fibre (g/100 g): 1-6,5
	Total protein (g/100 g): 3,5-6,0
	Miraculin (16) (g/100 g): 1,5-2,5
	Total fat (g/100 g): 0,50-3,50
	Microbiological criteria:
	Total aerobic colony count: < 10 ⁴ CFU (⁷)/g
	Bacillus cereus (presumptive): < 100 CFU/g
	Sulfite-reducing Clostridia: ≤ 30 CFU/g
	Total Enterobacteriaceae: < 100 CFU/g
	Yeasts and moulds: < 500 CFU/g

▼M70

Authorised Novel Food	Specifications	
	Pesticides:	
	Pesticide levels in accordance with Code number 0820990 ('others' in the group of fruit spices) set out in Regulation (EC) No 396/2005 (17)	

▼<u>M63</u>

Dried *Tenebrio molito*r larva (yellow mealworm)

Description/Definition:

The novel food is the whole, thermally dried yellow mealworm, either whole (blanched, oven-dried larva) or in the form of a powder (blanched, oven-dried, ground larva). The term 'mealworm' refers to the larval form of *Tenebrio molitor*, an insect species that belongs to the family of *Tenebrionidae* (darkling beetles).

The entire mealworms are meant for human consumption and no parts are removed.

A minimum 24 hours fasting period is required before the thermal drying step, to allow the larvae to discard their bowel content.

Characteristics/Composition:

Ash (% w/w): 3,5 - 4,5

Moisture (% w/w): 1-8

Crude protein (N x 6,25) (% w/w): 56-61

Digestible Carbohydrates (15) (% w/w): 1-6

Fat (% w/w): 25-30

of which saturated (% w/w): 4-9

Peroxide value (Meq O_2/kg fat): ≤ 5

Dietary fibre (% w/w): 4-7

Chitin (% w/w): 4-7

Heavy metals:

Lead: $\leq 0.075 \text{ mg/kg}$

Cadmium: ≤ 0.1 mg/kg

Mycotoxins:

Aflatoxins (Sum of B1, B2, G1, G2): ≤ 4 µg/kg

Aflatoxin B1: \leq 2 $\mu g/kg$

Deoxynivalenol: ≤ 200 μg/kg

Ochratoxin A: $\leq 1 \mu g/kg$

Authorised Novel Food	Specifications	
	Microbiological criteria:	
	Total aerobic colony count: ≤ 10 ⁵ CFU (⁷)/g	
	Yeasts and moulds: ≤ 100 CFU/g	
	Escherichia coli: ≤ 50 CFU/g	
	Salmonella spp.: Not detected in 25 g	
	Listeria monocytogenes: Not detected in 25 g	
	Sulfite-reducing Anaerobes: ≤ 30 CFU/g	
	Bacillus cereus (presumptive): ≤ 100 CFU/g	
	Enterobacteriaceae (presumptive): < 10 CFU/g	
	Coagulase-positive <i>staphylococci</i> : ≤ 100 CFU/g	
en, dried and powder forms		

▼<u>M78</u>

Frozen, dried and powder forms of yellow mealworm (*Tenebrio molitor* larva)

The novel food are frozen, dried and powder forms of yellow mealworm (*Tenebrio molitor* larva). The term 'mealworm' refers to the larval form of *Tenebrio molitor*, an insect species that belongs to the family of Tenebrionidae (darkling beetles). Another identified scientific synonym is *Tenebrio molitor* Linnaeus.

The entire mealworms are meant for human consumption, no parts are removed.

A minimum 24 hours fasting period is required before killing the insects by freezing, to allow the larvae to discard their bowel content.

The novel food is intended to be placed on the market in three different forms, namely: whole, blanched and frozen *T. molitor* larva (frozen); whole, blanched and freeze-dried *T. molitor* larva (dried) which may be in powder form (powder).

Parameters	Frozen	Dried or powder		
Characteristics/Composition				
Ash	0,9-1,10	3,6-4,1		
Moisture (% w/w)	69-75	≤ 5		
Crude protein (N x 6,25) (% w/w)	14-19	54-60		

▼<u>M78</u>

Authorised Novel Food	Specifications			
	Fat (% w/w) — of which saturated fatty acids (% fat)	7-12,5 20-29	27-30 20-29	
	Digestible carbohydrates (% w/w)	1-2	4-8	
	Dietary fibre (% w/w)	1,2-3,5	4-6	
	Chitin(*) (% w/w)	≤ 3	4-9	
	Peroxide value (Meq O ₂ /kg fat)	≤ 5	≤ 5	
	Contaminants			
	Heavy metals			
	Lead (mg/kg)	≤ 0,01	≤ 0,075	
	Cadmium (mg/kg)	≤ 0,05	≤ 0,1	
	Mycotoxins			
	Aflatoxins (Sum of B1, B2, G1, G2) (µg/kg)	≤ 4	≤ 4	
	Aflatoxin B1 (μg/kg)	≤ 2	≤ 2	
	Deoxynivalenol (μg/kg)	≤ 200	≤ 200	
	Ochratoxin A (µg/kg)	≤ 1	≤ 1	
	Dioxins and PCBs			
	Sum of dioxins and dl-PCBs (UB, WHO-TEQ2005)(**) (pg/g fat)	≤ 0,75	≤ 0,75	

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Authorised Novel Food	Specifications			
	Microbiological criteria			
	Total aerobic colony count (CFU/g)	≤ 10 ⁵	≤ 10 ⁵	
	Enterobacteriaceae (presumptive) (CFU/g)	≤ 100	≤ 100	
	Escherichia coli (CFU/g)	≤ 50	≤ 50	
	Listeria monocytogenes	Absence in 25g	Absence in 25g	
	Salmonella spp.	Absence in 25g	Absence in 25g	
	Bacillus cereus (presumptive) (CFU/g)	≤ 100	≤ 100	
	Coagulase positive Staphylococci (CFU/g	≤ 100	≤ 100	
	Sulfite-reducing Anaerobes (CFU/g)	≤ 30	≤ 30	
	Yeasts and moulds (CFU/g)	≤ 100	≤ 100	
	40 -444 4 4 4 400			

^(*) Chitin calculated as the difference between the Acid Detergent Fibre fraction and the Acid Detergent Lignin fraction (ADF-ADL), as described by Hahn et al. (2018).

CFU: colony forming units.

▼<u>M9</u>

Dried Tetraselmis chuii microalgae

Description/Definition:

The dried product is obtained from the marine microalgae *Tetraselmis chuii*, belonging to the *Chlorodendraceae* family, cultivated in sterile sea water in closed photobioreactors insulated from the outside air.

Purity/Composition:

Identified by means of nuclear marker rDNA 18~S (sequence analysed no less than 1~600 base pairs) in the National Centre for Biotechnology information (NCBI) database: Not less than 99.9~%

^(**) Upper bound sum of polychlorinated dibenzo-para-dioxins (PCDDs)-polychlorinated dibenzo-furans (PCDFs) and dioxin-like polychlorinated biphenyls (PCBs) expressed as World Health Organization toxic equivalent (using WHO-TEFs of 2005)).

Authorised Novel Food	Specifications		
	Humidity: ≤ 7,0 %		
	Proteins: 35-40 %		
	Ashes: 14-16 %		
	Carbohydrates: 30-32 %		
	Fibre: 2-3 %		
	Fat: 5-8 %		
	Saturated fatty acids: 29-31 % of total fatty acids		
	Monounsaturated fatty acids: 21-24 % of total fatty acids		
	Polyunsaturated fatty acids: 44-49 % of total fatty acids		
	Iodine: ≤ 15 mg/kg		
Therapon barcoo/Scortum	Description/Definition:		
	Scortum/Therapon barcoo is a species of fish in the family Terapontidae. It is an endemic fresh water species from Australia. It is now reared in fish farms.		
	Taxonomic Identification: Class: Actinopterygii > order: Perciformes > family: Terapontidae > genus: Therapon or Scortum barcoo		
	Composition of fish flesh:		
	Protein (%): 18-25		
	Moisture (%): 65-75		
	Ash (%): 0,5-2,0		
	Energy (KJ/Kg): 6000-11500		
	Carbohydrates (%): 0,0		
	Fat (%): 5-15		
	Fatty acids (mg FA/g fillet):		
	Σ PUFA n-3: 1,2-20,0		
	Σ PUFA n-6: 0,3-2,0		
	PUFA n-3/n-6: 1,5-15,0		
	Total omega 3 acids: 1,6-40,0		
	Total omega 6 acids: 2,6-10,0		
D-Tagatose	Description/Definition:		
	Tagatose is produced by isomerization of galactose by means of chemical or enzymatic conversion, or by epimerization of fructose by means of enzymatic conversion. These are single-step conversions.		
	Appearance: White or almost white crystals		
	Chemical name: D-tagatose		

Authorised Novel Food	Specifications		
	Synonym: D-lyxo-Hexulose		
	CAS number: 87-81-0		
	Chemical formula: $C_6H_{12}O_6$ Formula weight: $180,16$ (g/mol) Purity: Assay: ≥ 98 % on a dry weight basis Loss on drying: ≤ 0.5 % (102 °C, 2 hours)		
	Specific Rotation: $\left[\alpha\right]_{D}^{20}$: -4 to -5.6° (1 % aqueous solution)(1)		
	Melting range: 133–137 °C		
	Heavy metals:		
	Lead: $\leq 1.0 \text{ mg/kg}(*)$		
	(*) Determine using an atomic absorption technique appropriate to the specified level. The selection of sample size and method of sample preparation may be based on the principles of the method described in FNP 5. 'Instrumental methods' (1).		
	(1) Food and nutrition paper 5 Rev 2 – Guide to specifications for general notices, general analytical techniques, identification tests, test solutions and other reference materials (JECFA) 1991, 307 p.; English – ISBN 92-5-102991-1		
► <u>M50</u> Taxifolin-rich extract ◀	Description: Taxifolin-rich extract from the wood of Dahurian Larch (<i>Larix gmelinii</i> (Rupr.) Rupr) is a white to pale-yellow powder that crystallizes from hot aqueous solutions. ▶ M50 Definition: Chemical name: [(2R,3R)-2-(3,4 dihydroxyphenyl)-3,5,7-trihydroxy-2,3-dihydrochromen-4-one, also called (+) trans (2R,3R)- dihydroquercetin] and with no more than 2 % of the cis-form Specifications: Physical parameter Moisture: ≤ 10 %Compound analysis Taxifolin (m/m): ≥ 90,0 % of the dry weight		

Authorised Novel Food	Specifications		
	W. J. D. C. J.		
	Heavy Metals, Pesticide		
	Lead: $\leq 0.5 \text{ mg/kg}$		
	Arsenic: ≤ 0,02 mg/kg		
	Cadmium: ≤ 0,5 mg/kg Mercury: ≤ 0,1 mg/kg Dichlorodiphenyltrichloroethane (DDT): ≤ 0,05 mg/kg **Residual solvents** Ethanol: < 5 000 mg/kg		
	Microbiological criteria Total Plate Count (TPC): ≤ 10 ⁴ CFU/g Enterobacteria: ≤ 100/g Yeast and Mould: ≤ 100 CFU/g Escherichia coli: Absence/1 g		
	_	Salmonella: Absence/10 g	
	Staphylococcus aureus: Absence	/1 g	
		Pseudomonas: Absence/1g Usual range of components of the Taxifolin-rich extract (as per dry substance)	
	Usual range of components of		
	Extract component	Content, usual observed range (%)	
	Taxifolin	90 - 93	
	Aromadendrin	2,5 - 3,5	
	Eriodictyol	0,1-0,3	
	Quercetin	0,3-0,5	
	Naringenin	0,2-0,3	
	Kaempferol	0,01-0,1	
	Pinocembrin	0,05-0,12	
	Unidentified flavonoids	1 – 3	
	Water(*)	1,5	
	(*) Taxifolin in its hydrated form an	nd during the drying process is a crystal. This results on the inclusion of water of crystallisation in a quantity of 1,5 %.	

Trehalose

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Description/Definition:

Authorised Novel Food

A non-reducing disaccharide that consists of two glucose moieties linkes by an α -1,1-glucosidic bond. It is obtained from liquefied starch or from sucrose by a multistep enzymatic process. The commercial product is the dihydrate. Virtually odourless, white or almost white crystals with a sweet taste

Specifications

Synonyms: α,α-trehalose

Chemical name: α-D-glucopyranosyl-α-D-glucopyranoside, dihydrate

CAS No.: 6138-23-4 (dihydrate)

Chemical formula: C₁₂H₂₂O₁₁ · 2H₂O (dihydrate)

Formula weight: 378,33 (dihydrate) Assay: \geq 98 % on the dry basis

Determine using an atomic absorption technique appropriate to the specified level. The selection of sample size and method of sample preparation may be based on the principles of the method described in FNP 5 (1), 'Instrumental methods'

Method of assay:

Principle: trehalose is identified by liquid chromatography and quantified by comparison to a reference standard containing standard trehalose

Preparation of sample solution: weigh accurately about 3 g of dry sample into a 100 ml volumetric flask and add about 80 ml of purified, deionised water. Bring sample to complete dissolution and dilute to mark with purified deionised water. Filter through a 0,45 micron filter

Preparation of standard solution: dissolve accurately weighed quantities of dry standard reference trehalose in water to obtain a solution having known concentration of about 30 mg of trehalose per ml.

Apparatus: liquid chromatography equipped with a refractive index detector and integrating recorder

Conditions:

Column: Shodex Ionpack KS-801 (Showa Denko Co.) or equivalent

length: 300 mm
diameter: 10 mm
temperature: 50 °C
Mobile phase: water
flow rate: 0,4 ml/min
Injection volume: 8 µl

Procedure: inject separately equal volumes of the sample solution and the standard solution into the chromatograph.

Record the chromatograms and measure the size of response of the trehalose peak

Calculate the quantity, in mg, of trehalose in 1 ml of the sample solution by the following formula:

Authorised Novel Food	Specifications
	% trehalose = $100 \times (R_U/R_S) (W_S/W_U)$
	where
	R_S = peak area of trehalose in the standard preparation
	R_U = peak area of trehalose in the sample preparation
	W_S = weight in mg of trehalose in the standard preparation
	W_U = weight of dry sample in mg
Characteristics:	
	Identification:
	Solubility: Freely soluble in water, very slightly soluble in ethanol
	Specific rotation: $[\alpha]_D^{20} = +179^\circ$ (5 % aqueous solution, dihydrate), +199° (5 % aqueous solution, anhydrous substance)
	Melting point: 97 °C (dihydrate)
	Purity:
	Loss on drying: $\leq 1.5 \%$ (60 °C, 5h)
	Total ash: $\leq 0.05 \%$
	Heavy metals:
	Lead: ≤ 1,0 mg/kg
<u>50</u>	
	Description/Definition
bisporus)	Commercially grown Agaricus bisporus to which UV light treatment is applied to harvested mushrooms.
	UV radiation: a process of radiation in ultraviolet light within the wavelength of 200-800 nm.
	Vitamin D ₂
	Chemical name: (3β,5Z,7E,22E)-9,10-secoergosta-5,7,10(19),22-tetraen-3-ol
	Synonym: Ergocalciferol
	CAS No: 50-14-6
	Molecular weight: 396,65 g/mol
	Contents Vitamin D_2 in the final product: 5-20 μ g/100 g fresh weight at the expiration of shelf life.

	Authorised Novel Food	Specifications
▼ <u>M50</u>		
	UV-treated baker's yeast (Saccharomyces cerevisiae)	Description/Definition Baker's yeast (Saccharomyces cerevisiae) is treated with ultraviolet light to induce the conversion of ergosterol to vitamin D ₂ (ergocalciferol). Vitamin D ₂
		content in the yeast concentrate varies between 800 000-3 500 000 IU vitamin D/100 g (200-875 µg/g). The yeast may be inactivated.
		The yeast concentrate is blended with regular baker's yeast in order not to exceed the maximum level in the pre-packed fresh or dry yeast for home baking.
		Tan-coloured, free-flowing granules.
		Vitamin D ₂
		Chemical name: (5Z,7E,22E)-(3S)-9,10-secoergosta-5,7,10(19),22-tetraen-3-ol
		Synonym: Ergocalciferol
		CAS No.: 50-14-6
		Molecular weight: 396,65 g/mol
		Microbiological criteria for the yeast concentrate
		Coliforms: $\leq 10^3/g$
		Escherichia coli: ≤ 10/g
		Salmonella: Absence in 25 g
▼ <u>M9</u>		
	UV-treated bread	Description/Definition:
		UV-treated bread is yeast leavened bread and rolls (without toppings) to which a treatment with ultraviolet radiation is applied after baking in order to convert ergosterol to vitamin D ₂ (ergocalciferol).
		UV radiation: A process of radiation in ultraviolet light within the wavelength of 240-315 nm for maximum of 5 seconds with energy input of 10-50 mJ/cm ² .
		Vitamin D ₂ :
		Chemical name: (5Z,7E,22E)-3S-9,10-secoergosta-5,7,10(19),22-tetraen-3-ol
		Synonym: Ergocalciferol
		CAS No: 50-14-6
		Molecular weight: 396,65 g/mol
		Contents:
		Vitamin D ₂ (ergocalciferol) in the final product: 0,75-3 μg/100 g(¹)
		Yeast in dough: 1-5 g/100 g (²)
		(¹) EN 12821, 2009, European Standard.
		(²) Recipe calculation.

Authorised Novel Food

Description/Definition:

UV-treated milk

	e i treated man	Description Definition.	
		UV-treated milk is cow's milk (whole and semi-skimmed) to which a treatment with ultraviolet (UV) radiation via turbulent flow is applied after pasteurisation. The treatment of the pasteurised milk with UV radiation results in an increase in the vitamin D ₃ (cholecalciferol) concentrations by conversion of 7-dehydrocholesterol to vitamin D ₃ .	
		UV radiation: A process of radiation in ultraviolet light within the wavelength of 200-310 nm with energy input of 1 045 J/l.	
		Vitamin D ₃ :	
		Chemical name: (1S,3Z)-3-[(2E)-2-[(1R,3aS,7aR)-7a-methyl-1-[(2R)-6-methylheptan-2-yl]-2,3,3a,5,6,7-hexahydro-1H-inden-4-ylidene]ethylidene]-4-methyl-idenecyclohexan-1-ol	
		Synonym: Cholecalciferol	
		CAS No: 67-97-0	
		Molecular weight: 384,6377 g/mol	
		Contents:	
		Vitamin D ₃ in the final product:	
		Whole $milk(^{1})0,5-3,2 \mu g/100 g(^{2})$	
		Semi-skimmed milk(1): $0,1-1,5 \mu g/100 g(^2)$	
		(1) As defined by Regulation (EU) No 1308/2013 of the European Parliament and of the Council of 17 December 2013 establishing a common organisation of the markets in agricultural products and repealing Council Regulations (EEC) No 922/72, (EEC) No 234/79, (EC) No 1037/2001 and (EC) No 1234/2007 (OJ L 347, 20.12.2013, p. 671).	
		(²) HPLC	
▼ <u>M49</u>			
	Vitamin D ₂ mushroom powder	Description/Definition	
		Vitamin D ₂ mushroom powder is a granular powder made from homogenised Agaricus bisporus mushrooms that have been exposed to UV light.	
		The mushrooms are washed, homogenised and suspended in water to produce a mushroom slurry. The mushroom slurry is passed under a UV lamp. The slurry is then filtered, dried and ground, producing vitamin D_2 mushroom powder.	
		UV radiation: A process of radiation in ultraviolet light within a range of wavelength similar to those UV-treated novel foods authorised under the novel food regulation.	
		Characteristics/Composition	
		Vitamin D ₂ content: 1 000–1 300 μg/g of mushroom powder (12)	
		Moisture: ≤ 10,0 %	
		Ash: $\leq 13,5 \%$	

Specifications

Authorised Novel Food	Specifications
	Heavy Metals
	Lead (as Pb): ≤ 0,5 mg/kg
	Cadmium: ≤ 0,5 mg/kg
	Mercury: ≤ 0,1 mg/kg
	Arsenic: ≤ 0,3 mg/kg
	Mycotoxins
	Aflatoxins (sum of B1+B2+G1+G2): $< 4 \mu g/kg$
	Microbiological criteria:
	Total plate count: $\leq 5~000~\text{CFU}~(^7)/\text{g}$
	Yeast and mould: ≤ 100 CFU/g
	Salmonella sp.: Absent in 25 g
	Staphylococcus aureus: ≤ 10 CFU/g
	Escherichia coli: ≤ 10 CFU/g
	Coliforms: ≤ 10 CFU/g
	Enterobacteriaceae: ≤ 10 CFU/g
	Listeria monocytogenes: Absent in 25 g
3	
Vitamin D ₂ mushroom powder	Description/Definition:
	The novel food is mushroom powder produced from the dried whole <i>Agaricus bisporus</i> mushrooms. The process includes drying, milling and the controlled exposure of the mushroom powder to UV irradiation.
	UV radiation: A process of radiation in ultraviolet light within a range of wavelength similar to those UV-treated novel foods authorised under Regulation (EU) 2015/2283.
	Characteristics/composition:
	Vitamin D ₂ content: 580-595 μg/g of mushroom powder
	Ash: ≤ 13,5 %
	Water activity: < 0,5
	Moisture content: ≤ 7,5 %
	Carbohydrates: ≤ 35,0 %

Authorised Novel Food

Total Dietary Fibre: ≥ 15 %

		Crude protein (N × 6,25): \geq 22 %
		Fat: ≤ 4,5 %
		Heavy metals:
		Lead: $\leq 0.5 \text{ mg/kg}$
		Cadmium: $\leq 0.5 \text{ mg/kg}$
		Mercury: ≤ 0.1 mg/kg
		Arsenic: ≤ 0,3 mg/kg
		Mycotoxins:
		Aflatoxin B1: $\leq 0.10 \mu \text{g/kg}$
		Aflatoxins (sum of B1 + B2 + G1 + G2): $< 4 \mu g/kg$
		Microbiological criteria:
		Total plate count: $\leq 5000\text{CFU}(^{17})$
		Total yeast and mould count: < 100 CFU/g E. coli: < 10 CFU/g
		Salmonella spp.: Absence in 25 g
		Staphylococcus aureus: ≤ 10 CFU/g
		Coliforms: ≤ 10 CFU/g
		Listeria spp.: Absence in 25 g
		Enterobacteriaceae: < 10 CFU/g
▼ <u>M9</u>		
	Vitamin K ₂ (menaquinone)	This novel food is produced by a synthetic or microbiological process.
		Vitamin K ₂ (2-methyl-3-all-trans-polyprenyl-1,4-naphthoquinones), or the menaquinone series, is a group of prenylated naphthoquinone derivatives. The number of isoprene residues, where 1 isoprene unit consists of 5 carbons comprising the side chain, is used to characterise the menaquinone homologues containing primarily MK-7 and to a smaller extent MK-6.
		Vitamin K_2 (menaquinones) series with menaquinone-7 (MK-7)(n = 6) being $C_{46}H_{64}O_2$, menaquinone-6 (MK-6)(n = 5) being $C_{41}H_{56}O_2$ and menaquinone-4 (MK-4)(n = 3) being $C_{31}H_{40}O_2$.
		Chemical Name: (all-E)-2-(3,7,11,15,19,23,27-Heptamethyl-2,6,10,14,18,22,26-octacosaheptaenyl)-3-methyl-1,4-naphtalenedione
		CAS Number: 2124-57-4
		Molecular formula: C ₄₆ H ₆₄ O ₂
		1

Specifications

Authorised Novel Food	Specifications		
	Molecular weight: 649 g/mol		
	CH_3		
	(menadione moiety)		
	Specification of synthetic Vitamin K_2 (menaquinone-7)		
	Appearance: Yellow powder		
	Purity: Max 6,0 % cis-isomer, max 2,0 % other impurities		
	Content: 97-102 % Menaquinone-7 (including at least 92 % all-trans Menaquinone-7)		
	Specifications of microbiologically produced Vitamin K_2 (menaquinone-7)		
	Source: Bacillus subtilis spp. natto and Bacillus licheniformis		
	Appearance: Yellow powder or oil suspension		
Wheat bran extract	Description/Definition:		
	White crystalline powder obtained by enzymatic extraction from <i>Triticum aestivum</i> L. bran, rich in arabinoxylan oligosaccharides		
	Dry matter: Min. 94 %		
	Arabinoxylan oligosaccharides: Min 70 % of dry matter		
	Average degree of polymerisation of arabinoxylan oligosaccharides: 3-8		
	Ferulic acid (bound to arabinoxylan oligosaccharides): 1-3 % of dry matter		
	Total poly/oligosaccharides: Min 90 %		
	Protein: Max 2 % of dry matter		
	Ash: Max 2 % of dry matter		

▼<u>M9</u>

	Authorised Novel Food	Specifications
		Microbiological parameters:
		Mesophilic bacteria – total count: Max 10 000/g
		Yeasts: Max 100/g
		Fungi: Max 100/g
		Salmonella: Absence in 25g
		Bacillus cereus: Max 1000/g
		Clostridium perfringens: Max 1000/g
▼ <u>M7</u> 5	5	
	Wolffia arrhiza and/or Wolffia	Description/Definition:
	globosa fresh plants (Traditional food from a third country)	The traditional food consists of fresh plants of Wolffia arrhiza (L.) Horkel ex Wimm. and/or of Wolffia globosa (Roxb.) Hartog & Plas (family: Araceae).
	,	Microbiological criteria:
		Total plate count: < 10 ³ CFU/g
		Total yeast and mould count: < 100 CFU/g Total Enterobacteriaceae: < 100 CFU/g
		Escherichia coli: < 100 CFU/g
		Salmonella: Absence in 25 g
		Listeria monocytogenes: Absence in 25 g
		Staphylococcus aureus: Absence/10 g
		Heavy metals:
		Lead: < 0,3 mg/kg
		Arsenic (inorganic): < 0,10 mg/kg
		Cadmium: < 0,2 mg/kg
		Chromium: < 1 mg/kg
		Mercury: < 0,10 mg/kg
		Trace elements:
		Copper: < 0,8 mg/kg
		Molybdenum: < 0,3 mg/kg
		Zinc: < 5 mg/kg

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▼<u>M19</u>

Xylo-oligosaccharides

Authorised Novel Food

Description:

Boron: < 5 mg/kg

Manganese: < 6 mg/kg

Microcystins: 0,006 μg/g

Regulation (EC) No 396/2005 (17).

Cyanotoxins:

Pesticides:

The novel food is a mixture of xylo-oligosaccharides (XOS) which are obtained from corncobs (Zea mays subsp. mays) via hydrolysis by a xylanase from Trichoderma reesei followed by a purification process.

Pesticide levels in accordance with Code number 0254000 ('Subgroup (d) watercresses' in the group of Leaf vegetables, herbs and edible flowers)

Specifications

Characteristics/Composition

Parameter	Powder form 1	Powder form 2	Syrup form
Moisture (%)	≤ 5,0	≤ 5,0	70-75
Protein (g/100 g)		< 0,2	
Ash (%)		≤ 0,3	
pH		3,5-5,0	
Total carbohydrate content (g/100 g)	≥ 97	≥ 95	≥ 70
XOS content (dry basis) (g/100 g)	≥ 95	≥ 70	≥ 70
Other carbohydrates (g/100 g) (a)	2,5-7,5	2-16	1,5-31,5
Monosaccharides total (g/100 g)	0-4,5	0-13	0-29
Glucose (g/100 g)	0-2	0-5	0-4
Arabinose (g/100 g)	0-1,5	0-3	0-10
Xylose (g/100 g)	0-1,0	0-5	0-15
Disaccharides total (g/100 g)	27,5-48	25-43	26,5-42,5

Authorised Novel Food	Specifications				
	Xylobiose (XOS DP2) (g/100 g)	25-45	23-40	25-40	
	Cellobiose (g/100 g)	2,5-3	2-3	1,5-2,5	
	Oligosaccharides total (g/100 g)	41-77	36-72	32-71	
	xylotriose (XOS DP3) (g/100 g)	27-35	18-30	18-30	
	xylotetraose (XOS DP4) (g/100 g)	10-20	10-20	8-20	
	xylopentaose (XOS DP5) (g/100 g)	3-10	5-10	3-10	
	xylohexaose (XOS DP6) (g/100 g)	1-5	1-5	1-5	
	Xyloheptaose (XOS DP7) (g/100 g)	0-7	2-7	2-6	
	Maltodextrin (g/100 g) (^b)	0	20-25	0	
	Copper (mg/kg)		< 5,0		
	Lead (mg/kg)		< 0,5		
	Arsenic (mg/kg)		< 0,3		
	Salmonella (CFU (°)/25 g)		Negative		
	E, coli (MPN (d)/100 g)		Negative		
	Yeast (CFU/g)		< 10		
	Mould (CFU/g)		< 10		
	DP: Degree of polymerization (a) Other carbohydrates include monosaccharides (glue (b) Maltodextrin content is calculated according to the (c) CFU: Colony Forming Units. (d) MPN: Most Probable Number.	cose, xylose and arabinose) and cellot amount added in the process.	biose.		

	Authorised Novel Food	Specifications	
▼ <u>M30</u>			
	Yarrowia lipolytica yeast biomass	Description/Definition:	
		The novel food is the dried and heat-killed biomass of the yeast Yarrowia lipolytica.	
		Characteristics/Composition:	
		Protein: 45-55 g/100 g	
		Dietary fibre: 24-30 g/100 g	
		Sugars: < 1,0 g/100 g	
		Fat: 7-10 g/100 g	
		Total ash: $\leq 12 \%$	
		Water content: ≤ 5 %	
		Dry matter content: ≥ 95 %	
		Microbiological criteria:	
		Total Aerobic Microbial Count: $\leq 5 \times 10^3$ CFU/g	
		Total Yeast and Mould Count: $\leq 10^2$ CFU/g	
		Viable Yarrowia lipolytica cells (10): < 10 CFU/g (i.e. limit of detection)	
		Coliforms: $\leq 10 \text{ CFU/g}$	
		Salmonella spp.: Absence in 25 g	
▼ M9	-		
	Yeast beta-glucans	Description/Definition:	
		Beta-glucans are complex, high molecular mass (100-200 kDa) polysaccharides, found in the cell wall of many yeasts and cereals.	
		The chemical name for 'yeast beta-glucans' is (1-3),(1-6)-\(\beta\)-D-glucans.	
		Beta-glucans consist of a backbone of β-1-3-linked glucose residues that are branched by β-1-6-linkages, to which chitin and mannoproteins are linked by β-1-4-bonds.	
		Beta-glucans are isolated from yeast Saccharomyces cerevisiae.	
		The tertiary structure of the glucan cell wall of <i>Saccharomyces cerevisiae</i> consists of chains of β-1,3-linked glucose residues, branched by β-1,6-linkages, forming a backbone to which are linked chitin via β-1,4- bonds, β-1,6-glucans and some mannoproteins.	

Authorised Novel Food

This novel food is available in three different forms: soluble, insoluble and insoluble in water, but dispersible in many liquid matrices.
Chemical characteristics yeast (Saccharomyces cerevisiae) beta-glucans:
Soluble form:
Total carbohydrates: > 75 %
Beta-glucans (1,3/1,6): > 75 %
Ash: < 4,0 %
Moisture: < 8,0 %
Protein: < 3,5 %
Fat: < 10 %
Insoluble form:
Total carbohydrates: > 70 %
Beta-glucans (1,3/1,6): > 70 %
Ash: ≤ 12 %
Moisture: < 8,0 %
Protein: < 10 %
Fat: < 20 %
Insoluble in water, but dispersible in many liquid matrices:
(1,3)-(1,6)-\(\beta\)-Glucans: > 80 \(\psi\)
Ash: < 2,0 %
Moisture: < 6,0 %
Protein: < 4,0 %
Total fat: < 3,0 %
Microbiological data for insoluble in water, but dispersible in many liquid matrices:
Total plate count: < 1 000 CFU/g
Enterobacteriaceae: < 100 CFU/g
Total coliforms: < 10 CFU/g
Yeast: < 25 CFU/g

Specifications

Authorised Novel Food	Specifications
	Mould: < 25 CFU/g
	Salmonella: Absence in 25 g
	Escherichia coli: Absence in 1 g
	Bacillus cereus: < 100 CFU/g
	Staphylococcus aureus: Absence in 1 g
	Heavy metals for insoluble in water, but dispersible in many liquid matrices:
	▶ <u>M31</u> Lead: < 0,2 mg/kg
	Arsenic: < 0,2 mg/kg
	Mercury: < 0,1 mg/kg
	Cadmium: < 0,1 mg/kg ◀
Zeaxanthin	Description/Definition:
Zeaxantiiii	Zeaxanthin is a naturally occurring xanthophyll pigment, it is an oxygenated carotenoid.
	The synthetic zeaxanthin is presented either as a spray-dried powder of gelatin or starch base ('beadlets') with added α-tocopherol and ascorbyl palmitate or as a corn oil suspension with added α-tocopherol. Synthetic zeaxanthin is produced by a multi-step chemical synthesis from smaller molecules.
	Orange-red crystalline powder with little or no odour.
	Chemical formula: C ₄₀ H ₅₆ O ₂
	CAS No: 144-68-3
	Molecular weight: 568,9 daltons
	Physical-chemical properties:
	Loss on drying: < 0,2 %
	All-trans zeaxanthin: > 96 %
	Cis-zeaxanthin: < 2,0 %
	Other carotenoids: < 1,5 %
	Triphenylphosphine oxid (CAS No 791-28-6): < 50 mg/kg

Authorised Novel Food	Specifications			
Zinc L-pidolate	Description/Definition:			
	Zinc L-pidolate is a white to off-white powder, with characteristic odour.			
	International non-proprietary name (INN): L-pyroglutamic acid, Zinc salt			
	Synonyms: Zinc 5-oxoproline, Zinc pyroglutamate, Zinc pyrrolidone carboxylate, Zinc PCA, L-Zinc pidolate			
	CAS No.: 15454-75-8			
	Molecular formula: (C ₅ H ₆ NO ₃) ₂ Zn			
	Relative anhydrous molecular mass: 321,4			
	Appearance: White to slightly white powder			
	Purity:			
	Zinc L-pidolate (purity): ≥ 98 %			
	pH (10 % aqueous sol.): 5,0-6,0			
	Specific rotation: 19,6°- 22,8°			
	Water: ≤ 10,0 %			
	Glutamic acid: < 2,0 %			
	Heavy metals:			
	Lead: ≤ 3,0 ppm			
	Arsenic: ≤ 2,0 ppm			
	Cadmium: ≤ 1,0 ppm			
	Mercury: ≤ 0,1 ppm			

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(1) Commission Regulation (EU) No 231/2012	of 9 March 2012 laying down specifications for food additive	es listed in Annexes II and III to Regulation (EC)	No 1333/2008 of the European Parliament and of the Council
(OJ L 83, 22.3.2012, p. 1).			

Specifications

- (2) Commission Implementing Regulation (EU) 2015/175 of 5 February 2015 laying down special conditions applicable to the import of guar gum originating in or consigned from India due to contamination risks by pentachlorophenol and dioxins (OJ L 30, 6.2.2015, p. 10).
- ▶ M15 (3) OSC-DMAC (4-dimethylaminocinnamaldehyde) method (Ocean Spray Cranberries, Inc) Martin MA, Ramos S, Mateos R, Marais JPJ, Bravo-Clemente, L, Khoo C and Goya L. Food Res Intl 2015 71: 68-82.

 Modified from Cunningham DG, Vannozzi S, O'Shea E, Turk R (2002) In: Ho C-T, Zheng QY (eds) Quality Management of Nutraceuticals ACS Symposium series 803, Washington DC. Quantitation of PACs by DMAC Color Reaction pp 151-166.
- (4) BL-DMAC 4-dimethylaminocinnamaldehyde) method (Brunswick Lab) Multi-laboratory validation of a standard method for quantifying proanthocyanidins in cranberry powders. Prior RL, Fan E, Ji H, Howell A, Nio C, Pavne MJ. Reed J. J Sci Food Agric. 2010 Jul;90(9):1473-8.
- (5) The different values for these three parameters are due to the different methods used.
- (6) GAE: Gallic Acid Equivalents.
- (7) CFU: Colony Forming Units.

Authorised Novel Food

▶ M29 (8) HPLC/RI: High-performance liquid chromatography coupled with refractive index detection.

Microbiological criteria:

Pathogen: Absence

Yeasts and moulds: ≤ 100 CFU/g

- (9) CFU: Colony-forming unit. ◀
- (10) To be tested immediately after the heat-treatment step. Measures have to be in place to prevent cross-contamination with viable Yarrowia lipolytica cells during packaging and/or storage of the NF.
- (11) 2'-Fucosyl-galactose, Glucose, Galactose, Mannitol, Sorbitol, Galactitol, Trihexose, Allo-lactose and other structurally related carbohydrates.

Total viable mesophilic count: ≤ 1 000 CFU/g

- ► M49 (12) Converted from International Units (IU) using the conversion factor of 0.025 µg = 1 IU.
- (13) Expressed as selenium.
- (14) Applicable at all stages after the heat-treatment step to guarantee the absence of viable Yarrowia lipolytica cells and to be first tested immediately after the heat-treatment step. Measures have to be in place to prevent cross-contamination with viable Yarrowia lipolytica cells during packaging and/or storage of the novel food.
- (15) Digestible carbohydrates = 100 (crude protein + fat + dietary fibre + ash + moisture).
- (16) Miraculin is part of the total protein content.
- (17) Regulation (EC) No 396/2005 of the European Parliament and of the Council of 23 February 2005 on maximum residue levels of pesticides in or on food and feed of plant and animal origin and amending Council Directive 91/414/EEC (OJ L 70, 16.3.2005, p. 1).
- (18) Dietary fibre may not include chitin due to different analytical methods.
- (19) Upper bound sum of polychlorinated dibenzo-para-dioxins (PCDDs)-polychlorinated dibenzofurans (PCDFs) and dioxin-like polychlorinated biphenyls (PCBs) expressed as World Health Organization toxic equivalent (using WHO-TEFs of 2005)).