

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. 2015–08602 Filed 4–13–15; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2015–0025]

Secretary's Advisory Committee on Animal Health; Meeting

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Notice of meeting.

SUMMARY: This is a notice to inform the public of an upcoming meeting of the Secretary's Advisory Committee on Animal Health. The meeting is being organized by the Animal and Plant Health Inspection Service to discuss matters of animal health.

DATES: The meeting will be held on April 28 and 29, 2015, from 9 a.m. to 5 p.m. each day.

ADDRESSES: The meeting will be held at the Bethesda Marriott, 5151 Pooks Hill Road, Bethesda, MD 20814.

FOR FURTHER INFORMATION CONTACT: Mrs. R.J. Cabrera, Designated Federal Officer, VS, APHIS, 4700 River Road Unit 34, Riverdale, MD 20737; phone (301) 851–3478, email SACAH.Management@aphis.usda.gov.

SUPPLEMENTARY INFORMATION:

The Secretary's Advisory Committee on Animal Health (the Committee) advises the Secretary of Agriculture on matters of animal health, including means to prevent, conduct surveillance on, monitor, control, or eradicate animal diseases of national importance. In doing so, the Committee will consider public health, conservation of natural resources, and the stability of livestock economies.

Tentative topics for discussion at the meeting include:

- Follow-on discussion of antimicrobial resistance, mitigations, and the U.S. Department of Agriculture (USDA) action plan,
- Comprehensive discussion on porcine epidemic diarrhea,
- Follow-on discussion on foot-and-mouth disease,
- USDA draft framework for emerging diseases,
- Proposed national list of reportable animal diseases,
- Avian influenza, and
- Bovine tuberculosis program—understanding the disease.

A final agenda will be posted on the Committee Web site by April 13, 2015.

Those wishing to attend the meeting in person must complete a brief registration form by clicking on the “SACAH Meeting Sign-Up” button on the Committee's Web site (<http://www.aphis.usda.gov/animalhealth/sacah>). Members of the public may also join the meeting via teleconference in “listen-only” mode. Participants who wish to listen in on the teleconference may do so by dialing 1–888–469–3079 and then entering the public passcode, 2061888#.

Due to time constraints, members of the public will not have an opportunity to participate in the Committee's discussions. However, questions and written statements for the Committee's consideration may be submitted up to 5 working days before the meeting. They may be sent to SACAH.Management@aphis.usda.gov or mailed to the person listed on the notice under **FOR FURTHER INFORMATION CONTACT**. Statements filed with the Committee should specify that they pertain to the April 2015 Committee meeting.

This notice of meeting is given pursuant to section 10 of the Federal Advisory Committee Act (5 U.S.C. App. 2).

Done in Washington, DC, this 8th day of April 2015.

Kevin Shea,

Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2015–08603 Filed 4–13–15; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2015–0003]

Availability of an Environmental Assessment for Field Testing a Marek's Disease Vaccine, Serotype 1, Live Virus

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Notice of availability.

SUMMARY: We are advising the public that the Animal and Plant Health Inspection Service has prepared an environmental assessment concerning authorization to ship for the purpose of field testing, and then to field test, an unlicensed Marek's disease vaccine, serotype 1, live virus. The environmental assessment, which is based on a risk analysis prepared to assess the risks associated with the field testing of this vaccine, examines the potential effects that field testing this veterinary vaccine could have on the quality of the human environment. Based on the risk analysis and other relevant data, we have reached a preliminary determination that field testing this veterinary vaccine will not have a significant impact on the quality of the human environment, and that an environmental impact statement need not be prepared. We intend to authorize shipment of this vaccine for field testing following the close of the comment period for this notice unless new substantial issues bearing on the effects of this action are brought to our attention. We also intend to issue a U.S. Veterinary Biological Product license for this vaccine, provided the field test data support the conclusions of the environmental assessment and the issuance of a finding of no significant impact and the product meets all other requirements for licensing.

DATES: We will consider all comments that we receive on or before May 14, 2015.

ADDRESSES: You may submit comments by either of the following methods:

- Federal eRulemaking Portal: Go to <http://www.regulations.gov/#/docketDetail;D=APHIS-2015-0003>.
- Postal Mail/Commercial Delivery:

Send your comment to Docket No. APHIS–2015–0003, Regulatory Analysis and Development, PPD, APHIS, Station 3A–03.8, 4700 River Road Unit 118, Riverdale, MD 20737–1238.

Supporting documents and any comments we receive on this docket may be viewed at <http://>

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.