

the petition must contain sufficient facts to merit an FDA investigation. (See H. Rept. 857, part 1, 98th Cong., 2d sess., pp. 41–42, 1984.) Petitions should be in the format specified in 21 CFR 10.30.

Comments and petitions should be submitted to the Division of Dockets Management. Three copies of any mailed information are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket numbers found in brackets in the heading of this document. Comments and petitions may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: March 22, 2010.

**Jane A. Axelrad,**

*Associate Director for Policy, Center for Drug Evaluation and Research.*

[FR Doc. 2010–9509 Filed 4–22–10; 8:45 am]

**BILLING CODE 4160–01–S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA–2010–N–0159]

#### North American Bioproducts Corp.; Filing of Food Additive Petition (Animal Use); Erythromycin Thiocyanate

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that North American Bioproducts Corp. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of erythromycin thiocyanate as an antimicrobial processing aid in fuel-ethanol fermentations with respect to its consequent presence in by-product distiller grains used as an animal feed or feed ingredient.

**DATES:** Submit written or electronic comments on the petitioner's environmental assessment May 24, 2010.

**ADDRESSES:** You may submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.regulations.gov>.

**FOR FURTHER INFORMATION CONTACT:** Isabel W. Pocurull, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl.,

Rockville, MD 20855, 240–453–6853, email: [isabel.pocurull@fda.hhs.gov](mailto:isabel.pocurull@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** Under the Federal Food, Drug, and Cosmetic Act (section 409(b)(5) (21 U.S.C. 348(b)(5)), notice is given that a food additive petition (FAP 2263) has been filed by North American Bioproducts Corp., Corporate Support Center, 1815 Satellite Blvd., Building 200, Duluth, GA 30097. The petition proposes to amend the food additive regulations in 21 CFR Part 573 *Food Additives Permitted in Feed and Drinking Water of Animals* to provide for the safe use of erythromycin thiocyanate as an antimicrobial processing aid in fuel-ethanol fermentations with respect to its consequent presence in by-product distiller grains used as an animal feed or feed ingredient.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Division of Dockets Management (see **ADDRESSES**) for public review and comment.

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) electronic or written comments regarding this document. Submit a single copy of electronic comments or two paper copies of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.51(b).

Dated: April 14, 2010.

**Bernadette Dunham,**

*Director, Center for Veterinary Medicine.*

[FR Doc. 2010–9420 Filed 4–22–10; 8:45 am]

**BILLING CODE 4160–01–S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Substance Abuse and Mental Health Services Administration

#### Fiscal Year (FY) 2010 Funding Opportunity

**AGENCY:** Substance Abuse and Mental Health Services Administration, HHS.

**ACTION:** Notice of intent to award a Single Source Grant to the current grantee for the National Center for Child Traumatic Stress.

**SUMMARY:** This notice is to inform the public that the Substance Abuse and Mental Health Services Administration (SAMHSA) intends to award approximately \$1,000,000 (total costs) for up to one year to the current grantee for the National Center for Child Traumatic Stress (NCCTS). This is not a formal request for applications. Assistance will be provided only to the current grantee for the National Center for Child Traumatic Stress based on the receipt of a satisfactory application that is approved by an independent review group.

*Funding Opportunity Title:* SM–10–016.

*Catalog of Federal Domestic Assistance (CFDA) Number:* 93.243.

*Authority:* Section 582 of the Public Health Service Act, as amended.

*Justification:* Only an application from the current grantee for the National Center for Child Traumatic Stress will be considered for funding under this announcement. One-year funding has become available to assist SAMHSA in responding to data analysis and reporting activities that improve evidence-based practices and raise the standard of trauma care. It is considered most cost-effective and efficient to supplement the existing grantee because they have access to the existing National Child Traumatic Stress Network (NCTSN) datasets and data analytic expertise to conduct the required data analytic activities. There is no other potential organization with the required access and expertise.

Eligibility for this program supplement is restricted to the current grantee, National Center for Child Traumatic Stress in accordance with Congressional intent for 2010 SAMHSA appropriations.

The role of the NCCTS is to provide infrastructure and support for the National Child Traumatic Stress Network to achieve its goals of increasing access and raising the standard of care for traumatized children, adolescents, and their