(AMS) activities under authority of 7 CFR 250, regulations for the Donation of Food for Use in the United States, Its Territories and Possessions and Areas Under Its Jurisdiction, AMS will use a customer driven approach to maintain and improve the quality of food products and packaging. AMS will use AMS-11, "Customer Opinion Postcard," to collect information. Customers that use USDA procured commodities to prepare and serve meals retrieve these cards from the boxes and use them to rate their perception of product flavor, texture, and appearance as well as overall satisfaction.

Need and Use of the Information: AMS will collect information on the product type, production lot, and identify the location and type of facility in which the product was served. Without this information, AMS will not be able to obtain timely and accurate information about its products from customers that use them.

Description of Respondents: State, Local or Tribal Government; not-forprofit institutions.

Number of Respondents: 8,400. Frequency of Responses: Reporting: On occasion.

Total Burden Hours: 700.

Grain Inspection, Packers & Stockyards Administration

Title: "Clear Title"—Protection for Purchasers of Farm Products. OMB Control Number: 0580–0016. Summary of Collection: Grain Inspection, Packers and Stockyards Administration (GIPSA) have the responsibility for the Clear Title Program (Section 1324 of the Food Security Act of 1985). The Clear Title Program was enacted to facilitate interstate commerce in farm products and protect purchasers of farm products by enabling States to establish central filing systems. The Food Security Act of 1985 permits the states to establish "central filing systems". These central filing systems notify buyers of farm products of any mortgages or liens on the products. There are 19 states that currently have certified central filing systems.

Need and Use of the Information: A state submits information one time to GIPSA when applying for certification. GIPSA reviews the information submitted by the states to certify that those central filing systems meet the criteria set forth in section 1324 of the Food Security Act of 1985.

Description of Respondents: Business or other for-profit.

Number of Respondents: 1. Frequency of Responses: Reporting: On occasion. Total Burden Hours: 80.

Sondra Blakev,

Departmental Information Collection Clearance Officer.

[FR Doc. 04–22981 Filed 10–18–04; 8:45 am] BILLING CODE 3410–05–M; 3410–34–M; 3410–30–M; 3410–11–M; 3410–20–M; 3410–02–M; 3410–KD–M

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. 04-075-1]

Monsanto Co. and KWS SAAT AG; Availability of Petition and Environmental Assessment for Determination of Nonregulated Status for Sugar Beet Genetically Engineered for Tolerance to the Herbicide Glyphosate

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 received a petition from Monsanto Company and KWS SAAT AG seeking a determination of nonregulated status for sugar beet designated as event H7-1, which has been genetically engineered for tolerance to the herbicide glyphosate. The petition has been submitted in accordance with our regulations concerning the introduction of certain genetically engineered organisms and products. In accordance with those regulations, we are soliciting public comments on whether this sugar beet presents a plant pest risk. We are also making available for public comment an environmental assessment for the proposed determination of nonregulated status.

DATES: We will consider all comments we receive on or before December 20, 2004.

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

Postal Mail/Commercial Delivery: Please send four copies of your comment (an original and three copies) to Docket No. 04–075–1, Regulatory Analysis and Development, PPD, APHIS, Station 3C71, 4700 River Road Unit 118, Riverdale, MD 20737–1238. Please state that your comment refers to Docket No. 04–075–1.

E-mail: Address your comment to regulations@aphis.usda.gov. Your comment must be contained in the body of your message; do not send attached files. Please include your name and

address in your message and "Docket No. 04–075–1" on the subject line.

Agency Web site: Go to http://www.aphis.usda.gov/ppd/rad/cominst.html for a form you can use to submit an e-mail comment through the APHIS Web site.

Federal eRulemaking Portal: Go to http://www.regulations.gov and follow the instructions for locating this docket and submitting comments.

Reading Room: You may read the petition, the environmental assessment, and any comments that we receive on this docket 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.

Other Information: You may view APHIS documents published in the **Federal Register** and related information, including the names of groups and individuals who have commented on APHIS dockets, on the Internet at http://www.aphis.usda.gov/ppd/rad/webrepor.html.

FOR FURTHER INFORMATION CONTACT: Mr. John Cordts, Biotechnology Regulatory Services, APHIS, 4700 River Road Unit 147, Riverdale, MD 20737–1236; (301) 734–5531. To obtain copies of the petition or the environmental assessment (EA), contact Ms. Terry Hampton at (301) 734–5715; e-mail: Terry.A.Hampton@aphis.usda.gov. The petition and the EA are also available on the Internet at: http://www.aphis.usda.gov/brs/aphisdocs/03_32301p.pdf and http://www.aphis.usda.gov/brs/aphisdocs/03_32301p_ea.pdf.

SUPPLEMENTARY INFORMATION: The regulations in 7 CFR part 340, "Introduction of Organisms and Products Altered or Produced Through Genetic Engineering Which Are Plant Pests or Which There Is Reason to Believe Are Plant Pests," regulate, among other things, the introduction (importation, interstate movement, or release into the environment) of organisms and products altered or produced through genetic engineering that are plant pests or that there is reason to believe are plant pests. Such genetically engineered organisms and products are considered "regulated articles.'

The regulations in § 340.6(a) provide that any person may submit a petition to the Animal and Plant Health Inspection Service (APHIS) seeking a determination that an article should not be regulated under 7 CFR part 340. Paragraphs (b) and (c) of § 340.6 describe the form that a petition for a determination of nonregulated status must take and the information that must be included in the petition.

On November 19, 2003, APHIS received a petition (APHIS Petition Number 03–323–01p) from Monsanto Company of St. Louis, MO, and KWS SAAT AG of Einbeck, Germany (Monsanto/KWS), requesting a determination of nonregulated status under 7 CFR part 340 for sugar beet (Beta vulgaris ssp. vulgaris) designated as event H7-1, which has been genetically engineered for tolerance to the herbicide glyphosate. The Monsanto/KWS petition states that the subject sugar beet should not be regulated by APHIS because it does not present a plant pest risk. As described in the petition, sugar beet event H7-1 has been genetically engineered to express a 5-enolpyruvyshikimate-3phosphate synthase protein from Agrobacterium sp. strain CP4 (CP4 EPSPS), which confers tolerance to the herbicide glyphosate. Expression of the added genes is controlled in part by the 35S promoter derived from the plant pathogen figwort mosaic virus. The Agrobacterium tumefaciens transformation method was used to transfer the added genes into the KWS proprietary sugar beet line 3S0057.

Sugar beet event H7–1 has been considered a regulated article under the regulations in 7 CFR part 340 because it contains gene sequences from plant pathogens. In the process of reviewing the notifications for field trials of the subject sugar beet, APHIS determined that the vectors and other elements were disarmed and that the trials, which were conducted under conditions of reproductive and physical confinement or isolation, would not present a risk of plant pest introduction or dissemination.

In § 403 of the Plant Protection Act (7 U.S.C. 7701–7772), plant pest is defined as any living stage of any of the following that can directly or indirectly injure, cause damage to, or cause disease in any plant or plant product: A protozoan, a nonhuman animal, a parasitic plant, a bacterium, a fungus, a virus or viroid, an infectious agent or other pathogen, or any article similar to or allied with any of the foregoing. APHIS views this definition very broadly. The definition covers direct or indirect injury, disease, or damage not just to agricultural crops, but also to plants in general, for example, native species, as well as to organisms that

may be beneficial to plants, for example, honeybees, rhizobia, etc.

The U.S. Environmental Protection Agency (EPA) is responsible for the regulation of pesticides under the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), as amended (7 U.S.C. 136 et seq.). FIFRA requires that all pesticides, including herbicides, be registered prior to distribution or sale, unless exempt by EPA regulation. In cases in which genetically modified plants allow for a new use of a pesticide or involve a different use pattern for the pesticide, EPA must approve the new or different use. Accordingly, EPA has granted a registration for the use of glyphosate on glyphosate-tolerant sugar beet.

When the use of the pesticide on the genetically modified plant would result in an increase in the residues in a food or feed crop for which the pesticide is currently registered, or in new residues in a crop for which the pesticide is not currently registered, establishment of a new tolerance or a revision of the existing tolerance would be required. Residue tolerances for pesticides are established by EPA under the Federal Food, Drug, and Cosmetic Act (FFDCA), as amended (21 U.S.C. 301 et seq.), and the Food and Drug Administration (FDA) enforces tolerances set by EPA under the FFDCA. EPA has determined that the existing residue tolerance for glyphosate-tolerant sugar beet is sufficient to support future use of glyphosate on event H7-1.

FDA published a statement of policy on foods derived from new plant varieties in the Federal Register on May 29, 1992 (57 FR 22984-23005). The FDA statement of policy includes a discussion of FDA's authority for ensuring food safety under the FFDCA, and provides guidance to industry on the scientific considerations associated with the development of foods derived from new plant varieties, including those plants developed through the techniques of genetic engineering. Monsanto/KWS has begun consultation with FDA on the subject sugar beet event.

To provide the public with documentation of APHIS' review and analysis of the environmental impacts and plant pest risk associated with a proposed determination of nonregulated status for the Monsanto/KWS event H7–1 sugar beet, an environmental assessment (EA) has been prepared. The EA was 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).

In accordance with § 340.6(d) of the regulations, we are publishing this notice to inform the public that APHIS will accept written comments regarding the petition for a determination of nonregulated status from interested persons for a period of 60 days from the date of this notice. We are also soliciting written comments from interested persons on the EA prepared to examine any environmental impacts of the proposed determination for the subject sugar beet event. The petition and the EA and any comments received are available for public review, and copies of the petition and the EA are available as indicated in the FOR FURTHER **INFORMATION CONTACT** section of this notice.

After the comment period closes, APHIS will review the data submitted by the petitioner, all written comments received during the comment period, and any other relevant information. After reviewing and evaluating the comments on the petition and the EA and other data and information, APHIS will furnish a response to the petitioner, either approving the petition in whole or in part, or denying the petition. APHIS will then publish a notice in the Federal Register announcing the regulatory status of the Monsanto/KWS glyphosate-tolerant sugar beet event H7-1 and the availability of APHIS' written decision.

Authority: 7 U.S.C. 1622n and 7701–7772; 31 U.S.C. 9701; 7 CFR 2.22, 2.80, and 371.3.

Done in Washington, DC, this 14th day of October 2004.

Kevin Shea,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. E4–2710 Filed 10–18–04; 8:45 am] **BILLING CODE 3410–34–P**

DEPARTMENT OF AGRICULTURE

Forest Service

Madera County Resource Advisory Committee

AGENCY: Forest Service, USDA. **ACTION:** Notice of Resource Advisory Committee Meeting.

SUMMARY: Pursuant to the authorities in the Federal Advisory Committee Act of 1972 (Pub. L. 92–463) and under the secure Rural Schools and Community Self-Determination Act of 2000 (Pub. L. 106–393) the Sierra National Forest's