

proposed licensing action. Provided that the field test data support the conclusions of the original EA and the issuance of a FONSI, APHIS does not intend to issue a separate EA and FONSI to support the issuance of the associated product license, and would determine that an environmental impact statement need not be prepared. APHIS intends to issue a veterinary biological product license for this vaccine following satisfactory completion of the field test provided no adverse impacts on the human environment are identified and provided the product meets all other requirements for licensing.

Authority: 21 U.S.C. 151–159.

Done in Washington, DC, this 3rd day of October 2018.

Kevin Shea,

Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2018–21926 Filed 10–9–18; 8:45 am]

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DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2018–0033]

Oral Rabies Vaccine Trial; Availability of a Supplement to an Environmental Assessment and Finding of No Significant Impact

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Notice.

SUMMARY: We are advising the public that the Animal and Plant Health Inspection Service has prepared a supplement to an environmental assessment and finding of no significant impact relative to an oral rabies vaccination field trial in New Hampshire, New York, Ohio, Vermont, and West Virginia. Based on its finding of no significant impact, the Animal and Plant Health Inspection Service has determined that an environmental impact statement need not be prepared.

FOR FURTHER INFORMATION CONTACT: Mr. Richard Chipman, Rabies Program Coordinator, Wildlife Services, APHIS, 59 Chennell Drive, Suite 7, Concord, NH 03301; (603) 223–9623. To obtain copies of the supplement to the environmental assessment and the finding of no significant impact, contact Ms. Beth Kabert, Environmental Coordinator,

Wildlife Services, 140–C Locust Grove Road, Pittstown, NJ 08867; (908) 735–5654, fax (908) 735–0821, email: beth.e.kabert@aphis.usda.gov.

SUPPLEMENTARY INFORMATION: The Wildlife Services (WS) program in the Animal and Plant Health Inspection Service (APHIS) cooperates with Federal agencies, State and local governments, and private individuals to research and implement the best methods of managing conflicts between wildlife and human health and safety, agriculture, property, and natural resources. Wildlife-borne diseases that can affect domestic animals and humans are among the types of conflicts that APHIS–WS addresses. Wildlife is the dominant reservoir of rabies in the United States.

On July 3, 2018, we published in the **Federal Register** (83 FR 31117–31118, Docket No. APHIS–2018–0033) a notice¹ in which we announced the availability, for public review and comment, of a supplement to an environmental assessment (EA) that examined the potential environmental impacts associated with the proposed field trial to test the safety and efficacy of an experimental oral rabies vaccine (ORV) for wildlife in New Hampshire, New York, Ohio, Vermont, and West Virginia. In addition, the supplement analyzed the potential impacts of expanding the geographic range of the field trial zone to two additional counties in Ohio and four additional counties in West Virginia.

We solicited comments on the EA for 30 days ending August 2, 2018. We did not receive any comments.

In this document, we are advising the public of our finding of no significant impact (FONSI) relative to the ORV field trial in New Hampshire, New York, Ohio, Vermont, and West Virginia. The finding, which is based on the EA and the 2013, 2015, and 2017 supplements to the EA, reflects our determination that the distribution of this experimental wildlife rabies vaccine will not have a significant impact on the quality of the human environment.

The 2018 supplement to the EA and the FONSI may be viewed on the APHIS website at <http://www.aphis.usda.gov/wildlifedamage/nepa> and on the *Regulations.gov* website (see footnote 1). Copies of the 2018 supplement to the EA and the FONSI are also available for public inspection at USDA, Room 1141, South Building, 14th Street and

Independence Avenue SW, Washington, DC, between 8 a.m. and 4:30 p.m., Monday through Friday, except holidays. Persons wishing to inspect copies are requested to call ahead on (202) 799–7039 to facilitate entry into the reading room. In addition, copies may be obtained as described under **FOR FURTHER INFORMATION CONTACT**.

The 2018 supplement to the EA and the FONSI have been prepared in accordance with: (1) The National Environmental Policy Act of 1969 (NEPA), as amended (42 U.S.C. 4321 *et seq.*); (2) regulations of the Council on Environmental Quality for implementing the procedural provisions of NEPA (40 CFR parts 1500–1508); (3) USDA regulations implementing NEPA (7 CFR part 1b); and (4) APHIS' NEPA Implementing Procedures (7 CFR part 372).

Done in Washington, DC, this 3rd day of October 2018.

Kevin Shea,

Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2018–21924 Filed 10–9–18; 8:45 am]

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DEPARTMENT OF COMMERCE

Economic Development Administration

Notice of Petitions by Firms for Determination of Eligibility To Apply for Trade Adjustment Assistance

AGENCY: Economic Development Administration, U.S. Department of Commerce.

ACTION: Notice and opportunity for public comment.

SUMMARY: The Economic Development Administration (EDA) has received petitions for certification of eligibility to apply for Trade Adjustment Assistance from the firms listed below.

Accordingly, EDA has initiated investigations to determine whether increased imports into the United States of articles like or directly competitive with those produced by each of the firms contributed importantly to the total or partial separation of the firms' workers, or threat thereof, and to a decrease in sales or production of each petitioning firm.

SUPPLEMENTARY INFORMATION:

¹ To view the notice, the EA, and the FONSI, go to <http://www.regulations.gov/#!docketDetail;D=APHIS-2018-0033>.

LIST OF PETITIONS RECEIVED BY EDA FOR CERTIFICATION OF ELIGIBILITY TO APPLY FOR TRADE ADJUSTMENT
ASSISTANCE
[9/18/2018 through 10/1/2018]

Firm name	Firm address	Date accepted for investigation	Product(s)
Belta-4 Manufacturing, LLC	4300 Garland Drive, Haltom City, TX 76117.	9/19/2018	The firm manufactures metal parts and assemblies, including seat racks, manifolds, short rods with threads, base plates, and sleeves.
Tempel Steel Company, Inc	5500 North Wolcott Avenue, Chicago, IL 60640.	9/21/2018	The firm manufactures flat-rolled silicon electrical steel for the automotive, motor, generator, transformer, and lighting industries.
US Felt Company, Inc	61 Industrial Avenue, Sanford, ME 04073.	9/21/2018	The firm manufactures non-woven fabrics, felt, and composite materials.
Garage Graphics & Visuals, Inc. d/b/a Elemoose	2503 North Neergard Avenue, Springfield, MO 65803.	9/24/2018	The firm manufactures custom signs, exhibits, sculptures, stage sets, and architectural elements.
Environmental Advisors and Engineers, Inc.	19211 West 64th Terrace, Shawnee, KS 66218.	9/25/2018	The firm provides consulting services, including engineering and construction monitoring services.

Any party having a substantial interest in these proceedings may request a public hearing on the matter. A written request for a hearing must be submitted to the Trade Adjustment Assistance Division, Room 71030, Economic Development Administration, U.S. Department of Commerce, Washington, DC 20230, no later than ten (10) calendar days following publication of this notice. These petitions are received pursuant to section 251 of the Trade Act of 1974, as amended.

Please follow the requirements set forth in EDA's regulations at 13 CFR 315.9 for procedures to request a public hearing. The Catalog of Federal Domestic Assistance official number and title for the program under which these petitions are submitted is 11.313, Trade Adjustment Assistance for Firms.

Irette Patterson,
Program Analyst.

[FR Doc. 2018-21929 Filed 10-9-18; 8:45 am]

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DEPARTMENT OF COMMERCE

Foreign-Trade Zones Board

[B-34-2018]

Foreign-Trade Zone (FTZ) 230—Greensboro, North Carolina, Authorization of Production Activity, Patheon Softgels (Pharmaceutical Products), High Point, North Carolina

On May 31, 2018, The Piedmont Triad Partnership, grantee of FTZ 230, submitted a notification of proposed production activity to the FTZ Board on behalf of Patheon Softgels, within Subzone 230C, in High Point, North Carolina.

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 26947, June 11, 2018). On September 28, 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.

Dated: October 1, 2018.

Andrew McGilvray,
Executive Secretary.

[FR Doc. 2018-21981 Filed 10-9-18; 8:45 am]

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DEPARTMENT OF COMMERCE

International Trade Administration

[A-533-843]

Certain Lined Paper Products From India: Preliminary Results of Antidumping Duty Administrative Review and Preliminary Determination of No Shipments; 2016-2017

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

SUMMARY: The Department of Commerce (Commerce) preliminarily determines that sales of certain lined paper products were not made at less than normal value during the September 1, 2016, through August 31, 2017, period of review (POR). We invite interested parties to comment on these preliminary results.

DATES: Applicable October 10, 2018.

FOR FURTHER INFORMATION CONTACT:

Cindy Robinson or Joy Zhang, AD/CVD Operations, Office III, Enforcement and Compliance, International Trade Administration, U.S. Department of Commerce, 1401 Constitution Avenue NW, Washington DC 20230; telephone (202) 482-3797 or (202) 482-1168, respectively.

Background

On September 28, 2006, Commerce published the *CLPP from India Order* in the **Federal Register**.¹ On November 13, 2017, pursuant to section 751(a)(1) of the Tariff Act of 1930, as amended (the Act), Commerce initiated an administrative review of the antidumping duty order on certain lined paper products from India.² On January 23, 2018, Commerce exercised its discretion to toll all deadlines affected by the closure of the Federal Government from January 20 through 22, 2018.³ On May 10, 2018, we extended the deadline for the preliminary results to October 3, 2018.⁴

¹ See *Notice of Amended Final Determination of Sales at Less Than Fair Value: Certain Lined Paper Products from the People's Republic of China; Notice of Antidumping Duty Orders: Certain Lined Paper Products from India, Indonesia and the People's Republic of China; and Notice of Countervailing Duty Orders: Certain Lined Paper Products from India and Indonesia*, 71 FR 56949 (September 28, 2006) (*CLPP from India Order*).

² See *Initiation of Antidumping and Countervailing Duty Administrative Reviews*, 82 FR 52268 (November 13, 2017) (*Initiation Notice*).

³ See memorandum, "Deadlines Affected by the Shutdown of the Federal Government," dated January 23, 2018. All deadlines in this segment of the proceeding have been extended by three days.

⁴ See Memorandum, "Certain Lined Paper Products from India: Extension of Time Limit for Preliminary Results of Antidumping Duty Administrative Review; 2016-2017," dated May 10, 2018.