application form and the forms associated with submitting new petitions to the Commission. Also included in the proposed generic clearance are the institution notices for the five-year reviews of antidumping and countervailing duty orders and suspended investigations. The Commission evaluates responses to the institution notices, which will form much of the record supporting the Commission's determinations to conduct either expedited or full five-year reviews of existing antidumping and countervailing duty orders.

(2) Information Collection Plan

The Commission sends questionnaires for specific investigations to all identified domestic producers of the product(s) in question subject to the Commission proceeding. The Commission also sends importer and purchaser questionnaires to all substantial U.S. importers and purchasers of the product(s). Further, the Commission sends questionnaires to all foreign manufacturers of the product(s) in question that are represented by counsel, and, in addition, it attempts to contact any other foreign manufacturers, especially if they export the product(s) in question to the United States. Firms receiving questionnaires include businesses, farms, and other for-profit institutions; responses by domestic firms are mandatory. The Commission publishes institution notices for the five-year reviews in the **Federal Register** and solicits comments from interested parties (*e.g.,* U.S. producers within the industry in question, as well as labor unions or representative groups of workers, U.S. importers and foreign exporters, and involved foreign country governments).

(3) Description of the Information To Be Collected

As it relates to import injury questionnaires, the content of each questionnaire will differ based on the needs of a particular investigation; questionnaires are based on long-established, generic formats, that align the data being gathered to the specific points of analysis that the statutes direct the Commission to analyze. Producer questionnaires generally consist of the following four parts: (part I) general questions relating to the organization and activities of the firm; (part II) data on capacity, production, inventories, employment, and the quantity and value of the firm's shipments and purchases from various sources; (part III) financial data, including income-and-loss data on the product in question, data on asset valuation, research and development expenses, and capital expenditures; and (part IV) pricing and market factors. Questionnaires may, on occasion, also contain additional parts depending on the facts of the case and the arguments raised by interested parties, the most frequent of which relate to information to assess proposed alternative definitions of the domestic like product.

Importer questionnaires generally consist of three parts: (part I) general questions relating to the organization and activities of the firm; (part II) data on the firm's imports and the shipment and inventories of its imports; and (part III) pricing and market factors similar to that requested in the domestic producer questionnaire. Purchaser questionnaires generally consist of four parts: (part I) general questions relating to the organization and activities of the firm; (part II) data concerning the purchases of the product by the firm and the names of the firm's vendors; (part III) market characteristics and purchasing practices; and (part IV) comparisons between imported and U.S.-produced product. The Commission may send an abbreviated purchaser questionnaire: (1) in a preliminary phase investigation, consisting of two parts: (part I) data concerning the purchases of the product by the firm; and (part II) questions regarding purchasing practices; or (2) in an adequacy phase of a review investigation, consisting of one part: (part I) general questions regarding the industry. Foreign producer questionnaires generally consist of: (part I) general questions relating to the organization and activities of the firm; (part II) data concerning the firm's manufacturing operations; and may include (part III) market factors. The notices of institution for the five-year reviews include 11 specific requests for information that firms are to provide if their response is to be considered by the Commission.

(4) Estimated Burden of the Proposed Information Collection

The Commission estimates that information collections issued under the requested generic clearance will impose an average annual burden of 409,050 hours on 12,935 respondents (*i.e.,* recipients that provide a response to the Commission's questionnaires, notices of institution of five-year reviews, and other investigations and forms).

(5) Minimization of Burden

The Commission periodically reviews its investigative processes, including data collection, to reduce the information burden. Questionnaires clearly state that reasonable estimates are acceptable for certain items. The questionnaires are designed in part with check-in type formats to simplify the response. The reporting burden is reduced by limiting data to a terminal year when a time series is not required. Moreover, the reporting burden for smaller firms is reduced in that the sections of the questionnaire that are applicable to their operations are typically more limited and, when pertinent, there are fewer requested data points. The Commission will not accept requests by parties to expand the data collection or add items to the questionnaire for specific investigations if it believes that such requests will increase the response burden without substantially adding to the investigative record. Respondents submit the information provided in response to the Commission's notices of institution for the five-year reviews electronically to the Commission's Electronic Data Information System (EDIS) and Electronic Docket. In addition, the Commission has reduced the information burden by streamlining the questionnaires. For example, the Commission removed redundant fields, added auto-calculated reconciliation fields, enabled population of whole data tables, and reduced the number of years for which data is collected in certain five-year reviews. In addition, the Commission ceased collecting nonsubject pricing data in preliminary proceedings.

No record keeping burden is known to result from the proposed collection of information.

By order of the Commission.

Issued: December 29, 2022.

Jessica Mullan,

Acting Supervisory Attorney.

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INTERNATIONAL TRADE COMMISSION

Notice of Receipt of Complaint; Solicitation of Comments Relating to the Public Interest

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has received a complaint entitled *Certain Selective Thyroid Hormone Receptor-Beta Agonists, Processes for Manufacturing or Relating to Same, and Products Containing Same, DN 3662;* the Commission is soliciting comments on any public

interest issues raised by the complaint or complainant's filing pursuant to the Commission's Rules of Practice and Procedure.

FOR FURTHER INFORMATION CONTACT: Katherine M. Hiner, Acting Secretary to the Commission, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205–2000. The public version of the complaint can be accessed on the Commission's Electronic Document Information System (EDIS) at *https://edis.usitc.gov.* For help accessing EDIS, please email *EDIS3Help@usitc.gov.* General information concerning the Commission may also be obtained by accessing its internet server at United States International Trade Commission (USITC) at *https://www.usitc.gov*. The public record for this investigation may be viewed on the Commission's Electronic Document Information System (EDIS) at *https://edis.usitc.gov.* Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810.

SUPPLEMENTARY INFORMATION: The Commission has received a complaint and a submission pursuant to § 210.8(b) of the Commission's Rules of Practice and Procedure filed on behalf of Viking Therapeutics, Inc. on December 29, 2022. The complaint alleges violations of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) in the importation into the United States, the sale for importation, and the sale within the United States after importation of regarding certain selective thyroid hormone receptor-beta agonists, processes for manufacturing or relating to same, and products containing same. The complainant names as respondents: Ascletis Pharma Inc. of China; Ascletis Pharmaceuticals Co. Ltd. of China; Ascletis Bioscience Co., Ltd. of China; Gannex Pharma Co., Ltd. of China; Jinzi Jason Wu of Seattle, WA. The complainant requests that the Commission issue a permanent exclusion order, a cease and desist order, and impose a bond upon respondent's alleged infringing articles during the 60-day Presidential review period pursuant to 19 U.S.C. 1337(j).

Proposed respondents, other interested parties, and members of the public are invited to file comments on any public interest issues raised by the complaint or § 210.8(b) filing. Comments should address whether issuance of the relief specifically requested by the complainant in this investigation would affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

(i) explain how the articles potentially subject to the requested remedial orders are used in the United States;

(ii) identify any public health, safety, or welfare concerns in the United States relating to the requested remedial orders;

(iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;

(iv) indicate whether complainant, complainant's licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to the requested exclusion order and/or a cease and desist order within a commercially reasonable time; and

(v) explain how the requested remedial orders would impact United States consumers.

Written submissions on the public interest must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the **Federal Register**. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation. Any written submissions on other issues must also be filed by no later than the close of business, eight calendar days after publication of this notice in the **Federal Register**. Complainant may file replies to any written submissions no later than three calendar days after the date on which any initial submissions were due. No other submissions will be accepted, unless requested by the Commission. Any submissions and replies filed in response to this Notice are limited to five (5) pages in length, inclusive of attachments.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. Submissions should refer to the docket number (''Docket No. 3662'') in a prominent place on the cover page and/or the first page. (*See* Handbook for Electronic Filing Procedures, Electronic Filing Procedures).[1] Please note the Secretary's Office will accept only electronic filings during this time. Filings must be made through the Commission's Electronic Document Information System (EDIS, *https://edis.usitc.gov.*) No in-person paper-based filings or paper copies of any electronic filings will be accepted until further notice. Persons with questions regarding filing should contact the Secretary at *EDIS3Help@usitc.gov.*

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. *See* 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for purposes of this Investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. appendix 3; or (ii) by U.S. government employees and contract personnel,[2] solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.[3]

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

By order of the Commission.

Issued: December 29, 2022.

Jessica Mullan,

Acting Supervisory Attorney.

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[1] Handbook for Electronic Filing Procedures: *https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf.*

[2] All contract personnel will sign appropriate nondisclosure agreements.

[3] Electronic Document Information System (EDIS): *https://edis.usitc.gov.*