

## II

(Non-legislative acts)

## REGULATIONS

## COMMISSION IMPLEMENTING REGULATION (EU) 2021/1209

of 22 July 2021

**initiating ‘new exporter’ reviews of Implementing Regulation (EU) 2017/2230 imposing a definitive anti-dumping duty on imports of trichloroisocyanuric acid originating in the People’s Republic of China for three Chinese exporting producers, repealing the duty with regard to imports from these exporting producers and making these imports subject to registration**

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EU) 2016/1036 of the European Parliament and of the Council of 8 June 2016 on protection against dumped imports from countries not members of the European Union <sup>(1)</sup> (‘the basic Regulation’), and in particular Articles 11(4) and 14(5) thereof,

After having informed the Member States,

Whereas:

### 1. REQUEST

- (1) The Commission received three requests for a ‘new exporter’ review under Article 11(4) of the basic Regulation.
- (2) The requests were lodged by Hebei Xingfei Chemical Co., Ltd on 13 July 2020, by Inner Mongolia Likang Bio-Tech Co., Ltd (Likang) on 29 July 2019 and updated on 12 February 2021, and by Shandong Lantian Disinfection Technology Co., Ltd on 13 April 2021 (‘the applicants’), exporting producers of trichloroisocyanuric in the People’s Republic of China (‘the PRC’).

### 2. PRODUCT UNDER REVIEW

- (3) The product under review is trichloroisocyanuric acid and preparations thereof, also referred to as ‘symclosene’ under the international non-proprietary name (INN), currently falling under CN codes ex 2933 69 80 and ex 3808 94 20 (TARIC codes 2933 69 80 70 and 3808 94 20 20) and originating in the PRC.

### 3. EXISTING MEASURES

- (4) The measures currently in force are a definitive anti-dumping duty imposed by Commission Implementing Regulation (EU) 2017/2230 <sup>(2)</sup> under which imports of the product under review originating in the PRC, including the product produced by the applicant, are subject to a definitive anti-dumping duty of 42,6 % with the exception of several companies specifically mentioned in Article 1(2) of that Regulation which are subject to individual duty rates.

<sup>(1)</sup> OJ L 176, 30.6.2016, p. 21.

<sup>(2)</sup> Commission Implementing Regulation (EU) 2017/2230 of 4 December 2017 imposing a definitive anti-dumping duty on imports of trichloroisocyanuric acid originating in the People’s Republic of China following an expiry review pursuant to Article 11(2) of Regulation (EU) 2016/1036 of the European Parliament and of the Council (OJ L 319, 5.12.2017, p. 10).

#### 4. GROUNDS FOR THE REVIEWS

- (5) The applicants provided sufficient evidence that they did not export the product under review to the Union during the investigation period on which the anti-dumping measures were originally based (1 April 2003 to 31 March 2004).
- (6) The applicants provided sufficient evidence that they are not related to any of the exporting producer of the product under review which are subject to the anti-dumping duties in force.
- (7) Finally, the applicants provided sufficient evidence that they have begun exporting the product under review to the Union after the end of the original investigation period.

#### 5. PROCEDURE

##### 5.1. Initiation

- (8) The Commission examined the evidence available and concluded that there was sufficient evidence to justify the initiation of a 'new exporter' review pursuant to Article 11(4) of the basic Regulation, with a view to determining an individual margin of dumping for each of the applicants. Should dumping be found, the Commission will determine the level of the duty to which the imports of the product under review produced by each of the applicants should be subject.
- (9) In accordance with Articles 11(3) and 11(4) of the basic Regulation, normal value for the applicants shall be determined following the methodology laid down in Article 2(7) of the basic Regulation as it was in force on 19 December 2017 as the latest expiry review of the measures was initiated before 20 December 2017.
- (10) Union producers known to be concerned were informed of the request for a review on 8 April 2021 and were given an opportunity to comment until 23 April 2021.
- (11) The Commission also draws the attention of the parties that further to the COVID-19 outbreak a Notice <sup>(3)</sup> has been published on the consequences of the COVID-19 outbreak on anti-dumping and anti-subsidy investigations that may be applicable to this proceeding.

##### 5.2. Repeal of the existing measures and registration of imports

- (12) Pursuant to Article 11(4) of the basic Regulation, the anti-dumping duty in force should be repealed with regard to imports of the product under review produced by the applicants. At the same time, such imports should be made subject to registration in accordance with Article 14(5) of the basic Regulation, in order to ensure that anti-dumping duties can be levied from the date of the registration of these imports should the review result in a finding of dumping in respect of each of the applicants. Furthermore, the Commission notes that it is not possible, at this stage, to provide a reliable estimate of the amount of possible future liability, without prejudice to Article 9(4) of the basic Regulation.

##### 5.3. Review investigation period

- (13) In view of the small number of transactions on file and in order to make representative findings, the investigation will cover the period from 1 January 2019 to 30 June 2021 ('review investigation period').

##### 5.4. Investigating the applicants

- (14) In order to obtain information it deems necessary for its investigation, the Commission has made a questionnaire for the applicants available in the file for inspection by interested parties and on the website of the Directorate-General for Trade <https://trade.ec.europa.eu/tdi/>. The applicants must submit the completed questionnaire within the time limit specified in Article 4(2) of this Regulation.

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<sup>(3)</sup> <https://eur-lex.europa.eu/legal-content/EN/TXT/?uri=CELEX%3A52020XC0316%2802%29>

### 5.5. Other written submissions

- (15) Subject to the provisions of this Regulation, all interested parties are invited to make their views known, submit information and provide supporting evidence. Unless otherwise specified, this information and supporting evidence must reach the Commission within the time-limit specified in Article 4(2) of this Regulation.

### 5.6. Possibility to be heard by the Commission investigation services

- (16) All interested parties may request to be heard by the Commission investigation services within the time-limits specified in Article 4(3) of this Regulation. Any request to be heard must be made in writing and must specify the reasons for the request. For hearings on issues pertaining to the initiation stage of the investigation the request must be submitted within 15 days of the date of entry into force of this Regulation. Thereafter, a request to be heard must be submitted within the specific deadlines set by the Commission in its communication with the parties.

### 5.7. Instructions for making written submissions and sending completed questionnaires and correspondence

- (17) Information submitted to the Commission for the purpose of trade defence investigations shall be free from copyrights. Parties, before submitting to the Commission information and/or data which is subject to third party copyrights, must request specific permission to the copyright holder explicitly allowing a) the Commission to use the information and data for the purpose of this trade defence proceeding and b) to provide the information and/or data to interested parties to this investigation in a form that allows them to exercise their rights of defence.
- (18) All written submissions, including the information requested in this Regulation, completed questionnaires and correspondence provided by interested parties for which confidential treatment is requested shall be labelled 'Sensitive' <sup>(4)</sup>. Interested parties submitting information in the course of this investigation are invited to reason their request for confidential treatment.
- (19) Parties providing 'Sensitive' information are required to furnish non-confidential summaries of it pursuant to Article 19(2) of the basic Regulation, which will be labelled 'For inspection by interested parties'. Those summaries should be sufficiently detailed to permit a reasonable understanding of the substance of the information submitted in confidence.
- (20) If a party providing confidential information fails to show good cause for a confidential treatment request or does not furnish a non-confidential summary of it in the requested format and quality, the Commission may disregard such information unless it can be satisfactorily demonstrated from appropriate sources that the information is correct.
- (21) Interested parties are invited to make all submissions and requests via TRON.tdi (<https://webgate.ec.europa.eu/tron/TDI>) including scanned powers of attorney and certification sheets.
- (22) In order to have access to TRON.tdi, interested parties need an EU Login account. Full instructions on how to register and use TRON.tdi are available on <https://webgate.ec.europa.eu/tron/resources/documents/gettingStarted.pdf>
- (23) By using TRON.tdi or email, interested parties express their agreement with the rules applicable to electronic submissions contained in the document 'CORRESPONDENCE WITH THE EUROPEAN COMMISSION IN TRADE DEFENCE CASES' published on the website of the Directorate-General for Trade: [http://trade.ec.europa.eu/doclib/docs/2011/june/tradoc\\_148003.pdf](http://trade.ec.europa.eu/doclib/docs/2011/june/tradoc_148003.pdf)

<sup>(4)</sup> A 'Sensitive' document is a document which is considered confidential pursuant to Article 19 of the basic Regulation and Article 6 of the WTO Agreement on Implementation of Article VI of the GATT 1994 (Anti-Dumping Agreement). It is also a document protected pursuant to Article 4 of Regulation (EC) No 1049/2001 of the European Parliament and of the Council (OJ L 145, 31.5.2001, p. 43).

- (24) The interested parties must indicate their name, address, telephone and a valid email address and they should ensure that the provided email address is a functioning official business email which is checked on a daily basis. Once contact details are provided, the Commission will communicate with interested parties by TRON.tdi or email only, unless they explicitly request to receive all documents from the Commission by another means of communication or unless the nature of the document to be sent requires the use of a registered mail. For further rules and information concerning correspondence with the Commission including principles that apply to submissions including principles that apply to submissions via TRON.tdi and by email, interested parties should consult the communication instructions with interested parties referred to above.

Commission address for correspondence:

European Commission  
Directorate-General for Trade  
Directorate G  
Office: CHAR 04/039  
1049 Bruxelles/Brussel  
BELGIQUE/BELGIË

TRON.tdi: <https://webgate.ec.europa.eu/tron/tdi>

Email: TRADE-R746-TCCA@ec.europa.eu

## 6. NON-COOPERATION

- (25) If any interested party refuses access to or does not provide the necessary information within the time limits, or significantly impedes the investigation, findings, affirmative or negative, may be made on the basis of facts available, in accordance with Article 18 of the basic Regulation.
- (26) Where it is found that any interested party has supplied false or misleading information, the information shall be disregarded and use may be made of facts available in accordance with Article 18 of the basic Regulation.
- (27) If an interested party does not cooperate or cooperates only partially and findings are therefore based on facts available in accordance with Article 18 of the basic Regulation, the result may be less favourable to that party than if it had cooperated.

## 7. HEARING OFFICER

- (28) Interested parties may request the intervention of the Hearing Officer for trade proceedings. The Hearing Officer reviews requests for access to the file, disputes regarding the confidentiality of documents, requests for extension of time limits and any other request concerning the rights of defence of interested parties and third parties as may arise during the proceeding.
- (29) The Hearing Officer may organise hearings and mediate between the interested party/-ies and Commissions services to ensure that the interested parties' rights of defence are being fully exercised. A request for a hearing with the Hearing Officer should be made in writing and should specify the reasons for the request. The Hearing Officer will examine the reasons for the requests. These hearings should only take place if the issues have not been settled with the Commission services in the due course.
- (30) Any request must be submitted in good time and expeditiously so as not to jeopardise the orderly conduct of proceedings. To that effect, interested parties should request the intervention of the Hearing Officer at the earliest possible time following the occurrence of the event justifying such intervention. Where hearing requests are submitted outside the relevant timeframes, the Hearing Officer will also examine the reasons for such late requests, the nature of the issues raised and the impact of those issues on the rights of defence, having due regard to the interests of good administration and the timely completion of the investigation..

- (31) For further information and contact details interested parties may consult the Hearing Officer's web pages on DG Trade's website: <http://ec.europa.eu/trade/trade-policy-and-you/contacts/hearing-officer/>.

#### 8. SCHEDULE OF THE INVESTIGATION

- (32) The investigation will be concluded, pursuant to Article 11(5) of the basic Regulation, within nine months of the date of the entry into force of this Regulation.

#### 9. PROCESSING OF PERSONAL DATA

- (33) Any personal data collected in this investigation will be treated in accordance with Regulation (EU) 2018/1725 of the European Parliament and of the Council <sup>(5)</sup>.
- (34) A data protection notice that informs all individuals of the processing of personal data in the framework of Commission's trade defence activities is available on DG TRADE's website: <http://ec.europa.eu/trade/policy/accessing-markets/trade-defence/>

HAS ADOPTED THIS REGULATION:

#### *Article 1*

1. A review of Implementing Regulation (EU) 2017/2230 is hereby initiated under Article 11(4) of Regulation (EU) 2016/1036 in order to determine if an individual anti-dumping duty should be imposed on the imports of trichloroisocyanuric acid and preparations thereof, also referred to as 'symclosene' under the international non-proprietary name (INN), currently falling under CN codes ex 2933 69 80 and ex 3808 94 20 (TARIC codes 2933 69 80 70 and 3808 94 20 20), originating in the People's Republic of China ('the product under review'), produced for export to the Union by Inner Mongolia Likang Bio-Tech Co., Ltd (Likang) (TARIC additional code C630).

2. A review as referred to in paragraph 1 above is hereby also initiated concerning imports of the product under review produced for export to the Union by Hebei Xingfei Chemical Co., Ltd (TARIC additional code C629).

3. A review as referred to in paragraph 1 above is hereby also initiated concerning imports of the product under review produced for export to the Union by Shandong Lantian Disinfection Technology Co., Ltd (TARIC additional code C695).

#### *Article 2*

The anti-dumping duty imposed by Implementing Regulation (EU) 2017/2230 is hereby repealed with regard to the imports identified in Article 1 of this Regulation.

#### *Article 3*

The national customs authorities shall take the appropriate steps to register the imports identified in Article 1 of this Regulation, pursuant to Articles 11(4) and 14(5) of Regulation (EU) 2016/1036.

Registration shall expire nine months following the date of entry into force of this Regulation.

#### *Article 4*

1. Interested parties must make themselves known by contacting the Commission within 15 days from the date of entry into force of this Regulation.

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<sup>(5)</sup> Regulation (EU) 2018/1725 of the European Parliament and of the Council of 23 October 2018 on the protection of natural persons with regard to the processing of personal data by the Union institutions, bodies, offices and agencies and on the free movement of such data, and repealing Regulation (EC) No 45/2001 and Decision No 1247/2002/EC (OJ L 295, 21.11.2018, p. 39).

2. Interested parties, if their representations are to be taken into account during the investigation, must present their views in writing and submit questionnaire replies or any other information within 37 days from the date of the publication of this Regulation in the *Official Journal of the European Union*, unless otherwise specified.

3. Interested parties may also apply to be heard by the Commission within the same 37-day time limit. For hearings on issues pertaining to the initiation stage of the investigation the request must be submitted within 15 days of the date of entry into force of this Regulation. Any request to be heard must be made in writing and must specify the reasons for the request.

#### *Article 5*

This Regulation shall enter into force on the 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.

Done at Brussels, 22 July 2021.

*For the Commission*  
*The President*  
Ursula VON DER LEYEN

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