administrative supplement requests, change of organization requests, and change of grantee/training institution requests submitted September 25, 2013 and beyond. Multi-project applications that are transitioning to electronic submission beginning with the September 25, 2013 due dates (see the NIH Guide Notice *NOT–OD–13–075*) will also use FORMS–C packages.

Exceptions

The programs noted below will move to FORMS–C application packages as follows:

- Individual Research Career Development Award Programs (Ks), Institutional Training and Career Development Programs (Ts and Ds) and Individual National Research Service Awards (Fs) applicants will be required to use FORMS–C packages for due dates on or after January 25, 2014.
- Small Business programs (SBIR/STTR) applicants will transition to FORMS–C packages later in 2014, so that anticipated form changes relating to the Small Business Authorization Act can be incorporated.

Instructions

- If presented with more than one forms package, applicants should download and use the most recent set of forms to complete their submission.
- Verify you have the correct application package by checking the Competition ID field for FORMS–C. The Competition ID field can be found when downloading the application package from Grants.gov, in the application header information of the downloaded package or in FOA summary information for multi-project applications.
- Learn more about choosing the correct forms packages at: *http://grants.nih.gov/grants/ElectronicReceipt/files/right_forms.pdf.*
- All applicants should carefully read their FOA and the appropriate ''C Series'' Application Guide for program-specific instructions before completing their application.

Inquiries

Please direct all inquiries to: Technical Information Management Section, Procurement and Grants Office, Centers for Disease Control and Prevention, Telephone: 770–488–2700, Email: *pgotim@cdc.gov.*

Dated: July 26, 2013.

William P. Nichols,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 2013–18608 Filed 8–1–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0536]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Medical Device User Fee Cover Sheet, Form FDA 3601

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Medical Device User Fee Cover Sheet, Form FDA 3601'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Daniel Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, *Daniel.Gittleson@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: On February 8, 2013, the Agency submitted a proposed collection of information entitled ''Medical Device User Fee Cover Sheet, Form FDA 3601'' to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0511. The approval expires on April 30, 2016. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: July 29, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–18638 Filed 8–1–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0868]

Agency Information Collection Activities; Proposed Collection; Comment Request; Draft Guidance for Industry: Use of Serological Tests To Reduce the Risk of Transmission of Trypanosoma cruzi Infection in Whole Blood and Blood Components for Transfusion

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments concerning establishment notification of a consignee and consignee notification of a recipient's physician of record regarding a possible increased risk of *Trypanosoma cruzi (T. cruzi)* infection.

DATES: Submit either electronic or written comments on the collection of information by October 1, 2013.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov.* Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–7726, *Ila.mizrachi@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests