States of America, as represented by the Secretary of Agriculture. The prospective exclusive license will be royalty-bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. The prospective exclusive license may be granted unless, within ninety (90) days from the date of this published Notice, Agricultural Research Service receives written evidence and argument which establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR 404.7.

Michael D. Ruff,

Assistant Administrator.

[FR Doc. 01–16128 Filed 6–26–01; 8:45 am]

BILLING CODE 3410–03–P

DEPARTMENT OF AGRICULTURE

Agricultural Research Service

Notice of Intent To Grant Exclusive License

AGENCY: Agricultural Research Service, USDA.

ACTION: Notice of intent.

SUMMARY: Notice is hereby given that the U.S. Department of Agriculture, Agricultural Research Service, intends to grant to Triple ''F,'' Inc. of Des Moines, Iowa, an exclusive license to U.S. Patent Application Serial No. 09/611,615, ''COBY Products and a Process for Their Manufacture,'' filed on July 7, 2000. Notice of Availability for U.S. Patent Application Serial No. 09/611,615 was published in the **Federal Register** on March 13, 2001.

DATES: Comments must be received on or before August 27, 2001.

ADDRESSES: Send comments to: USDA, ARS, Office of Technology Transfer, 5601 Sunnyside Avenue, Rm. 4–1158, Beltsville, Maryland 20705–5131.

FOR FURTHER INFORMATION CONTACT: June Blalock of the Office of Technology Transfer at the Beltsville address given above; telephone: 301–504–5257.

SUPPLEMENTARY INFORMATION: The Federal Government's patent rights in this invention are assigned to the United States of America, as represented by the Secretary of Agriculture. It is in the public interest to so license this invention as Triple ''F,'' Inc. has submitted a complete and sufficient application for a license. The prospective exclusive license will be royalty-bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. The prospective exclusive license may be granted unless, within sixty (60) days from the date of this published Notice, the Agricultural Research Service receives written evidence and argument which establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR 404.7.

Michael D. Ruff,

Assistant Administrator.

[FR Doc. 01–16129 Filed 6–26–01; 8:45 am]

BILLING CODE 3410–03–P

DEPARTMENT OF AGRICULTURE

Agricultural Research Service

Notice of Federal Invention Available for Licensing and Intent To Grant Exclusive License

AGENCY: Agricultural Research Service, USDA.

ACTION: Notice of availability and intent.

SUMMARY: Notice is hereby given that the thickspike wheatgrass variety designated ''Bannock'' is available for licensing and that the Department of Agriculture, Agricultural Research Service, intends to grant to the University of Idaho of Moscow, Idaho, an exclusive license to this variety.

DATES: Comments must be received on or before September 25, 2001.

ADDRESSES: Send comments to: USDA, ARS, Office of Technology Transfer, 5601 Sunnyside Avenue, Room 4–1158, Beltsville, Maryland 20705–5131.

FOR FURTHER INFORMATION CONTACT: June Blalock of the Office of Technology Transfer at the Beltsville address given above; telephone: 301–504–5257.

SUPPLEMENTARY INFORMATION: The Federal Government's intellectual property rights to this invention are assigned to the United States of America, as represented by the Secretary of Agriculture. The prospective exclusive license will be royalty-bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. The prospective exclusive license may be granted unless, within ninety (90) days from the date of this published Notice, Agricultural Research Service receives written evidence and argument which establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR 404.7.

Michael D. Ruff,

Assistant Administrator.

[FR Doc. 01–16132 Filed 6–26–01; 8:45 am]

BILLING CODE 3410–03–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. 01–051–1]

Availability of a Draft Environmental Assessment for Field Testing Avian Encephalomyelitis-Fowl Pox-Laryngotracheitis Vaccine, Live Virus, Fowl Pox Vector

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 draft environmental assessment concerning authorization to ship for the purpose of field testing, and then to field test, an unlicensed avian encephalomyelitis-fowl pox-laryngotracheitis vaccine for use in poultry. 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, 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 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 licensure.

DATES: We invite you to comment on this docket. We will consider all comments that we receive by July 27, 2001.

ADDRESSES: Please send four copies of your comment (an original and three copies) to: Docket No. 01–051–1, 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. 01–051–1.

Copies of the draft environmental assessment may be obtained by contacting the person listed under **FOR**

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