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8 November 2007

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[1] Text with EEA relevance

(Continued overleaf)

EN

II

(Information)

INFORMATION FROM EUROPEAN UNION INSTITUTIONS AND BODIES

COMMISSION

Non-opposition to a notified concentration

(Case COMP/M.4838 — SLP/TPG V/Avaya)

(Text with EEA relevance)

(2007/C 266/01)

On 26 September 2007, the Commission decided not to oppose the above notified concentration and to declare it compatible with the common market. This decision is based on Article 6(1)(b) of Council Regulation (EC) No 139/2004. The full text of the decision is available only in English and will be made public after it is cleared of any business secrets it may contain. It will be available:

— from the Europa competition website (http://ec.europa.eu/comm/competition/mergers/cases/). This website provides various facilities to help locate individual merger decisions, including company, case number, date and sectoral indexes,

— in electronic form on the EUR-Lex website under document number 32007M4838. EUR-Lex is the on-line access to European law (http://eur-lex.europa.eu).

IV

(Notices)

NOTICES FROM EUROPEAN UNION INSTITUTIONS AND BODIES

COMMISSION

Euro exchange rates [1]

7 November 2007

(2007/C 266/02)

1 euro =

	Currency	Exchange rate		Currency	Exchange rate
USD	US dollar	1,4722	RON	Romanian leu	3,4001
JPY	Japanese yen	166,07	SKK	Slovak koruna	33,183
DKK	Danish krone	7,4540	TRY	Turkish lira	1,7345
GBP	Pound sterling	0,69960	AUD	Australian dollar	1,5703
SEK	Swedish krona	9,2535	CAD	Canadian dollar	1,3354
CHF	Swiss franc	1,6589	HKD	Hong Kong dollar	11,4372
ISK	Iceland króna	86,40	NZD	New Zealand dollar	1,8767
NOK	Norwegian krone	7,7660	SGD	Singapore dollar	2,1180
BGN	Bulgarian lev	1,9558	KRW	South Korean won	1 333,89
CYP	Cyprus pound	0,5842	ZAR	South African rand	9,5611
CZK	Czech koruna	26,934	CNY	Chinese yuan renminbi	10,9563
EEK	Estonian kroon	15,6466	HRK	Croatian kuna	7,3315
HUF	Hungarian forint	253,64	IDR	Indonesian rupiah	13 426,46
LTL	Lithuanian litas	3,4528	MYR	Malaysian ringgit	4,9032
LVL	Latvian lats	0,7021	PHP	Philippine peso	63,709
MTL	Maltese lira	0,4293	RUB	Russian rouble	35,9600
PLN	Polish zloty	3,6435	THB	Thai baht	46,571

[1] *Source:* reference exchange rate published by the ECB.

NOTICES FROM MEMBER STATES

Information communicated by Member States regarding State aid granted under Commission Regulation (EC) No 70/2001 on the application of Articles 87 and 88 of the EC Treaty to State aid to small and medium-sized enterprises

(Text with EEA relevance)

(2007/C 266/03)

Aid No	XS 268/07
Member State	Malta
Region	—
Title of aid scheme or name of company receiving individual aid	Skema ghall-Promozzjoni ta' l-Innovazzjoni
Legal basis	Att dwar il-Korporazzjoni Maltija ghall-Intrapriza (KAP 463)
Type of measure	Aid scheme
Budget	Annual budget: MTL 0,3 million; overall budget: —
Maximum aid intensity	In conformity with Articles 4(2)-(6) and 5 of the Regulation
Date of implementation	1.9.2007
Duration	31.12.2013
Objective	Small and medium-sized enterprises
Economic sectors	All sectors eligible for aid to SMEs
Name and address of the granting authority	Malta Enterprise Corporation Enterprise Centre San Gwann SGN 09 Malta

Aid No	XS 269/07
Member State	Denmark
Region	—
Title of aid scheme or name of company receiving individual aid	Program for brugerdreven innovation støtter udvikling og afprøvning af den brugerdrevne tilgang til innovation til SMV'ere
Legal basis	Lov om erhvervsfremme § 2, stk. 2 og 3, stk. 2, § 4 stk. 1 og § 22 stk. 1,3 og 4 i lov nr. 602 af 24. juni 2005. Bekendtgørelse nr. 241 af 20. marts 2007

Type of measure	Aid scheme
Budget	Annual budget: DKK 94 million; overall budget: —
Maximum aid intensity	In conformity with Articles 4(2)-(6) and 5 of the Regulation
Date of implementation	15.6.2007
Duration	31.12.2011
Objective	Small and medium-sized enterprises
Economic sectors	All sectors eligible for aid to SMEs
Name and address of the granting authority	Dahlerups Pakhus Langelinie Alle 17 DK-2100 København Ø

Aid No	XS 270/07
Member State	Spain
Region	Islas Canarias
Title of aid scheme or name of company receiving individual aid	Fomento de la creación y expansión de empresas innovadoras de base tecnológica
Legal basis	Orden de la Consejería de Empleo, Industria y Comercio del Gobierno de Canarias, de 20 de septiembre de 2007, por la que se aprueban las bases reguladoras para el período 2007-2013, que han de regir en la concesión de subvenciones, mediante concurso, para fomentar la creación y expansión de empresas innovadoras de base tecnológica, y se efectúa convocatoria para 2007 (BOC nº 197, de 2 de octubre de 2007) www.gobiernodecanarias.org/boc/anexos/2007/197/00023-00063.pdf
Type of measure	Aid scheme
Budget	Annual budget: EUR 0,9 million; overall budget: —
Maximum aid intensity	In conformity with Articles 4(2)-(6) and 5 of the Regulation
Date of implementation	2.10.2007
Duration	31.12.2013
Objective	Small and medium-sized enterprises
Economic sectors	All sectors eligible for aid to SMEs
Name and address of the granting authority	Consejería de Empleo, Industria y Comercio del Gobierno de Canarias Avda de Anaga 35 — Edifício de Servicios Múltiples I, planta 8ª E-38071 Santa Cruz de Tenerife o C/ León y Castillo, 200 — Edifício de Servicios Múltiples III, planta 4ª E-35071 Las Palmas de Gran Canaria

Aid No	XS 271/07
Member State	Spain
Region	Islas Canarias
Title of aid scheme or name of company receiving individual aid	Innovación y desarrollo tecnológico de empresas industriales y de base tecnológica
Legal basis	Orden de la Consejería de Industria, Comercio y Nuevas Tecnologías del Gobierno de Canarias, de 13 de julio de 2007, por la que se convoca mediante concurso la concesión de subvenciones para la innovación y desarrollo tecnológico de empresas industriales y de base tecnológica (BOC nº 150, de 26 de julio de 2007) www.gobiernodecanarias.org/boc/anexos/2007/150/00015-00056.pdf
Type of measure	Aid scheme
Budget	Annual budget: EUR 2,98 million; overall budget: —
Maximum aid intensity	In conformity with Articles 4(2)-(6) and 5 of the Regulation
Date of implementation	26.7.2007
Duration	31.12.2007
Objective	Small and medium-sized enterprises
Economic sectors	All manufacturing
Name and address of the granting authority	Consejería de Empleo, Industria y Comercio del Gobierno de Canarias Avda de Anaga 35 — Edificio de Servicios Múltiples I, planta 8ª E-38071 Santa Cruz de Tenerife o C/ León y Castillo, 200 — Edificio de Servicios Múltiples III, planta 4ª E-35071 Las Palmas de Gran Canaria

NOTICES CONCERNING THE EUROPEAN ECONOMIC AREA

EFTA SURVEILLANCE AUTHORITY

Authorisation of State aid pursuant to Article 61 of the EEA Agreement and Article 1(3) in Part I of Protocol 3 to the Surveillance and Court Agreement

EFTA Surveillance Authority decision not to raise objections

(2007/C 266/04)

Date of adoption:	24 April 2007
EFTA State:	Norway
Aid No:	61274
Title:	Regional direct transport aid scheme
Objective:	The objective of the scheme is to offset competitive disadvantages resulting from extra transport costs for firms located in peripheral areas and within sparsely populated regions and thus situated long distances away from their markets.
Legal basis:	The legal basis for the scheme can be found in the State aid guidelines on National regional aid 2007-2013, National guidelines for regional transport aid ('Nasjonale retningslinjer for regional transportstøtte (Ytre rammeverk)') and in the State budget for 2007 (St.prp. nr. 1 (2006-2007) Kommunal- og regionaldepartementet).
Budget/Duration:	The annual budget of the scheme will not exceed NOK 70 million (approximately EUR 9 million). The duration of the scheme is from 1 January 2007 until 31 December 2013.

The authentic text of the decision, from which all confidential information has been removed, can be found at:

http://www.eftasurv.int/fieldsofwork/fieldstateaid/stateaidregistry

Medicinal products — List of marketing authorisations granted by the EEA EFTA States for the first half of 2006

(2007/C 266/05)

Subcommittee I — On the free movement of goods

With reference to EEA Joint Committee Decision No 74/1999 of 28 May 1999, the EEA Joint Committee is invited to note, at the meeting on 8 December 2006, the following lists concerning marketing authorisations for medicinal products for the period 1 January-30 June 2006:

ANNEX I List of new marketing authorisations

ANNEX II List of renewed marketing authorisations

ANNEX III List of extended marketing authorisations

ANNEX IV List of withdrawn marketing authorisations

ANNEX V List of suspended marketing authorisations

ANNEX I

List of new marketing authorisations

The following marketing authorisations have been granted in the EEA EFTA States during the period 1 January-30 June 2006:

EU-Number	Product	Country	Date of authorisation
EU/1/01/183/020-029	HBVAXPRO	Liechtenstein	28.2.2006
EU/1/01/187/001	DepoCyte	Liechtenstein	31.1.2006
EU/1/03/258/015-022	Avandamet	Liechtenstein	28.2.2006
EU/1/03/262/001-006	Emend	Liechtenstein	31.1.2006
EU/1/03/265/003-004	Bonviva	Liechtenstein	31.1.2006
EU/1/03/266/001-002	Bondenza	Liechtenstein	28.2.2006
EU/1/04/307/005-010	Zonegran	Liechtenstein	31.1.2006
EU/1/05/322/001	Yttriga	Liechtenstein	31.1.2006
EU/1/05/322/001/IS	Yttriga	Iceland	17.2.2006
EU/1/05/322/001/NO	Yttriga	Norway	10.2.2006
EU/1/05/323/001/NO-011/NO	ProQuad	Norway	24.4.2006
EU/1/05/323/001-011	ProQuad	Liechtenstein	30.4.2006
EU/1/05/323/001-011/IS	ProQuad	Iceland	25.4.2006
EU/1/05/324/001	Naglazyme	Liechtenstein	31.1.2006
EU/1/05/324/001/IS	Naglazyme	Iceland	22.6.2006
EU/1/05/324/01/NO	Naglazyme	Norway	23.2.2006
EU/1/05/325/001	Macugen	Liechtenstein	28.2.2006
EU/1/05/325/001/IS	Macugen	Iceland	10.3.2006
EU/1/05/325/001/NO	Macugen	Norway	2.3.2006
EU/1/05/326/001	Ionsys	Liechtenstein	31.1.2006
EU/1/05/326/001/IS	Ionsys	Iceland	22.2.2006
EU/1/05/326/001/NO	Ionsys	Norway	16.2.2006
EU/1/05/327/001/NO-017/NO	Exubera	Norway	16.2.2006
EU/1/05/327/001-017	Exubera	Liechtenstein	31.1.2006
EU/1/05/327/001-017/IS	Exubera	Iceland	22.3.2006
EU/1/05/328/001/NO-002/NO	Cubicin	Norway	16.2.2006
EU/1/05/328/001-002	Cubicin	Liechtenstein	28.2.2006
EU/1/05/328/001-002/IS	Cubicin	Iceland	13.2.2006
EU/1/05/329/001/NO-005/NO	Kiovig	Norway	27.1.2006
EU/1/05/329/001-005	Kiovig	Liechtenstein	31.1.2006

EU-Number	Product	Country	Date of authorisation
EU/1/05/329/001-005/IS	Kiovig	Iceland	10.2.2006
EU/1/05/330/001/NO-004/NO	Rotarix	Norway	8.3.2006
EU/1/05/330/001-004	Rotarix	Liechtenstein	28.2.2006
EU/1/05/330/001-004/IS	Rotarix	Iceland	27.4.2006
EU/1/05/331/001/NO-013/NO	Neupro	Norway	17.3.2006
EU/1/05/331/001-013	Neupro	Liechtenstein	28.2.2006
EU/1/05/331/001-013/IS	Neupro	Iceland	10.4.2006
EU/1/06/332/001/NO-003/NO	Omnitrope	Norway	5.5.2006
EU/1/06/332/001-003	Omnitrope	Liechtenstein	30.4.2006
EU/1/06/332/001-003/IS	Omnitrope	Iceland	11.5.2006
EU/1/06/333/001/NO-003/NO	Myozyme	Norway	8.5.2006
EU/1/06/333/001-003	Myozyme	Liechtenstein	30.4.2006
EU/1/06/334/001/NO-004/NO	Evoltra	Norway	27.6.2006
EU/1/06/334/001-004/IS	Evoltra	Iceland	26.6.2006
EU/1/06/335/001	Valtropin	Liechtenstein	30.6.2006
EU/1/06/335/001/IS	Valtropin	Iceland	12.5.2006
EU/1/06/335/001/NO	Valtropin	Norway	24.5.2006
EU/1/06/336/001	Tygacil	Liechtenstein	30.6.2006
EU/1/06/336/001/IS	Tygacil	Iceland	1.6.2006
EU/1/06/336/001/NO	Tygacil	Norway	24.5.2006
EU/1/06/337/001/NO-013/NO	M-M-Rvaxpro	Norway	19.5.2006
EU/1/06/337/001-013	M-M-Rvaxpro	Liechtenstein	30.6.2006
EU/1/06/337/001-013/IS	M-M-Rvaxpro	Iceland	13.6.2006
EU/1/06/338/001/NO-003/NO	DuoTrav	Norway	10.5.2006
EU/1/06/338/001-003	DuoTrav	Liechtenstein	30.6.2006
EU/1/06/338/001-003/IS	DuoTrav	Iceland	12.5.2006
EU/1/06/339/001/NO-002/NO	Preotact	Norway	19.5.2006
EU/1/06/339/001-002	Preotact	Liechtenstein	30.6.2006
EU/1/06/339/001-002/IS	Preotact	Iceland	18.5.2006
EU/1/06/340/001/NO-002/NO	Ganfort	Norway	1.6.2006
EU/1/06/340/001-002	Ganfort	Liechtenstein	30.6.2006
EU/1/06/340/001-002/IS	Ganfort	Iceland	15.6.2006
EU/1/06/341/001/NO-013/NO	Zostavax	Norway	6.6.2006
EU/1/06/341/001-013	Zostavax	Liechtenstein	30.6.2006
EU/1/06/341/001-013/IS	Zostavax	Iceland	19.6.2006

EU-Number	Product	Country	Date of authorisation
EU/1/06/344/001/NO-009/NO	Acomplia	Norway	30.6.2006
EU/1/06/344/001-009	Acomplia	Liechtenstein	30.6.2006
EU/1/06/345/001/NO-009/NO	Zimulti	Norway	30.6.2006
EU/1/06/345/001-009	Zimulti	Liechtenstein	30.6.2006
EU/1/96/022/019-022	Zyprexa	Liechtenstein	31.1.2006
EU/1/97/043/003	Revasc	Liechtenstein	31.1.2006
EU/1/97/044/007-008	Tasmar	Liechtenstein	28.2.2006
EU/1/97/046/031-033	Aprovel	Liechtenstein	31.1.2006
EU/1/97/049/031-033	Karvea	Liechtenstein	31.1.2006
EU/1/97/055/003	Viramune	Liechtenstein	31.1.2006
EU/1/98/063/007	Rebif	Liechtenstein	31.1.2006
EU/2/00/025/001-004	Advasure	Liechtenstein	28.2.2006
EU/2/03/037/005	ProteqFlu	Liechtenstein	28.2.2006
EU/2/03/038/005	ProteqFlu Te	Liechtenstein	28.2.2006
EU/2/04/043/001	Equilis StrepE	Liechtenstein	31.1.2006
EU/2/04/044/001	Aivlosin	Liechtenstein	31.1.2006
EU/2/04/044/002-005	Aivlosin	Liechtenstein	28.2.2006
EU/2/06/058/001/NO-003/NO	Flexicam	Norway	3.5.2006
EU/2/06/058/001-003	Flexicam	Liechtenstein	30.4.2006
EU/2/06/058/001-003/IS	Flexicam	Iceland	22.4.2006
EU/2/06/059/001	Convenia	Liechtenstein	30.6.2006
EU/2/97/004/014-015	Metacam	Liechtenstein	28.2.2006

ANNEX II

List of renewed marketing authorisations

The following marketing authorisations have been renewed in the EEA EFTA States during the period 1 January-30 June 2006:

EU-Number	Product	Country	Date of renewal
EU/1/00/154/001/NO-002/NO	Neospect	Norway	27.2.2006
EU/1/00/154/001-002	NeoSpect	Liechtenstein	28.2.2006
EU/1/00/154/001-002/IS	NeoSpect	Iceland	22.2.2006
EU/1/00/155/001/NO-006/NO	Luveris	Norway	27.2.2006
EU/1/00/155/001-006	Luveris	Liechtenstein	31.1.2006
EU/1/00/155/001-006/IS	Luveris	Iceland	25.4.2006
EU/1/00/156/002/NO-003/NO	Trizivir	Norway	27.4.2006
EU/1/00/156/002-003	Trizivir	Liechtenstein	30.4.2006
EU/1/00/157/001-034	Azomyr	Liechtenstein	28.2.2006
EU/1/00/160/001-034	Aerius	Liechtenstein	28.2.2006
EU/1/00/161/001-034	Neoclarityn	Liechtenstein	28.2.2006
EU/1/00/163/001/NO	Xeloda	Norway	3.3.2006
EU/1/00/163/001-002	Xeloda	Liechtenstein	28.2.2006
EU/1/00/163/001-002/IS	Xeloda	Iceland	7.3.2006
EU/1/00/163/002/NO	Xeloda	Norway	3.3.2006
EU/1/00/164/003/NO-006/NO	NutropinAq	Norway	20.4.2006
EU/1/00/164/003-005	NutropinAq	Liechtenstein	30.4.2006
EU/1/00/164/003-005/IS	NutropinAq	Iceland	18.5.2006
EU/1/00/165/001/NO-007/NO	Ovitrelle	Norway	2.3.2006
EU/1/00/165/001-007	Ovitrelle	Liechtenstein	28.2.2006
EU/1/00/165/007/IS	Ovitrelle	Iceland	11.4.2006
EU/1/00/166/001/NO-003/NO	NeuroBloc	Norway	3.5.2006
EU/1/00/166/001-003	NeuroBloc	Liechtenstein	30.4.2006
EU/1/00/166/001-003/IS	NeuroBloc	Iceland	9.6.2006
EU/1/00/167/001/NO-007/NO	Prevenar	Norway	3.5.2006
EU/1/00/167/001-007	Prevenar	Liechtenstein	30.4.2006
EU/1/00/167/001-007/IS	Prevenar	Iceland	30.5.2006
EU/1/00/169/002/NO-004/NO	Metalyse	Norway	14.3.2006
EU/1/00/169/004-006	Metalyse	Liechtenstein	30.4.2006
EU/1/00/169/004-006/IS	Metalyse	Iceland	11.4.2006

EU-Number	Product	Country	Date of renewal
EU/1/00/170/001/NO-002/NO	Fasturtec	Norway	27.2.2006
EU/1/00/170/001-002	Fasturtec	Liechtenstein	28.2.2006
EU/1/00/170/001-002(IS	Fasturtec	Iceland	7.3.2006
EU/1/01/171/001,007-012/IS	Rapamune	Iceland	8.6.2006
EU/1/01/171/001/NO, 007/NO-012/NO	Rapamune	Norway	5.5.2006
EU/1/01/172/001/NO-003/NO	Kaletra	Norway	26.4.2006
EU/1/01/172/001-003	Kaletra	Liechtenstein	30.4.2006
EU/1/01/172/001-003/IS	Kaletra	Iceland	23.5.2006
EU/1/01/173/001/NO-003/NO	Vaniqa	Norway	9.5.2006
EU/1/01/173/001-003	Vaniqa	Liechtenstein	30.4.2006
EU/1/01/173/001-003/IS	Vaniqa	Iceland	7.6.2006
EU/1/01/174/001/NO-007/NO	Starlix	Norway	23.5.2006
EU/1/01/174/001-021	Starlix	Liechtenstein	30.6.2006
EU/1/01/174/001-021/IS	Starlix	Iceland	7.6.2006
EU/1/01/175/001, 004-007, 008, 011-014, 015, 018-021/IS	Trazec	Iceland	7.6.2006
EU/1/01/175/001, 004-008, 011-015, 018-021	Trazec	Liechtenstein	30.6.2006
EU/1/01/175/001/NO, 004/NO-007/NO, 008/NO, 011/NO-014/NO, 015/NO, 018/NO-021/NO	Trazec	Norway	23.5.2006
EU/1/01/176/001/NO-003/NO	Zometa	Norway	2.5.2006
EU/1/01/176/001-006	Zometa	Liechtenstein	30.6.2006
EU/1/01/176/001-006/IS	Zometa	Iceland	22.5.2006
EU/1/01/177/001/NO-002/NO	SonoVue	Norway	4.5.2006
EU/1/01/177/001-002	SonoVue	Liechtenstein	30.4.2006
EU/1/01/177/002/IS	SonoVue	Iceland	8.6.2006
EU/1/01/178/001/IS	Targretin	Iceland	9.6.2006
EU/1/01/178/001/NO	Targretin	Norway	16.5.2006
EU/1/01/179/001	Osigraft	Liechtenstein	30.6.2006
EU/1/01/179/001/NO	Osigraft	Norway	30.5.2006
EU/1/01/184/001/NO-056/NO	Nespo	Norway	26.6.2006
EU/1/01/184/001-056	Nespo	Liechtenstein	30.6.2006
EU/1/01/185/001/NO-056/NO	Aranesp	Norway	26.6.2006
EU/1/01/185/001-056	Aranesp	Liechtenstein	30.6.2006
EU/1/01/187/001	DepoCyte	Liechtenstein	30.6.2006
EU/1/01/195/001/NO-015 /NO	Liprolog	Norway	14.3.2006

EU-Number	Product	Country	Date of renewal
EU/1/01/195/001-015	Liprolog	Liechtenstein	28.2.2006
EU/1/01/195/001-015/IS	Liprolog	Iceland	13.3.2006
EU/1/03/262/007-008/IS	Emend	Iceland	19.6.2006
EU/1/95/001/001, 003-005, 009, 012, 021-022, 025-028, 031-035	Gonal-f	Liechtenstein	31.1.2006
EU/1/95/002/001/NO-002/NO	Taxotere	Norway	23.2.2006
EU/1/95/002/001-002	Taxotere	Liechtenstein	31.1.2006
EU/1/95/002/001-002/IS	Taxotere	Iceland	23.2.2006
EU/1/95/003/003/NO-004/NO	Betaferon	Norway	2.3.2006
EU/1/95/003/003-004	Betaferon	Liechtenstein	28.2.2006
EU/1/95/003/003-004/IS	Betaferon	Iceland	10.4.2006
EU/1/96/004/001/NO-002/NO	Fareston	Norway	1.3.2006
EU/1/96/004/001-002	Fareston	Liechtenstein	28.2.2006
EU/1/96/005/001/NO-006/NO	CellCept	Norway	6.4.2006
EU/1/96/005/001-006	CellCept	Liechtenstein	30.4.2006
EU/1/96/005/001-006/IS	CellCept	Iceland	5.5.2006
EU/1/96/006/001/NO-003/NO	Novoseven	Norway	3.3.2006
EU/1/96/006/001-003	NovoSeven	Liechtenstein	28.2.2006
EU/1/96/006/001-003/IS	NovoSeven	Iceland	10.4.2006
EU/1/96/007/002, 004-006, 008, 010-011, 015-021, 023-030	Humalog	Liechtenstein	30.4.2006
EU/1/96/007/002, 004-006, 008, 010-011, 015-021, 023-030/IS	Humalog	Iceland	13.3.2006
EU/1/96/007/002/NO, 004/NO-006/NO, 008/NO, 010/NO-011/NO, 015/NO-021/NO, 023/NO-030/NO	Humalog	Norway	3.3.2006
EU/1/96/008/001/NO-040/NO	Puregon	Norway	26.6.2006
EU/1/96/008/001-041	Puregon	Liechtenstein	30.6.2006
EU/1/96/009/001/NO-002/NO	Zerit	Norway	2.5.2006
EU/1/96/009/001-009	Zerit	Liechtenstein	30.4.2006
EU/1/96/009/001-009/IS	Zerit	Iceland	24.5.2006
EU/1/96/011/001/NO-004/NO	Caelyx	Norway	26.5.2006
EU/1/96/011/001-004	Caelyx	Liechtenstein	30.6.2006
EU/1/96/011/001-004/IS	Caelyx	Iceland	16.6.2006
EU/1/98/093/002/IS	Forcaltonin	Iceland	31.5.2006
EU/1/98/093/002/NO	Forcaltonin	Norway	21.6.2006
EU/2/00/019/001-003/IS	Purevax FeLV	Iceland	9.3.2006

EU-Number	Product	Country	Date of renewal
EU/2/00/025/001/NO-004/NO	Advasure	Norway	15.3.2006
EU/2/00/026/001/NO-004/NO	Porcilis AR-T DF	Norway	10.3.2006
EU/2/00/026/001-004	Porcilis AR-T DF	Liechtenstein	28.2.2006
EU/2/00/026/001-004/IS	Porcilis AR-T DF	Iceland	8.3.2006
EU/2/00/027/001/NO-003/NO	Pirsue	Norway	10.3.2006
EU/2/00/027/001-003	Pirsue	Liechtenstein	28.2.2006
EU/2/00/028/002/NO-008 /NO	Zubrin	Norway	7.4.2006
EU/2/00/028/002-008	Zubrin	Liechtenstein	30.4.2006
EU/2/00/028/002-008/IS	Zubrin	Iceland	11.4.2006
EU/2/01/029/001/NO-003/NO	Eurican Herpes	Norway	18.5.2006
EU/2/01/029/001-003	Eurican Herpes	Liechtenstein	30.4.2006
EU/2/01/029/001-003/IS	Eurican Herpes	Iceland	8.6.2006
EU/2/96/001/001-010	Porcilis Porcoli	Liechtenstein	30.4.2006
EU/2/96/001/NO-010/NO	Porcilis Porcoli	Norway	5.5.2006

ANNEX III

List of extended marketing authorisations

The following marketing authorisations have been extended in the EEA EFTA States during the period 1 January-30 June 2006:

EU-Number	Product	Country	Date of extention
EU/1/00/137/013-018	Avandia	Liechtenstein	30.6.2006
EU/1/00/146/030	Keppra	Liechtenstein	30.4.2006
EU/1/00/146/030/IS	Keppra	Iceland	7.4.2006
EU/1/00/146/030/NO	Keppra	Norway	28.4.2006
EU/1/00/150/016-024	Actos	Liechtenstein	30.4.2006
EU/1/00/151/014-022	Glustin	Liechtenstein	30.4.2006
EU/1/00/162/019-021	Prandin	Liechtenstein	30.6.2006
EU/1/03/255/004-005	Ventavis	Liechtenstein	30.6.2006
EU/1/03/262/007/NO-008/NO	Emend	Norway	28.6.2006
EU/1/03/262/007-008	Emend	Liechtenstein	30.6.2006
EU/1/03/262/007-008/IS	Emend	Iceland	19.6.2006
EU/1/03/265/005/NO-006/NO	Bonviva	Norway	2.5.2006
EU/1/03/265/005-006/IS	Bonviva	Iceland	12.6.2006
EU/1/03/266/005/NO-006/NO	Bondenza	Norway	8.5.2006
EU/1/04/279/033/NO-035/NO	Lyrica	Norway	19.6.2006
EU/1/04/279/033-035	Lyrica	Liechtenstein	30.6.2006
EU/1/04/296/007	Cymbalta	Liechtenstein	30.4.2006
EU/1/04/296/008	Cymbalta	Liechtenstein	30.6.2006
EU/1/04/297/007	Xeristar	Liechtenstein	30.4.2006
EU/1/04/297/008	Xeristar	Liechtenstein	30.6.2006
EU/1/04/302/004	Prialt	Liechtenstein	30.4.2006
EU/1/04/302/004/IS	Prialt	Iceland	22.4.2006
EU/1/04/302/004/NO	Prialt	Norway	29.3.2006
EU/1/05/319/003-004	Xolair	Liechtenstein	30.6.2006
EU/1/96/007/029-030	Humalog	Liechtenstein	30.4.2006
EU/1/98/063/007/IS	Rebif	Iceland	30.5.2006
EU/1/98/063/007/NO	Rebif	Norway	16.2.2006
EU/1/98/076/022-024	NovoNorm	Liechtenstein	30.6.2006
EU/1/98/089/017-019	Pritor	Liechtenstein	30.4.2006
EU/1/99/107/005	Rebetol	Liechtenstein	30.4.2006

EU-Number	Product	Country	Date of extention
EU/2/00/019/005-007/IS	Purevax FeL V	Iceland	9.3.2006
EU/2/02/035/007	SevoFlo	Liechtenstein	30.4.2006
EU/2/03/037/005/NO	Proteqflu	Norway	15.3.2006
EU/2/03/038/005/NO	Proteqflu-Te	Norway	15.3.2006
EU/2/04/044/002/NO	Aivlosin	Norway	4.3.2006
EU/2/04/044/003-005/IS	Aivlosin	Iceland	22.3.2006
EU/2/97/004/016/NO-023/NO	Metacam	Norway	22.4.2006
EU/2/97/004/016-023	Metacam	Liechtenstein	30.4.2006
EU/2/97/004/016-023/IS	Metacam	Iceland	7.4.2006

ANNEX IV

List of withdrawn marketing authorisations

The following marketing authorisations have been withdrawn in the EEA EFTA States during the period 1 January-30 June 2006:

EU-Number	Product	Country	Date of withdrawal
EU/1/00/137/001	Avandia	Liechtenstein	30.6.2006
EU/1/01/183/002-003, 006, 009-010, 012, 014, 016-17	HBVAXPRO	Liechtenstein	28.2.2006
EU/1/02/245/001	Theryttrex	Liechtenstein	28.2.2006
EU/1/02/245/001/IS	Theryttrex	Iceland	10.2.2006
EU/1/06/024/006, 009	Crixivan	Liechstenstein	31.1.2006
EU/1/96/022/001, 003, 005, 007, 013, 015	Zyprexa	Liechstenstein	31.1.2006
EU/1/97/052/001/NO-006/NO, 009/NO-010/NO	Daquiran	Norway	22.2.2006
EU/1/97/052/001-006, 009-010	Daquiran	Liechtenstein	28.2.2006
EU/1/97/052/001-010/IS	Daquiran	Iceland	10.2.2006
EU/1/98/075/001-002	Fortovase	Liechtenstein	30.6.2006
EU/1/98/075/001-002/IS	Fortovase- Saquinavir	Iceland	26.6.2006
EU/1/98/087/001/NO-003/NO	Infergen	Norway	30.5.2006
EU/1/98/087/001-003	Infergen	Liechtenstein	30.6.2006
EU/1/98/087/001-003/IS	Infergen	Iceland	16.6.2006
EU/2/00/028/001	Zubrin	Liechtenstein	28.2.2006
EU/2/98/010/001-003, 015-016, 019-020	Econor	Liechtenstein	31.1.2006

ANNEX V

List of suspended marketing authorisations

The following marketing authorisations have been suspended in the EEA EFTA States during the period 1 January-30 June 2006:

EU-Number	Product	Country	Date of suspention
EU/1/02/239/001-030	Bextra	Liechtenstein	31.1.2006

Medicinal products — List of marketing authorisations granted by the EEA EFTA States for the second half of 2006

(2007/C 266/06)

Subcommittee I — On the free movement of goods

With reference to EEA Joint Committee Decision No 74/1999 of 28 May 1999, the EEA Joint Committee is invited to note, at the meeting on 8 December 2006, the following lists concerning marketing authorisations for medicinal products for the period 1 July-31 December 2006:

ANNEX I List of new marketing authorisations

ANNEX II List of renewed marketing authorisations

ANNEX III List of extended marketing authorisations

ANNEX IV List of withdrawn marketing authorisations

ANNEX I

List of new marketing authorisations

The following marketing authorisations have been granted in the EEA EFTA States during the period 1 July-31 December 2006:

EU-Number	Product	Country	Date of authorisation
EU/1/06/333/001-003/IS	Myozyme	Iceland	10.8.2006
EU/1/06/342/001	Nexavar	Liechtenstein	31.8.2006
EU/1/06/342/001/IS	Nexavar	Iceland	4.8.2006
EU/1/06/342/001/NO	Nexavar	Norway	28.7.2006
EU/1/06/343/001/NO-005/NO	Baraclude	Norway	25.7.2006
EU/1/06/343/001-006/IS	Baraclude	Iceland	24.7.2006
EU/1/06/344/001-009/IS	Acomplia	Iceland	19.7.2006
EU/1/06/345/001-009/IS	Zimulti	Iceland	19.7.2006
EU/1/06/346/001	Tysabri	Liechtenstein	31.8.2006
EU/1/06/346/001/IS	Tysabri	Iceland	24.7.2006
EU/1/06/346/001/NO	Tysabri	Norway	13.7.2006
EU/1/06/347/001/NO-003/NO	Sutent	Norway	27.7.2006
EU/1/06/347/001-003	Sutent	Liechtenstein	31.8.2006
EU/1/06/347/001-003/IS	Sutent	Iceland	3.8.2006
EU/1/06/348/001/NO-002/NO	RotaTeq	Norway	12.7.2006
EU/1/06/348/001-002	RotaTeq	Liechtenstein	31.8.2006
EU/1/06/348/001-002/IS	RotaTeq	Iceland	26.7.2006
EU/1/06/349/001/NO-008/NO	Avaglim	Norway	13.7.2006
EU/1/06/349/001-008	Avaglim	Liechtenstein	31.8.2006
EU/1/06/349/001-008/IS	Avaglim	Iceland	25.7.2006
EU/1/06/350/001	Savene	Liechtenstein	31.8.2006
EU/1/06/350/001/IS	Savene	Iceland	18.8.2006
EU/1/06/350/001/NO	Savene	Norway	16.8.2006
EU/1/06/351/001/NO-003/NO	Livensa	Norway	6.9.2006
EU/1/06/351/001-003	Livensa	Liechtenstein	31.8.2006
EU/1/06/351/001-003/IS	Livensa	Iceland	28.8.2006
EU/1/06/352/001/NO-003/NO	Intrinsa	Norway	25.8.2006

EU-Number	Product	Country	Date of authorisation
EU/1/06/352/001-003	Intrinsa	Liechtenstein	31.8.2006
EU/1/06/352/001-003/IS	Intrinsa	Iceland	18.8.2006
EU/1/06/353/001/NO-005/NO	Thelin	Norway	1.9.2006
EU/1/06/353/001-005	Thelin	Liechtenstein	31.10.2006
EU/1/06/353/001-005/IS	Thelin	Iceland	8.9.2006
EU/1/06/354/001/NO-009/NO	Competact	Norway	22.8.2006
EU/1/06/354/001-009	Competact	Liechtenstein	31.8.2006
EU/1/06/354/001-009/IS	Competact	Iceland	28.8.2006
EU/1/06/355/001/NO-003/NO	Atryn	Norway	21.8.2006
EU/1/06/355/001-003	Atryn	Liechtenstein	31.8.2006
EU/1/06/355/001-003/IS	Atryn	Iceland	17.8.2006
EU/1/06/356/001/NO-006/NO	Exjade	Norway	20.9.2006
EU/1/06/356/001-006	Exjade	Liechtenstein	31.10.2006
EU/1/06/356/001-006/IS	Exjade	Iceland	21.9.2006
EU/1/06/357/001/NO-017/NO	Gardasil	Norway	28.9.2006
EU/1/06/357/001-017	Gardasil	Liechtenstein	31.10.2006
EU/1/06/357/001-017/IS	Gardasil	Iceland	17.10.2006
EU/1/06/358/001/NO-017/NO	Silgard	Norway	29.9.2006
EU/1/06/358/001-017	Silgard	Liechtenstein	31.10.2006
EU/1/06/358/001-017/IS	Silgard	Iceland	19.10.2006
EU/1/06/359/001/NO-004/NO	Suboxone	Norway	26.10.2006
EU/1/06/359/001-004	Suboxone	Liechtenstein	31.10.2006
EU/1/06/359/001-004/IS	Suboxone	Iceland	24.10.2006
EU/1/06/360/001/NO-010/NO	Champix	Norway	20.10.2006
EU/1/06/360/001-010/IS	Champix	Iceland	24.10.2006
EU/1/06/361/001/IS	Luminity	Iceland	20.10.2006
EU/1/06/362/001/NO-004/NO	Byetta	Norway	13.12.2006

EU-Number	Product	Country	Date of authorisation
EU/1/06/362/001-004	Byetta	Liechtenstein	31.12.2006
EU/1/06/362/001-004/IS	Byetta	Iceland	19.12.2006
EU/1/06/363/001/NO-009/NO	Sprycel	Norway	20.12.2006
EU/1/06/363/001-009/IS	Sprycel	Iceland	15.12.2006
EU/2/02/036/001-002/IS	Nobilis OR inac	Iceland	3.8.2006
EU/2/06/059/001/IS	Convenia	Iceland	19.7.2006
EU/2/06/059/001/NO	Convenia	Norway	6.7.2006
EU/2/06/060/001/NO-002/NO	Poulvac FluFend H5N3	Norway	2.10.2006
EU/2/06/060/001-002	Poulvac Flufend H5N3 RG	Liechtenstein	31.10.2006
EU/2/06/060/001-002/IS	Poulvac Flufend H5N3 RG	Iceland	18.10.2006
EU/2/06/061/001/NO-004/NO	Nobilis Influenza H5N2	Norway	2.10.2006
EU/2/06/061/001-004	Nobilis Influenza H5N2	Liechtenstein	31.10.2006
EU/2/06/061/001-004/IS	Nobilis Influenza	Iceland	19.9.2006
EU/2/06/062/001/NO-005/NO	Cerenia	Norway	31.10.2006
EU/2/06/062/001-005	Cerenia	Liechtenstein	31.10.2006
EU/2/06/062/001-005/IS	Cerenia	Iceland	19.10.2006
EU/2/06/063/001/NO-003/NO	Yarvitan	Norway	14.12.2006
EU/2/06/063/001-003	Yarvitan	Liechtenstein	31.12.2006
EU/2/06/063/001-003/IS	Yarvitan	Iceland	7.12.2006

ANNEX II

List of renewed marketing authorisations

The following marketing authorisations have been renewed in the EEA EFTA States during the period 1 July-31 December 2006:

EU-Number	Product	Country	Date of renewal
EU/1/00/156/002-003/IS	Trizivir	Iceland	17.7.2006
EU/1/01/171/001, 007-012	Rapamune	Liechtenstein	31.8.2006
EU/1/01/179/001/IS	Osigraft	Iceland	4.9.2006
EU/1/01/183/001, 004-005, 007-008, 011, 013, 015, 018-029	HBVAXPRO	Liechtenstein	31.8.2006
EU/1/01/183/001/NO, 004/NO-005/NO, 007/NO-008/NO, 011/NO, 013/NO, 015/NO, 018/NO-029/NO	HBVAXPRO	Norway	1.9.2006
EU/1/01/184/001-056/IS	Nespo	Iceland	17.7.2006
EU/1/01/185/001-056/IS	Aranesp	Iceland	13.7.2006
EU/1/01/186/001/NO-002/NO	Nonafact	Norway	21.8.2006
EU/1/01/186/001-002	Nonafact	Liechtenstein	31.8.2006
EU/1/01/186/001-002/IS	Nonafact	Iceland	29.8.2006
EU/1/01/187/001/IS	Depocyte	Iceland	1.9.2006
EU/1/01/188/001/NO-006/NO	Fabrazyme	Norway	28.8.2006
EU/1/01/188/001-003/IS	Fabrazyme	Iceland	28.8.2006
EU/1/01/188/004-006	Fabrazyme	Liechtenstein	31.8.2006
EU/1/01/189/001-006	Replagal	Liechtenstein	31.8.2006
EU/1/01/189/001/IS	Replagal	Iceland	28.8.2006
EU/1/01/189/001/NO-006/NO	Replagal	Norway	28.8.2006
EU/1/01/190/001/NO-002/NO	Ceprotin	Norway	22.8.2006
EU/1/01/190/001-002	Ceprotin	Liechtenstein	31.8.2006
EU/1/01/190/001-002/IS	Ceprotin	Iceland	21.9.2006
EU/1/01/191/001/NO-005/NO	Ketek	Norway	16.10.2006
EU/1/01/191/001-005	Ketek	Liechtenstein	31.10.2006
EU/1/01/191/001-005/IS	Ketek	Iceland	18.9.2006
EU/1/01/192/001/NO-005/NO	Levviax	Norway	16.10.2006
EU/1/01/192/001-005	Levviax	Liechtenstein	31.10.2006
EU/1/01/192/001-005/IS	Levviax	Iceland	18.9.2006
EU/1/01/193/001/NO-002/NO	MabCampath	Norway	10.8.2006

EU-Number	Product	Country	Date of renewal
EU/1/01/193/001-002	MabCampath	Liechtenstein	31.8.2006
EU/1/01/193/001-002/IS	MabCampath	Iceland	28.8.2006
EU/1/01/194/001/NO-002/NO	INOmax	Norway	28.8.2006
EU/1/01/194/001-002	INOmax	Liechtenstein	31.8.2006
EU/1/01/194/001-002/IS	INOmax	Iceland	8.11.2006
EU/1/01/196/001/NO-003/NO	Cancidas	Norway	1.12.2006
EU/1/01/196/001-003	Cancidas	Liechtenstein	31.10.2006
EU/1/01/196/001-003/IS	Cancidas	Iceland	13.10.2006
EU/1/01/197/001/NO-002/NO	Foscan	Norway	11.12.2006
EU/1/01/197/001-002	Foscan	Liechtenstein	31.12.2006
EU/1/01/197/001-002/IS	Foscan	Iceland	1.12.2006
EU/1/01/198/001/NO-013/NO	Glivec	Norway	16.10.2006
EU/1/01/198/001-013	Glivec	Liechtenstein	31.10.2006
EU/1/01/198/001-013/IS	Glivec	Iceland	16.10.2006
EU/1/01/199/001/NO-002/NO	Travatan	Norway	30.10.2006
EU/1/01/199/001-002	Travatan	Liechtenstein	31.10.2006
EU/1/01/199/001-002/IS	Travatan	Iceland	9.11.2006
EU/1/02/201/001/NO-006/NO	Protopic	Norway	5.12.2006
EU/1/02/201/001-006	Protopic	Liechtenstein	31.12.2006
EU/1/02/202/001/NO-006/NO	Protopy	Norway	5.12.2006
EU/1/02/202/001-006	Protopy	Liechtenstein	31.12.2006
EU/1/06/015/001/NO-005/NO	Epivir	Norway	17.8.2006
EU/1/96/004/001-002/IS	Fareston	Iceland	18.9.2006
EU/1/96/008/001-041/IS	Puregon	Iceland	30.8.2006
EU/1/96/010/001	Rilutek	Liechtenstein	31.8.2006
EU/1/96/010/001/IS	Rilutek	Iceland	4.9.2006
EU/1/96/010/001/NO	Rilutek	Norway	4.9.2006
EU/1/96/012/001-013/IS	Bondronat	Iceland	28.8.2006
EU/1/96/012/004, 009-013	Bondronat	Liechtenstein	31.8.2006
EU/1/96/012/004/NO, 009/NO-010/NO, 011/NO-013/NO	Bondronat	Norway	24.7.2006
EU/1/96/014/001/NO-003/NO	Tritanrix HepB	Norway	21.8.2006
EU/1/96/014/001-003	Tritanrix HepB	Liechtenstein	31.8.2006
EU/1/96/015/001-005	Epivir	Liechtenstein	31.8.2006

EU-Number	Product	Country	Date of renewal
EU/1/96/015/001-005/IS	Epivir	Iceland	22.9.2006
EU/1/96/016/001, 003	Norvir	Liechtenstein	31.12.2006
EU/1/96/016/001, 003/IS	Norvir	Iceland	29.11.2006
EU/1/96/016/001/NO, 003/NO	Norvir	Norway	6.12.2006
EU/1/96/018/001	Rapilysin	Liechtenstein	31.8.2006
EU/1/96/018/001/IS	Rapilysin	Iceland	4.9.2006
EU/1/96/018/001/NO	Rapilysin	Norway	6.9.2006
EU/1/96/020/001/NO-009/NO	Twinrix Adult	Norway	25.9.2006
EU/1/96/020/001-009	Twinrix Adult	Liechtenstein	31.10.2006
EU/1/96/020/001-009/IS	Twinrix Adult	Iceland	16.10.2006
EU/1/96/022/002, 004, 006, 008-012, 014, 016-017, 019-022	Zyprexa	Liechtenstein	31.10.2006
EU/1/96/022/002, 004, 006, 008-012, 014, 016-017, 019-022/IS	Zyprexa	Iceland	29.9.2006
EU/1/96/022/002/NO, 004/NO, 006/NO, 008/NO-012/NO, 014/NO, 016/NO-017/NO, 019/NO-022/NO	Zyprexa	Norway	26.10.2006
EU/1/96/024/001/NO-005/NO, 007/NO-008/NO, 010/NO	Crixivan	Norway	7.12.2006
EU/1/96/024/001-005, 007-008, 010	Crixivan	Liechtenstein	31.12.2006
EU/1/96/024/001-005, 007-008, 010/IS	Crixivan	Iceland	1.12.2006
EU/1/96/026/001/NO-002/NO	Invirase	Norway	12.12.2006
EU/1/96/026/001-002	Invirase	Liechtenstein	31.12.2006
EU/1/96/026/001-002/IS	Invirase	Iceland	19.12.2006
EU/1/96/027/001, 003-005	Hycamtin	Liechtenstein	31.12.2006
EU/1/96/027/001/NO-005/NO	Hycamtin	Norway	4.12.2006
EU/1/97/029/001/NO-010/NO	Twinrix Paediatric	Norway	25.9.2006
EU/1/97/029/001-0010/IS	Twinrix Paediatric	Iceland	16.10.2006
EU/1/97/029/001-010	Twinrix Paediatric	Liechtenstein	31.10.2006
EU/1/99/125/001/NO-008/NO	Zyprexa Velotab	Norway	26.10.2006
EU/1/99/125/001-008	Zyprexa Velotab	Liechtenstein	31.10.2006
EU/1/99/125/001-008/IS	Zyprexa Velotab	Iceland	29.9.2006
EU/2/00/023/001-004/IS	Advasure	Iceland	7.11.2006
EU/2/00/027/001-003/IS	Pirsue	Iceland	7.11.2006
EU/2/01/030/001-004	Virbagen Omega	Liechtenstein	31.12.2006
EU/2/01/030/001-004/IS	Virbagen Omega	Iceland	4.12.2006
EU/2/96/001/001-002, 009-010/IS	Porcilis Porcoli	Iceland	3.10.2006

ANNEX III

List of extended marketing authorisations

The following marketing authorisations have been extended in the EEA EFTA States during the period 1 July-31 December 2006:

EU-Number	Product	Country	Date of extention
EU/02/215/011-012	PritorPlus	Liechtenstein	31.12.2006
EU/1/00/143/007-009	Kogenate	Liechtenstein	31.8.2006
EU/1/00/160/035-036	Aerius	Liechtenstein	31.10.2006
EU/1/01/172/004/NO-005/NO	Kaletra	Norway	11.7.2006
EU/1/01/172/004-005	Kaletra	Liechtenstein	31.8.2006
EU/1/01/172/004-005/IS	Kaletra	Iceland	12.7.2006
EU/1/01/184/057-068	Nespo	Liechtenstein	31.12.2006
EU/1/01/185/057-068	Aranesp	Liechtenstein	31.12.2006
EU/1/01/188/001-003	Fabrazyme	Liechtenstein	31.8.2006
EU/1/02/206/018-020	Arixtra	Liechtenstein	31.10.2006
EU/1/02/207/018-020	Quixidar	Liechtenstein	31.10.2006
EU/1/02/213/011-012	MicardisPlus	Liechtenstein	31.8.2006
EU/1/02/219/014-015	Ebixa	Liechtenstein	31.12.2006
EU/1/03/256/007-010	Humira	Liechtenstein	31.12.2006
EU/1/03/257/007-010	Trudexa	Liechtenstein	31.12.2006
EU/1/03/260/013-015	Stalevo	Liechtenstein	31.8.2006
EU/1/03/266/005-006/IS	Bondenza	Iceland	25.8.2006
EU/1/04/276/036	Abilify	Liechtenstein	31.10.2006
EU/1/04/276/036/NO	Abilify	Norway	21.12.2006
EU/1/04/279/033-035/IS	LYRICA	Iceland	21.7.2006
EU/1/04/285/029-036	Apidra	Liechtenstein	31.10.2006
EU/1/05/308/002	Aclasta	Liechtenstein	31.10.2006
EU/1/05/323/012-013	ProQuad	Liechtenstein	31.10.2006
EU/1/95/003/005-006	Betaferon	Liechtenstein	31.10.2006
EU/1/98/082/005	Comtess	Liechtenstein	31.8.2006
EU/1/98/085/023/NO-028/NO	Karvezide	Norway	22.9.2006
EU/1/98/085/023-028	Karvezide	Liechtenstein	31.10.2006
EU/1/98/085/023-028/IS	Karvezide	Iceland	18.9.2006
EU/1/98/086/023/NO-028/NO	CoAprovel	Norway	22.9.2006

EU-Number	Product	Country	Date of extention
EU/1/98/086/023-028	CoAprovel	Liechtenstein	31.10.2006
EU/1/98/086/023-028/IS	CoAprovel	Iceland	18.9.2006
EU/1/98/090/015-016	Micardis	Liechtenstein	31.8.2006
EU/1/99/111/010-011	Stocrin	Liechtenstein	31.12.2006
EU/1/99/111/010-011/IS	Stocrin	Iceland	15.12.2006
EU/1/99/126/012	Enbrel	Liechtenstein	31.8.2006
EU/1/99/126/012/NO	Enbrel	Norway	23.8.2006
EU/1/99/126/012-18/IS	Enbrel	Iceland	11.10.2006
EU/1/99/126/013/NO-018/NO	Enbrel	Norway	23.10.2006
EU/1/99/126/013-018	Enbrel	Liechtenstein	31.10.2006
EU/2/05/053/002	Naxcel	Liechtenstein	31.10.2006
EU/2/05/056/003-004	Equilis Prequenza	Liechtenstein	31.8.2006
EU/2/05/057/003-004	Equilis Prequenza Te	Liechtenstein	31.8.2006

ANNEX IV

List of withdrawn marketing authorisations

The following marketing authorisations have been withdrawn in the EEA EFTA States during the period 1 July-31 December 2006:

EU-Number	Product	Country	Date of withdrawal
EU/1/02/235/001-004	Monotard	Liechtenstein	31.12.2006
EU/1/02/235/001-004/IS	Monotard	Iceland	16.11.2006
EU/1/02/235/001-004/NO	Monotard	Norway	20.12.2006
EU/1/02/236/001-004	Ultratard	Liechtenstein	31.12.2006
EU/1/02/236/001-004/IS	Ultratard	Iceland	20.11.2006
EU/1/02/236/001-004/NO	Ultratard	Norway	13.12.2006
EU/1/98/075/001/NO-002/NO	Fortovase	Norway	6.7.2006
EU/1/98/075/001-002/IS	Fortovase	Iceland	4.7.2006

V

(Announcements)

OTHER ACTS

COMMISSION

Publication of an application pursuant to Article 8(2) of Council Regulation (EC) No 509/2006 on agricultural products and foodstuffs as traditional specialities guaranteed

(2007/C 266/07)

This publication confers the right to object to the application pursuant to Article 9 of Council Regulation (EC) No 509/2006 [1]. Statements of objection must reach the Commission within six months from the date of this publication.

APPLICATION FOR REGISTRATION OF A TSG

COUNCIL REGULATION (EC) No 509/2006

'CZWÓRNIAK'

EC No PL/TSG/007/0035/06.09.2006

1. **Name and address of the applicant group**

Name: Krajowa Rada Winiarstwa i Miodosytnictwa przy Stowarzyszeniu Naukowo — Technicznym Inżynierów i Techników Przemysłu Spożywczego

Address: ul. Czackiego 3/6
PL-00-043 Warszawa

Tel. (48) 22 828 27 21

E-mail: krwim@sitspoz.pl

2. **Member State or Third Country**

Poland

3. **Product specification**

3.1. *Name to be registered*

'Czwórniak'

When the product is placed on the market, the label may contain the following information: 'miód pitny wytworzony zgodnie ze staropolską tradycją' (mead produced in accordance with an old Polish tradition). This information should be translated into other official languages.

3.2. *Whether the name*

☒ is specific in itself

☒ expresses the specific character of the agricultural product or foodstuff

[1] OJ L 93, 31.3.2006, p. 1.

The name 'czwórniak' derives from the numeral '4' (PL: 'cztery') and relates directly to the historically established composition and method of production of 'czwórniak' — the proportions of honey and water in the mead wort being one part honey to three parts water. The name therefore expresses the specific character of the product. Since the term 'czwórniak' is a word that is used solely to denote a specific type of mead, the name should also be considered to be specific in itself.

3.3. *Whether reservation of the name is sought under Article 13(2) of Regulation (EC) No 509/200*

☐ Registration with reservation of the name

☒ Registration without reservation of the name

3.4. *Type of product*

Class 1.8 — Other products of Annex I

3.5. *Description of the agricultural product or foodstuff to which the name under point 3.1 applies*

'Czwórniak' is a mead, a clear beverage fermented from mead wort, distinguished by its characteristic honey aroma and the taste of the raw material used.

The flavour of 'czwórniak' may be enriched by the taste of spices that are used. The colour of 'czwórniak' ranges from golden to dark amber and depends on the type of honey used for production.

The physico-chemical indicators typical for 'czwórniak' mead are:

— alcohol content: 9-12 % vol.,

— reducing sugars after inversion: 35-90 g/l,

— total acidity expressed as malic acid: 3,5-8 g/l,

— volatile acidity expressed as acetic acid: max. 1,4 g/l,

— total sugar (g) plus actual alcohol content (% vol.) multiplied by 18: min. 240 g,

— non-sugar extract: not less than:

  — 15 g/l,

  — 20 g/l in the case of fruit mead (melomel),

— ash: min. 1,3 g/l — in the case of fruit mead.

The use of preservatives, stabilisers and artificial colourings and flavourings is prohibited in the production of 'czwórniak'.

3.6. *Description of the production method of the agricultural product or foodstuff to which the name under point 3.1 applies*

Raw materials

— Natural honey with the following parameters:

  — water content: max. 20 % (m/m),

  — reducing sugar content: min. 70 % (m/m),

  — combined sucrose and melezitose content: max. 5 % (m/m),

  — total acidity — 1 mol/l NaOH solution per 100 g of honey: within the range 1-5 ml,

  — 5-hydroxy-methyl-furfurol (HMF) content: max. 4,0 mg per 100 g honey.

— High-attenuation mead yeast — suitable for attenuation of high extracts in pitched wort.

— Herbs and spices: cloves, cinnamon, nutmeg or ginger.

— Natural fruit juices or fresh fruit.

Production method

Stage 1

Brewing (boiling) of the mead wort at a temperature of 95-105 °C. The required proportions of honey and water for 'czwórniak' are one part honey to three parts water (or water mixed with fruit juice), to which herbs or spices may be added. In the case of fruit meads, at least 30 % of the water is replaced with fruit juice.

Strict adherence to the proportions of water and honey and obtaining the required extract in a wort kettle fitted with a steam jacket. This method of brewing prevents caramelisation of the sugars.

Stage 2

Cooling of the wort to 20-22 °C, the optimum temperature for yeast to propagate. The wort must be cooled on the day of production, and the cooling time depends on the efficiency of the cooler. Cooling guarantees the microbiological safety of the wort.

Stage 3

Pitching — addition of a yeast solution to the wort in a fermentation tank.

Stage 4

A. Violent fermentation: 6-10 days. Keeping the temperature at a maximum level of 28 °C ensures that the fermentation process runs properly.

B. Still fermentation: 3-6 weeks. The still fermentation period ensures that the proper physico-chemical parameters are attained.

Stage 5

Racking of the attenuated pitched wort.

After obtaining an alcohol content of at least 9 % vol., racking prior to ageing should be carried out. This guarantees that the 'czwórniak' has the appropriate physico-chemical and organoleptic properties. Leaving the pitched wort on the lees beyond the still fermentation period adversely affects the organoleptic properties, owing to yeast autolysis.

Stage 6

Ageing (maturing) and siphoning (decanting) — this is repeated as necessary to prevent unwanted processes from taking place in the lees (yeast autolysis). During ageing it is possible to carry out operations such as pasteurisation and filtration.

This stage is essential for ensuring that the product has the right organoleptic properties.

The minimum ageing time for 'czwórniak' is nine months.

Stage 7

Flavour-adjustment (composition) — this stage concerns the preparation of a final product having the organoleptic and physico-chemical properties appropriate to 'czwórniak', as specified in point 3.5 — Description of the agricultural product or foodstuff. In order to ensure that the required parameters are attained, it is possible to correct the organoleptic and physico-chemical properties by:

— adding honey to sweeten the mead,

— adding herbs and spices.

The aim of this stage is to obtain a product with the characteristic 'czwórniak' bouquet.

Stage 8

Pouring into unit containers at a temperature of 55-60 °C. It is recommended that 'czwórniak' be presented in traditional packaging, such as: carboys, ceramic containers or oak barrels.

3.7. *Specific character of the agricultural product or foodstuff*

The specific character of 'czwórniak' results from:

— the preparation of the wort (composition and proportion of raw materials),

— ageing and maturing,

— its physico-chemical and organoleptic properties.

Preparation of the wort (composition and proportion of raw materials)

The specific character of 'czwórniak' results in particular from the use of, and strict adherence to, the established proportions of honey and water — one part honey to three parts water — in the mead wort. This proportion is the determining factor in all further stages in the production of 'czwórniak' that impart its unique properties.

Ageing and maturing

According to the traditional old Polish recipe, the character of the product depends on its being aged and matured for a specified period of time. In the case of 'czwórniak' this period is at least nine months.

Physico-chemical and organoleptic properties

Observance of all the stages of production included in the specification ensures that a product of unique taste and aroma is obtained. The unique taste and odour of 'czwórniak' is the result of appropriate sugar and alcohol content:

— reducing sugars after inversion: > 35-90 g/l,

— total sugar (g) plus actual alcohol content (% vol.) multiplied by 18: min. 240 g,

— alcohol: 9-12 % vol.

Owing to strictly defined proportions of the ingredients used in its production, 'czwórniak' possesses a typically viscous and runny consistency which distinguishes it from other types of mead.

3.8. *Traditional character of the agricultural product or foodstuff*

Traditional production method

Mead production in Poland is a tradition which dates back over a thousand years and is characterised by great diversity. The development and improvement of the production method over the centuries has given rise to many types of mead. The history of mead production dates back to the beginnings of Poland's statehood. In 966 the Spanish diplomat, merchant and traveller, Ibrahim ibn Yaqub, wrote: 'Besides food, meat and land for ploughing, the country of Mieszko I abounds in mead, which is what the Slavic wines and intoxicating drinks are called' (Mieszko I was the first historic king of Poland). The Chronicles of Gallus Anonymus, who recorded Polish history at the turn of the 11th and 12th centuries, also contain numerous references to the production of mead.

The Polish national epic poem 'Pan Tadeusz' by Adam Mickiewicz, which tells the story of the nobility between 1811 and 1812, contains a good deal of information on the production, consumption and different types of mead. Mentions of mead can also be found in the poems of Tomasz Zan (1796-1855) and in Henryk Sienkiewicz's trilogy describing events in Poland in the 17th century ('*Ogniem i mieczem*', published in 1884; '*Potop*', published in 1886 and '*Pan Wołodyjowski*', published in 1887 and 1888).

Source materials describing Polish culinary traditions of the 17th and 18th centuries contain not only general references to mead, but also references to different types of mead. Depending on the production method, they were called 'półtorak', 'dwójniak', 'trójniak' and 'czwórniak'. Each of these names relates to a different type of mead, produced on the basis of different proportions of honey and water or juice, and different ageing times. The 'czwórniak' production technique has been used, with minor modifications, for centuries.

Traditional composition

The traditional division of mead into 'półtorak', 'dwójniak', 'trójniak' and 'czwórniak' has existed in Poland for centuries and still exists in consumers' consciousness to this day. After the Second World War attempts were made to regulate the traditional division of mead into four categories. This division was finally enshrined in Polish law in 1948 by means of the Act on the production of wines, wine musts, meads and trade in such products (Journal of Laws of the Republic of Poland of 18 November 1948). This Act contains rules on the production of meads, specifying the proportions of honey and water and the technological requirements. The proportion of water and honey for 'czwórniak' is given as follows: 'Only mead produced from one part natural honey and three parts water may be called "czwórniak"'.

3.9. *Minimum requirements and procedures to check the specific character*

Mandatory checking encompasses:

— adherence to the established proportions of ingredients in the mead wort,

— adherence to the length of the ageing time,

— organoleptic properties of the finished product (taste, odour, colour, clarity),

— physico-chemical indicators of the finished product: alcohol content, total sugar, reducing sugar after inversion, total acidity, volatile acidity, non-sugar extract, and ash in the case of fruit meads — the values should correspond to the values specified at point 3.5 of the specification.

Mandatory checks are carried out at least once a year.

It is recommended that checks also be carried out during the production stages listed below. Checks at the production stages listed below are not mandatory, but are advisable, because they help eliminate possible errors occurring at different stages of production:

Stage 4

During the fermentation process, regular laboratory tests should be carried out on organoleptic properties (taste and odour) and physico-chemical parameters such as alcohol content and content of sugars that are subject to change during the alcoholic fermentation process.

Stage 6

During ageing, regular checks should be carried out on the basic organoleptic properties of the product and physico-chemical indicators such as alcohol content, total sugar, total acidity and volatile acidity.

Stage 8

Before bottling, checks are carried out on the various physico-chemical and organoleptic parameters specified at 3.5 — Description of the agricultural product or foodstuff.

4. **Authorities or bodies verifying compliance with the product specification**

4.1. *Name and address*

Name: Główny Inspektorat Jakości Handlowej Artykułów Rolno — Spożywczych

Address: ul. Wspólna 30
PL-00-930 Warszawa

Tel. (48) 22 623 29 00

Fax (48) 22 623 29 98

E-mail: —

☒ Public ☐ Private

4.2. *Specific tasks of the authority or body*

The inspection authority above is responsible for the verification of the entirety of the specification.

CORRIGENDA

Corrigendum to the Communication from the EFTA Surveillance Authority under Article 7 of the Act referred to at point 18 of Annex VII to the EEA Agreement (Council Directive 85/384/EEC of 10 June 1985 on the mutual recognition of diplomas, certificates and other evidence of formal qualifications in architecture, including measures to facilitate the effective exercise of the right of establishment and freedom to provide services)

(Official Journal of the European Union C 38 of 22 February 2007 and EEA Supplement No 8 of 22 February 2007)

(2007/C 266/08)

On page 17, in the third paragraph, second sentence, and fourth paragraph:

for: 'This new title is to be recognised by the contracting parties to the EEA Agreement in respect of those students who began their studies in architecture in the academic year 2001/2002.

The following designation is to be deleted from the list of titles of diplomas and the bodies awarding such diplomas for Norway:

— "Sivilarkitekt"

and replaced by the designation:

— "Master i arkitektur".',

read: 'This new title is to be recognised by the contracting parties to the EEA Agreement in respect of those students who began their studies in architecture in the academic year 1999/2000 at the *Norges teknisk-naturvitenskaplige universitet (NTNU)*, in the academic year 1998/1999 at the *Arkitektur- og designhøgskolen i Oslo (AHO)* (before 29 October 2004 *Arkitekthøgskolen i Oslo*) and in the academic year 2001/2002 at the *Bergen Arkitekt Skole (BAS)*.

The following designation is to be added to the list of titles of diplomas and the bodies awarding such diplomas for Norway:

— "Master i arkitektur".'