service. Please address the extent to which wind turbine operators have already paid for components potentially covered by a remedy, and related service, through warranty and other contractual provisions. Please also address whether switching providers would cause delays or compatibility issues. Please explain how such additional costs, if any, would affect one or more of the four public interest considerations.

5. Please explain what products, if any, are still subject to the license agreement between the parties or whether SGRE otherwise retains the right under patent exhaustion principles to import components for the purpose of repairing products sold under the license. Please explain how the Commission or Customs and Border Protection could ascertain whether imported products are covered by the license or are otherwise authorized.

6. Please address whether SGRE's proven domestic inventories of products and components that are accused of infringing (a) claims 1, 6, and 12 of the '985 patent and (b) claims 29, 30, 33–35, and 37 of the '985 patent are commercially significant within an appropriate context and whether SGRE has other significant business operations in the United States. Please address the various product categories separately: Full-converter turbines using the earlier software, full-converter turbines using the later software, and DFIG turbines.

If the Commission orders some form of remedy, the U.S. Trade Representative, as delegated by the President, has 60 days to approve, disapprove, or take no position on the Commission's action. *See* Presidential Memorandum of July 21, 2005, 70 FR 43251 (July 26, 2005). During this period, the subject articles would be entitled to enter the United States under bond, in an amount determined by the Commission and prescribed by the Secretary of the Treasury. The Commission is therefore interested in receiving submissions concerning the amount of the bond that should be imposed if a remedy is ordered.

Written Submissions: The Commission requests that the parties to the investigation file written submissions on the remedy and public interest issues identified in this notice. The Commission encourages parties to the investigation, interested government agencies, and any other interested parties to file written submissions on the issues of remedy, the public interest, and bonding. Such submissions should address the recommended determination by the ALJ on remedy and bonding, which issued on September 10, 2021. The Commission further requests that GE submit proposed remedial orders, state the date when the '985 patent expires, provide the HTSUS subheadings under which the subject articles are imported, and supply a list of known importers of the subject article. The written submissions, exclusive of any exhibits, must not exceed 40 pages, and must be filed no later than close of business on December 7, 2021. Reply submissions must not exceed 20 pages, and must be filed no later than the close of business on December 14, 2021. No further submissions on these issues will be permitted unless otherwise ordered by the Commission.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. The Commission's paper filing requirements in 19 CFR 210.4(f) are currently waived. 85 FR 15798 (March 19, 2020). Submissions should refer to the investigation number (Inv. No. 337–TA–1218) in a prominent place on the cover page and/or the first page. (*See* Handbook for Electronic Filing Procedures, *https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf*). Persons with questions regarding filing should contact the Secretary, (202) 205–2000.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment by marking each document with a header indicating that the document contains confidential information. This marking will be deemed to satisfy the request procedure set forth in Rules 201.6(b) and 210.5(e)(2) (19 CFR 201.6(b) & 210.5(e)(2)). Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. A redacted non-confidential version of the document must also be filed simultaneously with any confidential filing. 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, solely for cybersecurity purposes. All contract personnel will sign appropriate nondisclosure agreements. All nonconfidential written submissions will be available for public inspection on EDIS.

The Commission vote for this determination took place on November 12, 2021.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: November 12, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021–25134 Filed 11–17–21; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–1070B (Third Review)]

Certain Tissue Paper Products From China

Determination

On the basis of the record [1] developed in the subject five-year review, the United States International Trade Commission (''Commission'') determines, pursuant to the Tariff Act of 1930 (''the Act''), that revocation of the antidumping duty order on certain tissue paper products from China would be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time.

Background

The Commission instituted this review on June 1, 2021 (86 FR 29289) and determined on September 7, 2021 that it would conduct an expedited review (86 FR 54238, September 30, 2021).

The Commission made this determination pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determination in this review on November 4, 2021. The views of the Commission are contained in USITC Publication 5236 (November 2021), entitled *Certain Tissue Paper Products from China: Investigation No. 731–TA–1070B (Third Review).*

By order of the Commission.

Issued: November 18, 2021.

Katherine Hiner,

Supervisory Attorney.

[FR Doc. 2021–25196 Filed 11–17–21; 8:45 am]

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[1] The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).