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**F. No. 6/14/2021-DGTR**  
**Government of India**  
**Department of Commerce**  
**Ministry of Commerce & Industry**  
**(Directorate General of Trade Remedies)**  
**Jeevan Tara Building, 5, Parliament Street, New Delhi – 110001**

**Dated 30.09.2021**

**INITIATION NOTIFICATION**

**(Case No. AD-OI-14/2021)**

**Subject: Initiation of anti-dumping investigation concerning imports of “Glycine” originating in or exported from China PR.**

Whereas M/s Paras Intermediates Private Limited and Avid Organics Private Limited (hereinafter referred to as the ‘Applicant’ or the ‘domestic industry’) has filed an application before the Designated Authority (hereinafter referred to as the “Authority”) on behalf of the domestic industry, in accordance with the Customs Tariff Act, 1975 as amended from time to time (hereinafter referred to as the “Act”) and the Customs Tariff (Identification, Assessment and Collection of Anti-Dumping Duty on Dumped Articles and for determination of injury) Rules, 1995 as amended from time to time (hereinafter referred to as the “Rules”) for Anti-dumping investigation concerning imports of “Ofloxacin and its intermediates” (hereinafter referred to as the “subject goods” or specifically as “product under consideration” or “PUC”), originating in or exported from China PR (hereinafter referred to as the “subject country”).

2. The Applicants have alleged dumping of the subject goods, originating in or exported from the subject country and consequent injury to the domestic industry and have requested for initiation of anti-dumping investigation and imposition of the anti-dumping duty on the imports of the subject goods, originating in or exported from the subject country.

**Domestic Industry and Standing**

3. The application has been filed jointly by M/s Paras Intermediates Private Limited and Avid Organics Private Limited. The Applicants have neither imported the subject goods during the period of investigation from the subject country nor is related to any exporter or producer of the subject goods in the subject country. Avid Organics Private Limited is not related to any importer of the product under consideration in India. M/s Paras Intermediates Private Limited is related to an importer in India that has made imports of the subject goods from the subject country during the POI, however, the same are not significant in quantity so as to affect the domestic market situation.

4. As per the evidence available on record, the production of the Applicants account for a major proportion of the total Indian production of the like article. In view of the above, the Authority notes that the Applicant constitutes eligible domestic industry in terms of Rule 2 (b), and the application satisfies the criteria of standing in terms of Rule 5(3) of the Rules supra.

#### **Product under Consideration**

5. The product under consideration in the present Application is "Glycine". Glycine's International Union of Pure and Applied Chemistry ("IUPAC") name is 2-aminoacetic acid. It is also known as Amino-acetic Acid, 2-aminoethanoic acid (IUPAC systematic), and lycocoll. The scope of this investigation covers "glycine" known by any name. Because glycine is produced at varying levels of purity or grades, it has a number of commercial names based on its level of purity or grade such as crude, industrial, technical-grade, feed-grade, food-grade, U.S. Pharmacopeia- or USP-grade, IP Grade, Food Grade, and pharmaceutical-grade. The scope of this investigation covers glycine of all grades. a

6. Glycine is a typically white material, odorless and sweet to the taste. In its dried form which is the form in which it is ordinarily sold, it is a free-flowing crystalline substance, like salt or sugar. All glycine has the same basic chemical structure, customers may distinguish between glycine of different purity levels for different applications, but neither glycine producers nor their customers perceive glycine that meets certain purity standards to be distinguishable, regardless of where it is produced.

7. Product under consideration falls under Chapter 29 of the Customs Tariff Act, 1975. Glycine comes under HS code 29224910. However, the PUC has been imported also under various other HS Codes namely 29224990.

#### **Like article**

8. The applicant has claimed that the goods produced by the domestic industry are identical to the subject goods exported from the subject country to India. The subject goods produced by the domestic industry are comparable to the imported goods from the subject country in terms of technical specifications, manufacturing process & technology, functions & uses, pricing, distribution & marketing and tariff classification of the goods. The two are technically and commercially substitutable and should be treated as 'like article' under the Rules. Therefore, for the purpose of the present investigation, the subject goods produced by the domestic industry are being treated as 'like article' to the subject goods being imported from the subject country.

### **Subject Country**

9. The subject country in the present investigation is China PR.

### **Period of Investigation**

10. The period of investigation (POI) for the present investigation is April 2020 – March 2021 (12 Months). The injury period of investigation will, however, cover the periods 2017 – 18, 2018 – 19, 2019 – 20 and the POI.

### **Normal Value**

11. The Applicant has cited and relied upon Article 15(a) (i) of China's Accession Protocol. The Applicant has claimed that producers in China PR must be asked to demonstrate that market economy conditions prevail in the industry producing the subject goods with regard to the manufacture, production and sale of the product under consideration. It has been stated by the Applicant that in case the responding Chinese Producers are not able to demonstrate that their costs and price information are market driven, the normal value should be calculated in terms of provisions of para 7 and 8 of Annexure I to the Rules. The normal value has accordingly been determined in terms of Para 7 of Annexure I to the Rules.

12. The applicants have proposed that for the purpose of initiation normal value may be taken from three of the options i.e., exports from Japan to USA or export from India to Canada or export from India to USA.

13. However, in absence of proper justification in support of the contention made by the applicants, the normal value for China PR has been constructed on the basis of cost of production of like article of the domestic industry and the selling, general and administrative expenses along with reasonable addition for profit.

### **Export Price**

14. The export price for the subject goods from the subject country has been computed based on the Directorate General of Commercial Intelligence and Statistics (DGCI&S) transaction-wise import data. Price adjustments have been made on account of ocean freight, inland freight, handling charges, marine insurance and bank charges.

### **Dumping Margin**

15. Considering the normal value and export price determined, dumping margin determined is above de-minimis level. There is *prima facie* evidence that normal value of the subject goods in the subject country is higher than the net export price, thereby indicating that the subject goods originating in or exported from the subject country have been exported at dumped prices.

### **Evidence of Injury and Causal Link**

16. Information furnished by the Applicant has been considered for assessment of injury to the domestic industry. The Applicant has claimed that the dumping of the product under consideration is causing material injury to the domestic industry. The imports are undercutting the prices of the domestic industry and causing price depression in the market. The performance of the domestic industry in respect of production, domestic sales, capacity utilization, market share, profits, return on capital employed, cash profits etc., have deteriorated. The overall performance of the domestic industry deteriorated in the POI. There is *prima facie* evidence of injury to the domestic industry by dumped imports from China PR.

### **Initiation of Anti-Dumping Investigation**

17. On the basis of the duly substantiated application by or on behalf of the domestic industry, and having satisfied itself, on the basis of the *prima facie* evidence submitted by the domestic industry, about dumping of the product under consideration originating in or exported from the subject country, injury to the domestic industry and causal link between such alleged dumping and injury, and in accordance with Section 9A of the Act read with Rule 5 of the Rules, the authority, hereby, initiates an investigation to determine the existence, degree and effect of any alleged dumping in respect of the product under consideration originating in or exported from the subject country and to recommend the amount of anti-dumping duty, which if levied, would be adequate to remove the injury to the domestic industry.

### **Procedure**

18. Principles as given in Rule 6 of the Rules will be followed for the present investigation.

### **Submission of information**

19. In view of the special circumstances arising out of COVID-19 pandemic, all communication should be sent to the Designated Authority via email at email addresses [adg12-dgtr@gov.in](mailto:adg12-dgtr@gov.in); [dir12-dgtr@gov.in](mailto:dir12-dgtr@gov.in) [jd12-dgtr@gov.in](mailto:jd12-dgtr@gov.in) and [jd14-dgtr@gov.in](mailto:jd14-dgtr@gov.in). It should be ensured that the narrative part of the submission is in searchable PDF/MS Word Format and data files are in MS Excel format.

20. The known producers/exporters in the subject country and their government through their Embassies in India, importers and users in India known to be concerned with the Product under consideration and the domestic industry are being informed separately to enable them to file all relevant information in the form and manner prescribed within the time-limit set out below.

21. Any other interested party may also make its submissions relevant to the investigation in the form and manner prescribed within the time-limit set out below.

22. Any party making any confidential submission before the Authority is required to make a non-confidential version of the same available to the other parties.

23. Interested parties are further advised to keep a regular watch on the official website of the Designated Authority <http://www.dgtr.gov.in/> for any updated information with respect to this investigation.

#### **Time-Limit**

24. Any information relating to the present investigation should be sent to the Designated Authority via email at the email addresses adg12-dgtr@gov.in; dir12-dgtr@gov.in jd12-dgtr@gov.in and jd14-dgtr@gov.in within thirty days from the date of receipt of the notice as per Rule 6(4) of the Anti-Dumping Rules. It may, however, be noted that in terms of explanation of the said Rule, the notice calling for information and other documents shall be deemed to have been received within one week from the date on which it was sent by the Designated Authority or transmitted to the appropriate diplomatic representative of the exporting country. If no information is received within the prescribed time limit or the information received is incomplete, the Authority may record its findings on the basis of the facts available on record in accordance with the Rules.

25. All the interested parties are hereby advised to intimate their interest (including the nature of interest) in the instant matter and file their questionnaire responses within the above time limit.

#### **Submission of information on confidential basis**

26. Any party making any confidential submission or providing information on confidential basis before the Authority, is required to simultaneously submit a non-confidential version of the same in terms of Rule 7(2) of the Rules and the Trade Notices issued in this regard. Failure to adhere to the above may lead to rejection of the response/submissions.

27. The parties making any submission (including Appendices/Annexures attached thereto), before the Authority including questionnaire response, are required to file Confidential and Non-Confidential versions separately.

28. The “confidential” or “non-confidential” submissions must be clearly marked as “confidential” or “non-confidential” at the top of each page. Any submission made without such marking shall be treated as non-confidential by the Authority, and the Authority shall be at liberty to allow the other interested parties to inspect such submissions.

29. The confidential version shall contain all information which is by nature confidential and/or other information which the supplier of such information claims as confidential. For information which are claimed to be confidential by nature or the information on which confidentiality is claimed because of other reasons, the supplier of the information is required to provide a good cause statement along with the supplied information as to why such information cannot be disclosed.

30. The non-confidential version is required to be a replica of the confidential version with the confidential information preferably indexed or blanked out (in case indexation is not feasible) and summarized depending upon the information on which confidentiality is claimed. The non-confidential summary must be in sufficient detail to permit a reasonable understanding of the substance of the information furnished on confidential basis. However, in exceptional circumstances, the party submitting the confidential information may indicate that such information is not susceptible to summary, and a statement of reasons why summarization is not possible must be provided to the satisfaction of the Authority.

31. The Authority may accept or reject the request for confidentiality on examination of the nature of the information submitted. If the Authority is satisfied that the request for confidentiality is not warranted or if the supplier of the information is either unwilling to make the information public or to authorize its disclosure in generalised or summary form, it may disregard such information.

32. Any submission made without a meaningful non-confidential version thereof or without good cause statement on the confidentiality claim shall not be taken on record by the Authority.

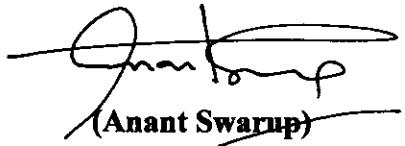
33. The Authority on being satisfied and accepting the need for confidentiality of the information provided, shall not disclose it to any party without specific authorization of the party providing such information.

#### **Sharing of responses/submission amongst interested parties**

34. A list of registered interested parties will be uploaded on DGTR's website along with the request therein to all of them to email the non-confidential version of their submissions to all other interested parties since the public file will not be accessible physically due to the ongoing global pandemic.

#### **Non-cooperation**

35. In case where an interested party refuses access to, or otherwise does not provide necessary information within a reasonable period, or significantly impedes the investigation, the Authority may declare such interested party as non-cooperative and record its findings on the basis of the facts available to it and make such recommendations to the Central Government as deemed fit.



(Anant Swarup)  
Designated Authority