

www.regulations.gov/
 #!docketDetail;D=APHIS-2015-0003 or
 in our reading room, which is located in
 room 1141 of the USDA South Building,
 14th Street and Independence Avenue
 SW., Washington, DC. Normal reading
 room hours are 8 a.m. to 4:30 p.m.,
 Monday through Friday, except
 holidays. To be sure someone is there to
 help you, please call (202) 799-7039
 before coming.

FOR FURTHER INFORMATION CONTACT: Dr.
 Donna Malloy, Operational Support
 Section, Center for Veterinary Biologics,
 Policy, Evaluation, and Licensing, VS,
 APHIS, 4700 River Road Unit 148,
 Riverdale, MD 20737-1231; phone (301)
 851-3426, fax (301) 734-4314.

For information regarding the
 environmental assessment or the risk
 analysis, or to request a copy of the
 environmental assessment (as well as
 the risk analysis with confidential
 business information removed), contact
 Dr. Patricia L. Foley, Risk Manager,
 Center for Veterinary Biologics, Policy,
 Evaluation, and Licensing, VS, APHIS,
 1920 Dayton Avenue, P.O. Box 844,
 Ames, IA 50010; phone (515) 337-6100,
 fax (515) 337-6120.

SUPPLEMENTARY INFORMATION:

Under the Virus-Serum-Toxin Act (21
 U.S.C. 151 *et seq.*), a veterinary
 biological product must be shown to be
 pure, safe, potent, and efficacious before
 a veterinary biological product license
 may be issued. A field test is generally
 necessary to satisfy prelicensing
 requirements for veterinary biological
 products. Prior to conducting a field test
 on an unlicensed product, an applicant
 must obtain approval from the Animal
 and Plant Health Inspection Service
 (APHIS), as well as obtain APHIS'
 authorization to ship the product for
 field testing.

To determine whether to authorize
 shipment and grant approval for the
 field testing of the unlicensed product
 referenced in this notice, APHIS
 considers the potential effects of this
 product on the safety of animals, public
 health, and the environment. Using the
 risk analysis and other relevant data,
 APHIS has prepared an environmental
 assessment (EA) concerning the field
 testing of the following unlicensed
 veterinary biological product:

Requester: Merial, Inc.

Product: Marek's Disease Vaccine,
 Serotype 1, Live Virus.

Possible Field Test Locations:
 Arkansas, Georgia, Kentucky, North
 Carolina, Tennessee, and Texas.

The above-mentioned product is a
 live Marek's Disease serotype 1 vaccine
 virus containing the long terminal
 repeat of the reticuloendotheliosis virus.

The attenuated vaccine is intended for
 use in healthy day-old chickens, as an
 aid in the prevention of Marek's disease
 caused by very virulent Marek's disease
 virus.

The EA has 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).

Unless substantial issues with adverse
 environmental impacts are raised in
 response to this notice, APHIS intends
 to issue a finding of no significant
 impact (FONSI) based on the EA and
 authorize shipment of the above product
 for the initiation of field tests following
 the close of the comment period for this
 notice.

Because the issues raised by field
 testing and by issuance of a license are
 identical, APHIS has concluded that the
 EA that is generated for field testing
 would also be applicable to the
 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 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 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 8th day of
 April 2015.

Kevin Shea,

*Administrator, Animal and Plant Health
 Inspection Service.*

[FR Doc. 2015-08604 Filed 4-13-15; 8:45 am]

BILLING CODE 3410-34-P

DEPARTMENT OF COMMERCE

**Submission for OMB Review;
 Comment Request**

The Department of Commerce will
 submit to the Office of Management and
 Budget (OMB) for clearance the
 following proposal for collection of
 information under the provisions of the

Paperwork Reduction Act (44 U.S.C.
 chapter 35).

Agency: U.S. Census Bureau.
Title: Comparing Health Insurance
 Measurement Error (CHIME).

OMB Control Number: 0607-XXXX.

Form Number(s): No forms;

respondent information collected by
 telephone interview.

Type of Request: Regular submission.

Number of Respondents: 5,000

Households.

Average Hours per Response: 13
 minutes.

Burden Hours: 3,028 hours.

Needs and Uses: The goal of the study
 is to assess measurement error in health
 coverage estimates that is ascribable to
 the questionnaire across the CPS and
 ACS health insurance modules using
 administrative records as a truth source.
 Both "absolute" reporting accuracy (the
 survey report compared to the
 administrative record data) and
 "relative" reporting accuracy
 (comparing absolute accuracy across
 questionnaire treatments) will be
 evaluated. The analysis will be used to
 understand the magnitude, direction
 and patterns of misreporting for three
 main purposes: (1) To provide Census
 program staff with empirical data to
 develop and refine edits and/or to
 include research notes for data users so
 they can make their own adjustments
 for misreporting; (2) to equip the wider
 research community with information
 that could serve as a guide for deciding
 which among the various surveys best
 suits their needs; and (3) to contribute
 to the general survey methods research
 literature on measurement error.

Analysis will also inform reporting
 accuracy of health coverage related to
 the Affordable Care Act (ACA).
 Specifically, for coverage that is known
 to be obtained from the marketplace, we
 will explore whether respondents report
 that coverage, the source they cite
 (direct-purchase, government, etc.), and
 the accuracy with which they answer a
 question on subsidized premiums.

Affected Public: Individuals or
 households.

Frequency: One time.

Respondent's Obligation: Voluntary.

Legal Authority: Title 13, United
 States Code, sections 141, 182 and 193.

*This information collection request
 may be viewed at www.reginfo.gov.*

Follow the instructions to view
 Department of Commerce collections
 currently under review by OMB.

Written comments and
 recommendations for the proposed
 information collection should be sent
 within 30 days of publication of this
 notice to [OIRA_Submission@](mailto:OIRA_Submission@omb.eop.gov)
omb.eop.gov or fax to (202)395-5806.

Dated: April 8, 2015.

Glenna Mickelson,

Management Analyst, Office of the Chief Information Officer.

[FR Doc. 2015-08473 Filed 4-13-15; 8:45 am]

BILLING CODE 3510-07-P

DEPARTMENT OF COMMERCE

Foreign-Trade Zones Board

[B-86-2014]

Production Activity Not Authorized; Foreign-Trade Zone 57—Charlotte, North Carolina; Gildan Yarns, LLC; (Cotton, Cotton/Polyester Yarns); Salisbury, North Carolina

On December 8, 2014, the Charlotte Regional Partnership, grantee of FTZ 57, submitted a notification of proposed production activity to the Foreign-Trade Zones (FTZ) Board on behalf of Gildan Yarns, LLC, in Salisbury, 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 (79 FR 75532, 12-18-2014). Pursuant to Section 400.37, the FTZ Board has determined that further review is warranted and has not authorized the proposed activity. If the applicant wishes to seek authorization for this activity, it will need to submit an application for production authority, pursuant to Section 400.23.

Dated: April 8, 2015.

Andrew McGilvray,

Executive Secretary.

[FR Doc. 2015-08592 Filed 4-13-15; 8:45 am]

BILLING CODE 3510-DS-P

DEPARTMENT OF COMMERCE

Foreign-Trade Zones Board

[B-20-2015]

Foreign-Trade Zone (FTZ) 50—Long Beach, California; Notification of Proposed Production Activity; Mercedes Benz USA, LLC; (Accessorizing Motor Vehicles); Long Beach, California

The Port of Long Beach, grantee of FTZ 50, submitted a notification of proposed production activity to the FTZ Board on behalf of Mercedes Benz USA, LLC (MBUSA), located in Long Beach, California. The notification conforming to the requirements of the regulations of the FTZ Board (15 CFR 400.22) was received on March 24, 2015.

The MBUSA facility is located within Site 41 of FTZ 50. The facility is used

for accessorizing passenger motor vehicles. Pursuant to 15 CFR 400.14(b), FTZ activity would be limited to the specific foreign-status materials and components and specific finished products described in the submitted notification (as described below) and subsequently authorized by the FTZ Board.

Production under FTZ procedures could exempt MBUSA from customs duty payments on the foreign status components used in export production. On its domestic sales, MBUSA would be able to choose the duty rate during customs entry procedures that applies to passenger motor vehicles (duty rate – 2.5%) for the foreign status components noted below. Customs duties also could possibly be deferred or reduced on foreign status production equipment.

The components sourced from abroad include: Plastic door sills and strips; wheel rim locks; metal door sills and strips; memory cards; navigation systems and related parts; entertainment systems; wiring sets and harnesses; storage compartments; spoilers; and, cup holders (duty rate ranges from free to 5%).

Public comment is invited from interested parties. Submissions shall be addressed to the FTZ Board's Executive Secretary at the address below. The closing period for their receipt is May 26, 2015.

A copy of the notification 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 FTZ Board's Web site, which is accessible via www.trade.gov/ftz.

For further information, contact Pierre Duy at Pierre.Duy@trade.gov or (202) 482-1378.

Dated: April 7, 2015.

Andrew McGilvray,

Executive Secretary.

[FR Doc. 2015-08590 Filed 4-13-15; 8:45 am]

BILLING CODE 3510-DS-P

DEPARTMENT OF COMMERCE

Foreign-Trade Zones Board

[Order No. 1974]

Reorganization of Foreign-Trade Zone 286; (Expansion of Service Area); Under Alternative Site Framework; Caledonia, Essex and Orleans Counties, Vermont

Pursuant to its authority under the Foreign-Trade Zones Act of June 18, 1934, as amended (19 U.S.C. 81a-81u), the Foreign-Trade Zones Board (the Board) adopts the following Order:

Whereas, the Board adopted the alternative site framework (ASF) (15 CFR Sec. 400.2(c)) as an option for the establishment or reorganization of zones;

Whereas, the Northeastern Vermont Development Association, grantee of FTZ 286, submitted an application to the Board (FTZ Docket B-60-2014, docketed 08-27-2014) for authority to expand the service area of the zone to include Lamoille County, as described in the application, adjacent to the Derby Line Customs and Border Protection port of entry;

Whereas, notice inviting public comment was given in the **Federal Register** (79 FR 52300, 09-03-14) and the application has been processed pursuant to the FTZ Act and the Board's regulations; and,

Whereas, the Board adopts the findings and recommendations of the examiner's report, and finds that the requirements of the FTZ Act and the Board's regulations are satisfied;

Now, therefore, the Board hereby orders:

The application to reorganize FTZ 286 to expand the service area under the ASF is approved, subject to the FTZ Act and the Board's regulations, including Section 400.13, and to the Board's standard 2,000-acre activation limit for the zone.

Signed at Washington, DC this 3rd day of April 2015.

Paul Piquado,

Assistant Secretary of Commerce for Enforcement and Compliance, Alternate Chairman, Foreign-Trade Zones Board.

ATTEST: _____

Andrew McGilvray,

Executive Secretary.

[FR Doc. 2015-08584 Filed 4-13-15; 8:45 am]

BILLING CODE 3510-DS-P