FURTHER INFORMATION CONTACT. Please refer to the docket number, date, and complete title of this notice when requesting copies. A copy of the draft environmental assessment (as well as the risk analysis with confidential business information removed) and any comments that we receive on this docket are available for public inspection in our reading room. The reading room is located in room 1141 of the 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) 690–2817 before coming.

APHIS documents published in the **Federal Register**, and related information, including the names of organizations and individuals who have commented on APHIS dockets, are available on the Internet at http://www.aphis.usda.gov/ppd/rad/webrepor.html.

FOR FURTHER INFORMATION CONTACT: Dr. Albert P. Morgan, Chief Staff Officer, Operational Support Section, Center for Veterinary Biologics, Licensing and Policy Development, VS, APHIS, USDA, 4700 River Road, Unit 148, Riverdale, MD 20737–1231; telephone (301) 734–8245; fax (301) 734–4314.

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 conducted a risk analysis to assess the potential effects of this product on the safety of animals, public health, and the environment.

Based on the risk analysis, APHIS has prepared a draft environmental assessment (EA) concerning the field testing of the combined unlicensed and licensed veterinary biological product:

Requester: Biomune Company.

Product: Avian Encephalomyelitis-Fowl Pox-Laryngotracheitis Vaccine, Live Virus, Fowl Pox Vector.

Field test locations: Georgia, Kentucky, Nebraska, Texas, and Virginia.

The above-mentioned product is a modified live avian encephalomyelitis vaccine in combination with a live, attenuated fowl pox virus that has been genetically modified to express fowl laryngotracheitis antigens. The vaccine is for use in chickens as an aid in the prevention of avian encephalomyelitis, fowl pox, and laryngotracheitis.

The draft 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 provision 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 environmental issues are raised in response to this notice, APHIS intends to issue a final EA and finding of no significant impact (FONSI) 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 licensure.

Authority: 21 U.S.C. 151–159.

Done in Washington, DC, this 21st day of June 2001.

Bobby R. Acord,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 01–16136 Filed 6–26–01; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. 97–093–8]

Scrapie Eradication Uniform Methods and Rules; Reopening and Extension of Comment Period

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Notice.

SUMMARY: We are reopening and extending the comment period for a notice seeking public comments on the draft Scrapie Eradication Uniform Methods and Rules. This action will allow interested persons additional time to prepare and submit comments.

DATES: We invite you to comment on the draft Scrapie Eradication Uniform Methods and Rules. We will consider all comments on Docket 97–093–7 that we receive by August 20, 2001.

ADDRESSES: Please send four copies of your comment (an original and three copies) to: Docket No. 97–093–7, Regulatory Analysis and Development, PPD, APHIS, Suite 3C03, 4700 River Road Unit 118, Riverdale, MD 20737–1238.

Please state that your comment refers to Docket No. 97–093–7.

You may read any comments that we receive on Docket No. 97–093–7 in our reading room. The reading room 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) 690–2817 before coming.

APHIS documents published in the **Federal Register**, and related information, including the names of organizations and individuals who have commented on APHIS dockets, are available on the Internet at http://www.aphis.usda.gov/ppd/rad/webrepor.html.

You may request a copy of the draft Scrapie Eradication Uniform Methods and Rules by writing to the person listed under **FOR FURTHER INFORMATION CONTACT**. The document is also available on the Internet at http://www.aphis.usda.gov/vs/scrapie, and we may post revised versions to this website for additional comment in the future.

FOR FURTHER INFORMATION CONTACT: Dr. Diane Sutton, National Scrapie Program Coordinator, National Animal Health

Programs Staff, VS, APHIS, 4700 River Road Unit 43, Riverdale, MD 20737–1231; (301) 734–6954.

SUPPLEMENTARY INFORMATION: Scrapie is a degenerative and eventually fatal disease affecting the central nervous systems of sheep and goats. To control the spread of scrapie within the United States, the Animal and Plant Health Inspection Service (APHIS), U.S. Department of Agriculture, administers regulations at 9 CFR part 79 that restrict the interstate movement of certain sheep and goats. APHIS also has regulations at 9 CFR part 54 that describe a voluntary scrapie control program.

On April 20, 2001, we published a notice in the **Federal Register** (66 FR 20231, Docket No. 97–093–7) soliciting comments on the draft Scrapie Eradication Uniform Methods and Rules(UM&R). The UM&R is a set of proposed cooperative procedures and standards to aid in the control and eradication of scrapie. The legal requirements for interstate movement of sheep and goats due to scrapie are contained in title 9 of the Code of Federal Regulations. The ScrapieEradication UM&R provides guidance to the States regarding the minimum standards necessary for a State to participate in the national eradication program.

Comments on the UM&R were required to be received on or before June 19, 2001. We are reopening and extending the comment period on Docket No. 97–093–7. This action will allow interested persons additional time to prepare and submit comments.

Authority: 21 U.S.C. 111–113, 114, 114a, 115, 117, 120, 121, 123–126, and 134a–134h; 7 CFR 2.22, 2.80, and 371.4.

Done in Washington, DC, this 21st day of June 2001.

Bobby R. Acord,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 01–16135 Filed 6–26–01; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Food and Nutrition Service

Child and Adult Care Food Program: National Average Payment Rates, Day Care Home Food Service Payment Rates, and Administrative Reimbursement Rates for Sponsoring Organizations of Day Care Homes for the Period July 1, 2001–June 30, 2002.

AGENCY: Food and Nutrition Service, USDA.

ACTION: Notice.

SUMMARY: This notice announces the annual adjustments to: the national average payment rates for meals and supplements served in child care centers, outside-school-hours care centers, at-risk afterschool care centers, and adult day care centers; the food service payment rates for meals and supplements served in day care homes; and the administrative reimbursement rates for sponsoring organizations of day care homes, to reflect changes in the Consumer Price Index. Further adjustments are made to these rates to reflect the higher costs of providing meals in the States of Alaska and Hawaii. The adjustments contained in this notice are made on an annual basis each July, as required by the statutes and regulations governing the Child and Adult Care Food Program (CACFP).

EFFECTIVE DATE: July 1, 2001.

FOR FURTHER INFORMATION CONTACT: Melissa Rothstein, Section Chief, Child and Adult Care and Summer Programs Section, Policy and Program Development Branch, Child Nutrition Division, Food and Nutrition Service, USDA, Alexandria, Virginia, 22302, (703) 305–2620.

SUPPLEMENTARY INFORMATION:

Definitions

The terms used in this notice shall have the meanings ascribed to them in the regulations governing the CACFP (7 CFR part 226).

Background

Pursuant to sections 4, 11 and 17 of the Richard B. Russell National School Lunch Act (NSLA) (42 U.S.C. 1753, 1759a and 1766), section 4 of the Child Nutrition Act of 1966 (CNA) (42 U.S.C. 1773) and sections 226.4, 226.12 and 226.13 of the regulations governing the CACFP (7 CFR Part 226), notice is hereby given of the new payment rates for institutions participating in CACFP. These rates shall be in effect during the period July 1, 2001 through June 30, 2002.

As provided for under the NSLA and the CNA, all rates in the CACFP must be revised annually on July 1 to reflect changes in the Consumer Price Index (CPI) for the most recent 12-month period. In accordance with this mandate, the Department last published the adjusted national average payment rates for centers, the food service payment rates for day care homes, and the administrative reimbursement rates for sponsors of day care homes on June 27, 2000, at 65 FR 39589 (for the period July 1, 2000–June 30, 2001).

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