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*Other Information:* These meetings are open to the public and press on a first-come, first-served basis. Space is limited. To ensure an accurate headcount, all expected attendees are asked to provide notice of intent to attend by sending an email to [BoardRSVP@firstnet.gov](mailto:BoardRSVP@firstnet.gov). If the number of RSVPs indicates that expected attendance has reached its capacity, FirstNet will respond to all subsequent notices indicating that capacity has been reached and that in-person viewing may no longer be available but that the meeting may still be viewed by webcast as detailed below. For access to the meetings, valid government issued photo identification may be requested for security reasons.

The Combined Committee and Board Meetings are accessible to people with disabilities. Individuals requiring accommodations, such as sign language interpretation or other ancillary aids, are asked to notify Ms. Miller-Kuwana by telephone (571) 665-6177 or email at [Karen.Miller-Kuwana@firstnet.gov](mailto:Karen.Miller-Kuwana@firstnet.gov) at least five (5) business days before the applicable meeting.

The meeting will also be webcast. Please refer to FirstNet's website at [www.firstnet.gov](http://www.firstnet.gov) for webcast instructions and other information. Viewers experiencing any issues with the live webcast may email [support@sparkstreetdigital.com](mailto:support@sparkstreetdigital.com) or call 202-684-3361 x3 for support. A variety of automated troubleshooting tests are also available via the "Troubleshooting Tips" button on the webcast player. The meetings will also be available to interested parties by phone. To be connected to the meetings in listen-only mode by telephone, please dial toll free 1-888-469-2980 and enter participant code 4810197#. If you experience technical difficulty, please contact the Conferencing Center customer service at 1-866-900-1011.

*Records:* FirstNet maintains records of all Board proceedings. Minutes of the Board Meeting and the Committee meetings will be available at [www.firstnet.gov](http://www.firstnet.gov).

Dated: December 3, 2018.

**Karen Miller-Kuwana,**

*Board Secretary, First Responder Network Authority.*

[FR Doc. 2018-26600 Filed 12-10-18; 8:45 am]

**BILLING CODE 3510-TL-P**

## DEPARTMENT OF COMMERCE

### Foreign-Trade Zones Board

[S-218-2018]

#### Foreign-Trade Zone 24—Pittston, Pennsylvania; Application for Subzone; adidas America, Inc.; Wilkes-Barre, Pennsylvania

An application has been submitted to the Foreign-Trade Zones Board (the Board) by the Eastern Distribution Center, Inc., grantee of FTZ 24, requesting subzone status for the facility of adidas America, Inc., located in Wilkes-Barre, Pennsylvania. The application was submitted pursuant to the provisions of the Foreign-Trade Zones Act, as amended (19 U.S.C. 81a-81u), and the regulations of the Board (15 CFR part 400). It was formally docketed on December 4, 2018.

The proposed subzone (89.39 acres) is located at 550 New Commerce Blvd., Wilkes-Barre. No authorization for production activity has been requested at this time. The proposed subzone would be subject to the existing activation limit of FTZ 24.

In accordance with the Board's regulations, Elizabeth Whiteman of the FTZ Staff is designated examiner to review the application and make recommendations to the Executive Secretary.

Public comment is invited from interested parties. Submissions shall be addressed to the Board's Executive Secretary at the address below. The closing period for their receipt is January 22, 2019. Rebuttal comments in response to material submitted during the foregoing period may be submitted during the subsequent 15-day period to February 4, 2019.

A copy of the application will be available for public inspection at the Office of the Executive Secretary, Foreign-Trade Zones Board, Room 21013, U.S. Department of Commerce, 1401 Constitution Avenue NW, Washington, DC 20230-0002, and in the "Reading Room" section of the Board's website, which is accessible via [www.trade.gov/ftz](http://www.trade.gov/ftz).

For further information, contact Elizabeth Whiteman at [Elizabeth.Whiteman@trade.gov](mailto:Elizabeth.Whiteman@trade.gov) or (202) 482-0473.

Dated: December 4, 2018.

**Andrew McGilvray,**  
*Executive Secretary.*

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**BILLING CODE 3510-DS-P**

## DEPARTMENT OF COMMERCE

### Foreign-Trade Zones Board

[B-50-2018]

#### Foreign-Trade Zone (FTZ) 41—Milwaukee, Wisconsin; Authorization of Production Activity; Generac Power Systems, Inc. (Outdoor Power Equipment, Pumps, and Lawn and Garden Equipment); Jefferson and Whitewater, Wisconsin

On August 6, 2018, Generac Power Systems, Inc. (Generac) submitted a notification of proposed production activity to the FTZ Board for its facilities within Subzone 41], in Jefferson and Whitewater, Wisconsin.

The notification was processed in accordance with the regulations of the FTZ Board (15 CFR part 400), including notice in the **Federal Register** inviting public comment (83 FR 42108-42109, August 20, 2018). On December 4, 2018, the applicant was notified of the FTZ Board's decision that no further review of the activity is warranted at this time. The production activity described in the notification was authorized, subject to the FTZ Act and the FTZ Board's regulations, including Section 400.14, and further subject to a restriction requiring that foreign-status disposable textile bag liners and lithium-ion batteries be admitted to the subzone in privileged foreign status (19 CFR 146.41).

Dated: December 4, 2018.

**Andrew McGilvray,**  
*Executive Secretary.*

[FR Doc. 2018-26768 Filed 12-10-18; 8:45 am]

**BILLING CODE 3510-DS-P**

## DEPARTMENT OF COMMERCE

### Bureau of Industry and Security

[Docket No. 181108999-8999-01]

RIN 0694-XC051

#### Impact of the Implementation of the Chemical Weapons Convention (CWC) on Legitimate Commercial Chemical, Biotechnology, and Pharmaceutical Activities Involving "Schedule 1" Chemicals (Including Schedule 1 Chemicals Produced as Intermediates) During Calendar Year 2018

**AGENCY:** Bureau of Industry and Security, Commerce.

**ACTION:** Notice of inquiry.

**SUMMARY:** The Bureau of Industry and Security (BIS) is seeking public comments on the impact that implementation of the Chemical

Weapons Convention (CWC), through the Chemical Weapons Convention Implementation Act and the Chemical Weapons Convention Regulations (CWCR), has had on commercial activities involving “Schedule 1” chemicals during calendar year 2018. The purpose of this notice of inquiry is to collect information to assist BIS in its preparation of the annual certification to the Congress on whether the legitimate commercial activities and interests of chemical, biotechnology, and pharmaceutical firms are harmed by such implementation. This certification is required under Condition 9 of Senate Resolution 75 (April 24, 1997), in which the Senate gave its advice and consent to the ratification of the CWC.

**DATES:** Comments must be received by January 10, 2019.

**ADDRESSES:** You may submit comments by any of the following methods (please refer to RIN 0694–XC051 in all comments and in the subject line of email comments):

- *Federal rulemaking portal* (<http://www.regulations.gov>)—you can find this notice by searching on its *regulations.gov* docket number, which is BIS–2018–0032;

- *Email*: [willard.fisher@bis.doc.gov](mailto:willard.fisher@bis.doc.gov)—include the phrase “Schedule 1 Notice of Inquiry” in the subject line;

- *Fax*: (202) 482–3355 (Attn: Willard Fisher);

- By mail or delivery to Regulatory Policy Division, Bureau of Industry and Security, U.S. Department of Commerce, Room 2099B, 14th Street and Pennsylvania Avenue NW, Washington, DC 20230.

**FOR FURTHER INFORMATION CONTACT:** For questions on the Chemical Weapons Convention requirements for “Schedule 1” chemicals, contact Douglas Brown, Treaty Compliance Division, Office of Nonproliferation and Treaty Compliance, Bureau of Industry and Security, U.S. Department of Commerce, Phone: (202) 482–2163. For questions on the submission of comments, contact Willard Fisher, Regulatory Policy Division, Office of Exporter Services, Bureau of Industry and Security, U.S. Department of Commerce, Phone: (202) 482–2440.

#### **SUPPLEMENTARY INFORMATION:**

##### **Background**

In providing its advice and consent to the ratification of the Convention on the Prohibition of the Development, Production, Stockpiling, and Use of Chemical Weapons and Their Destruction, commonly called the Chemical Weapons Convention (CWC or “the Convention”), the Senate included,

in Senate Resolution 75 (S. Res. 75, April 24, 1997), several conditions to its ratification. Condition 9, titled “Protection of Advanced Biotechnology,” calls for the President to certify to Congress on an annual basis that “the legitimate commercial activities and interests of chemical, biotechnology, and pharmaceutical firms in the United States are not being significantly harmed by the limitations of the Convention on access to, and production of, those chemicals and toxins listed in Schedule 1.” On July 8, 2004, President Bush, by Executive Order 13346, delegated his authority to make the annual certification to the Secretary of Commerce.

The CWC is an international arms control treaty that contains certain verification provisions. In order to implement these verification provisions, the CWC established the Organization for the Prohibition of Chemical Weapons (OPCW). The CWC imposes certain obligations on countries that have ratified the Convention (*i.e.*, States Parties), among which are the enactment of legislation to prohibit the production, storage, and use of chemical weapons, and the establishment of a National Authority to serve as the national focal point for effective liaison with the OPCW and other States Parties in order to achieve the object and purpose of the Convention and the implementation of its provisions. The CWC also requires each State Party to implement a comprehensive data declaration and inspection regime to provide transparency and to verify that both the public and private sectors of the State Party are not engaged in activities prohibited under the CWC.

“Schedule 1” chemicals consist of those toxic chemicals and precursors set forth in the CWC “Annex on Chemicals” and in “Supplement No. 1 to part 712—SCHEDULE 1 CHEMICALS” of the Chemical Weapons Convention Regulations (CWCR) (15 CFR parts 710–722). The CWC identified these toxic chemicals and precursors as posing a high risk to the object and purpose of the Convention.

The CWC (Part VI of the “Verification Annex”) restricts the production of “Schedule 1” chemicals for protective purposes to two facilities per State Party: A single small-scale facility (SSSF) and a facility for production in quantities not exceeding 10 kg per year. The CWC Article-by-Article Analysis submitted to the Senate in Treaty Doc. 103–21 defined the term “protective purposes” to mean “used for determining the adequacy of defense equipment and measures.” Consistent with this definition and as authorized

by Presidential Decision Directive (PDD) 70 (December 17, 1999), which specifies agency and departmental responsibilities as part of the U.S. implementation of the CWC, the Department of Defense (DOD) was assigned the responsibility to operate these two facilities. Although this assignment of responsibility to DOD under PDD–70 effectively precluded commercial production of “Schedule 1” chemicals for protective purposes in the United States, it did not establish any limitations on “Schedule 1” chemical activities that are not prohibited by the CWC. However, DOD does maintain strict controls on “Schedule 1” chemicals produced at its facilities in order to ensure accountability for such chemicals, as well as their proper use, consistent with the object and purpose of the Convention.

The provisions of the CWC that affect commercial activities involving “Schedule 1” chemicals are implemented in the CWCR (see 15 CFR 712) and in the Export Administration Regulations (EAR) (see 15 CFR 742.18 and 15 CFR 745), both of which are administered by the Bureau of Industry and Security (BIS). Pursuant to CWC requirements, the CWCR restrict commercial production of “Schedule 1” chemicals to research, medical, or pharmaceutical purposes. The CWCR prohibit commercial production of “Schedule 1” chemicals for “protective purposes” because such production is effectively precluded per PDD–70, as described above—see 15 CFR 712.2(a). The CWCR also contain other requirements and prohibitions that apply to “Schedule 1” chemicals and/or “Schedule 1” facilities. Specifically, the CWCR:

- (1) Prohibit the import of “Schedule 1” chemicals from States not Party to the Convention (15 CFR 712.2(b));

- (2) Require annual declarations by certain facilities engaged in the production of “Schedule 1” chemicals in excess of 100 grams aggregate per calendar year (*i.e.*, declared “Schedule 1” facilities) for purposes not prohibited by the Convention (15 CFR 712.5(a)(1) and (a)(2));

- (3) Provide for government approval of “declared Schedule 1” facilities (15 CFR 712.5(f));

- (4) Provide that “declared Schedule 1” facilities are subject to initial and routine inspection by the OPCW (15 CFR 712.5(e) and 716.1(b)(1));

- (5) Require 200 days advance notification of establishment of new “Schedule 1” production facilities producing greater than 100 grams aggregate of “Schedule 1” chemicals per calendar year (15 CFR 712.4);

(6) Require advance notification and annual reporting of all imports and exports of “Schedule 1” chemicals to, or from, other States Parties to the Convention (15 CFR 712.6, 742.18(a)(1) and 745.1); and

(7) Prohibit the export of “Schedule 1” chemicals to States not Party to the Convention (15 CFR 742.18(a)(1) and (b)(1)(ii)).

For purposes of the CWCR (see 15 CFR 710.1), “production of a Schedule 1 chemical” means the formation of “Schedule 1” chemicals through chemical synthesis, as well as processing to extract and isolate “Schedule 1” chemicals produced by a biochemical or biologically mediated reaction. Such production is understood, for CWCR declaration purposes, to include intermediates, by-products, or waste products that are produced and consumed within a defined chemical manufacturing sequence, where such intermediates, by-products, or waste products are chemically stable and therefore exist for a sufficient time to make isolation from the manufacturing stream possible, but where, under normal or design operating conditions, isolation does not occur.

#### Request for Comments

In order to assist in determining whether the legitimate commercial activities and interests of chemical, biotechnology, and pharmaceutical firms in the United States are significantly harmed by the limitations of the Convention on access to, and production of, “Schedule 1” chemicals as described in this notice, BIS is seeking public comments on any effects that implementation of the CWC, through the Chemical Weapons Convention Implementation Act and the CWCR, has had on commercial activities involving “Schedule 1” chemicals during calendar year 2018. To allow BIS to properly evaluate the significance of any harm to commercial activities involving “Schedule 1” chemicals, public comments submitted in response to this notice of inquiry should include both a quantitative and qualitative assessment of the impact of the CWC on such activities.

#### Submission of Comments

All comments must be submitted to one of the addresses indicated in this notice. The Department requires that all comments be submitted in written form. BIS will consider all comments received on or before January 10, 2019. All comments (including any personally identifying information or information for which a claim of confidentiality is

asserted either in those comments or their transmittal emails) will be made available for public inspection and copying. Parties who wish to comment anonymously may do so by submitting their comments via *Regulations.gov*, leaving the fields that would identify the commenter blank and including no identifying information in the comment itself.

Dated: December 3, 2018.

**Matthew S. Borman,**

*Deputy Assistant Secretary for Export Administration.*

[FR Doc. 2018–26734 Filed 12–10–18; 8:45 am]

**BILLING CODE 3510–33–P**

## DEPARTMENT OF COMMERCE

### International Trade Administration

#### Initiation of Antidumping and Countervailing Duty Administrative Reviews

**AGENCY:** Enforcement and Compliance, International Trade Administration, Department of Commerce.

**SUMMARY:** The Department of Commerce (Commerce) has received requests to conduct administrative reviews of various antidumping and countervailing duty orders and findings with October anniversary dates. In accordance with Commerce’s regulations, we are initiating those administrative reviews.

**DATES:** Applicable December 11, 2018.

**FOR FURTHER INFORMATION CONTACT:** Brenda E. Brown, Office of AD/CVD Operations, Customs Liaison Unit, Enforcement and Compliance, International Trade Administration, U.S. Department of Commerce, 1401 Constitution Avenue NW, Washington, DC 20230, telephone: (202) 482–4735.

#### SUPPLEMENTARY INFORMATION:

##### Background

Commerce has received timely requests, in accordance with 19 CFR 351.213(b), for administrative reviews of various antidumping and countervailing duty orders and findings with October anniversary dates.

All deadlines for the submission of various types of information, certifications, or comments or actions by Commerce discussed below refer to the number of calendar days from the applicable starting time.

##### Notice of No Sales

If a producer or exporter named in this notice of initiation had no exports, sales, or entries during the period of review (POR), it must notify Commerce within 30 days of publication of this

notice in the **Federal Register**. All submissions must be filed electronically at <http://access.trade.gov> in accordance with 19 CFR 351.303.<sup>1</sup> Such submissions are subject to verification in accordance with section 782(i) of the Tariff Act of 1930, as amended (the Act). Further, in accordance with 19 CFR 351.303(f)(1)(i), a copy must be served on every party on Commerce’s service list.

#### Respondent Selection

In the event Commerce limits the number of respondents for individual examination for administrative reviews initiated pursuant to requests made for the orders identified below, Commerce intends to select respondents based on U.S. Customs and Border Protection (CBP) data for U.S. imports during the period of review. We intend to place the CBP data on the record within five days of publication of the initiation notice and to make our decision regarding respondent selection within 30 days of publication of the initiation **Federal Register** notice. Comments regarding the CBP data and respondent selection should be submitted seven days after the placement of the CBP data on the record of this review. Parties wishing to submit rebuttal comments should submit those comments five days after the deadline for the initial comments.

In the event Commerce decides it is necessary to limit individual examination of respondents and conduct respondent selection under section 777A(c)(2) of the Act:

In general, Commerce has found that determinations concerning whether particular companies should be “collapsed” (e.g., treated as a single entity for purposes of calculating antidumping duty rates) require a substantial amount of detailed information and analysis, which often require follow-up questions and analysis. Accordingly, Commerce will not conduct collapsing analyses at the respondent selection phase of this review and will not collapse companies at the respondent selection phase unless there has been a determination to collapse certain companies in a previous segment of this antidumping proceeding (e.g., investigation, administrative review, new shipper review or changed circumstances review). For any company subject to this review, if Commerce determined, or continued to treat, that company as collapsed with others, Commerce will

<sup>1</sup> See *Antidumping and Countervailing Duty Proceedings: Electronic Filing Procedures; Administrative Protective Order Procedures*, 76 FR 39263 (July 6, 2011).