H. Should any provision of these Official Rules be or become illegal or unenforceable under applicable Federal law, such illegality or unenforceability shall leave the remainder of these Official Rules unaffected and valid. The illegal or unenforceable provision may be replaced by the Sponsor with a valid and enforceable provision that, in the Sponsor's sole judgment, comes closest and best reflects the Sponsor's intention in a legal and enforceable manner with respect to the invalid or unenforceable provision.

13. Disputes

Subject to the release provisions in these Official Rules, Contestant agrees that:

A. Any and all disputes, claims, and causes of action arising out of or connected with this Contest, any Prizes awarded, the administration of the Contest, the determination of Winners, or the construction, validity, interpretation, and enforceability of the Official Rules shall be resolved individually;

B. any and all disputes, claims, and causes of action arising out of or connected with this Contest, any Prizes awarded, the administration of the Contest, the determination of Winners, or the construction, validity, interpretation, and enforceability of the Official Rules shall be resolved pursuant to Federal law;

C. under no circumstances will Contestants be entitled to, and Contestants hereby waive, all rights to claim, any punitive, incidental, and consequential damages and any and all rights to have damages multiplied or otherwise increased.

14. Privacy

The Sponsor may collect personal information from the Contestant when he or she enters the Contest. Such personal information collected is subject to the privacy policy located here: *http://www.ftc.gov/site-information/privacy-policy.*

15. Contact Us

Please visit the Contest Web site for further Contest information and updates.

Jessica Rich,

Director, Bureau of Consumer Protection.

[FR Doc. 2015–05442 Filed 3–6–15; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0510]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Substances Prohibited From Use in Animal Food or Feed

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by April 8, 2015.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0627. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd.; COLE–14526, Silver Spring, MD 20993–0002 *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Substances Prohibited From Use in Animal Food or Feed—21 CFR Part 589 (OMB Control Number 0910–0627—Revision)

This regulation prohibits the use of certain cattle origin materials in the food or feed of all animals to help prevent the spread of bovine spongiform encephalopathy (BSE) in U.S. cattle. BSE is a progressive and fatal neurological disorder of cattle that results from an unconventional transmissible agent. BSE belongs to the family of diseases known as transmissible spongiform encephalopathies (TSEs). All TSEs affect the central nervous system of infected animals. These measures will further strengthen existing safeguards against BSE.

In the **Federal Register** of November 21, 2014 (79 FR 69493), FDA published a 60-day notice requesting public comment on the proposed collection of information. One comment was received however it did not respond to any of the four information collection topics solicited and is therefore not addressed by the Agency.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL RECORDKEEPING BURDEN [1]

21 CFR Section 589.2001; Substances prohibited from use in animal food or feed	Number of recordkeepers	Number of records per recordkeeper	Total annual records	Average burden per recordkeeper	Total hours	Operating and maintenance costs
589.2001(c)(2)(vi) and (c)(3)(i)	175	1	175	20	3,500	$59,500
589.2001(c)(2)(ii)	50	1	50	20	1,000	17,000
589.2001(c)(3)(i)(A)	175	1	175	26	4,550	80,580
Total					9,050	157,080

[1] There are no capital costs associated with this collection of information.

Description of Respondents for Recordkeeping: Rendering facilities, medicated feed manufacturers, livestock feeders.

The Agency's recordkeeping burden estimate was calculated by multiplying the number of recordkeepers times the number of records per recordkeeper to determine the total annual number of records. The total number of annual records were then multiplied by the average burden per recordkeeper to determine the total number of burden hours.

TABLE 2—ESTIMATED ANNUAL REPORTING BURDEN [1]

21 CFR section 589.2001(f)	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per respondent	Total hours
One-time (initial) burden	1	1	1	80	80
Burden from future review	1	1	1	26	26

[1] There are no capital costs or operating costs associated with the collection of information.

Description of Respondents for Reporting: The final regulation on BSE (73 FR 22720) included a provision that exempts cattle materials prohibited in animal feed (CMPAF) from designated countries from the prohibition on its use in animal feed. A foreign country seeking this designation will submit a written request to FDA that includes a variety of information about the country's BSE status (§ 589.2001(f)). During the past 6 years, FDA received 2 requests from countries to be exempted from CMPAF restrictions.

One-Time (Initial) Reporting Burden

There is a one-time burden to countries that apply to FDA seeking to be designated as not subject to restrictions applicable to CMPAF. We estimate that each country that applies for an exclusion will spend 80 hours putting information together to submit to FDA. Table 2 row 1 presents the one-time burden for the exclusion. (See final BSE regulation at 73 FR 22754).

Recurring Burden

Countries that successfully petition FDA to be designated as exempt from certain BSE-related restrictions applicable to animal feed will be subject to future review by FDA to ensure that their designation remains appropriate. As part of this process, FDA may ask designated countries from time-to-time to confirm that their BSE situation and the information submitted by them in support of their original application remains unchanged. We assume it will take FDA and the designated country undergoing a review in the future about one third the time and effort it did when the information was submitted. Table 2 row 2 presents the expected recurring burden.

Dated: March 2, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–05356 Filed 3–6–15; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Advancing Translational Sciences; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Center for Advancing Translational Sciences Special Emphasis Panel Small Business Innovation Research (SBIR).

Date: March 31, 2015.

Time: 8:00 a.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Bethesda Marriott Suites, 6711 Democracy Boulevard, Bethesda, MD 20817.

Contact Person: Rahat Khan, Ph.D., Scientific Review Officer, Office of Scientific Review, National Center for Advancing Translational Sciences, 6701 Democracy Blvd., Rm 1078, Bethesda, MD 20892, 301–894–7319, *khanr2@csr.nih.gov*.

Name of Committee: National Center for Advancing Translational Sciences Special Emphasis Panel CTSA Meeting 1.

Date: April 8–9, 2015.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Hilton Washington/Rockville 1750 Rockville Pike, Rockville, MD 20852.

Contact Person: Carol Lambert, Ph.D., Acting Deputy Director, Office of Grants Management & Scientific Review, National Center for Advancing Translational Sciences (NCATS) National Institutes of Health, 6701 Democracy Boulevard, Democracy 1, Room 1076, Bethesda, MD 20892, 301–435–0814, *lambert@mail.nih.gov*.

Name of Committee: National Center for Advancing Translational Sciences Special Emphasis Panel 2015 CTSA Application Review.

Date: April 15–16, 2015.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Hilton Washington/Rockville 1750 Rockville Pike, Rockville, MD 20852.

Contact Person: Guo He Zhang, Ph.D., MPH, Scientific Review Office, Office of Grants Management & Scientific Review, National Center for Advancing Translational Sciences (NCATS) National Institutes of Health, 6701 Democracy Boulevard, Democracy 1, Room 1064, Bethesda, MD 20892, 301–435–0812, *zhanggu@mail.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.859, Pharmacology, Physiology, and Biological Chemistry Research; 93.350, B–Cooperative Agreements; 93.859, Biomedical Research and Research Training, National Institutes of Health, HHS).

Dated: March 3, 2015.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–05304 Filed 3–6–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request Surveys and Interviews To Support an Evaluation of the Innovative Molecular Analysis Technologies (IMAT) Program (NCI)

SUMMARY: Under the provisions of section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on December 4, 2014, Vol. 79, Page 72004 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Cancer Institute (NCI), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an