of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

**Animal Drug User Fee Cover Sheet—OMB Control Number 0910–0539—Extension**

Under section 740 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 379j–12), FDA has the authority to assess and collect application fees from each person who submits certain new animal drug applications or certain supplemental animal drug applications. The Animal Drug User Fee cover sheet (Form FDA 3546) is designed to collect the minimum necessary information to determine whether a fee is required for the review of an application or supplement or whether an application fee waiver was granted, to determine the amount of the fee required, and to assure that each animal drug user fee payment is appropriately linked to the animal drug application for which payment is made. The form, when completed electronically, will result in the generation of a unique payment identification number used by FDA to track the payment. FDA's Center for Veterinary Medicine and FDA's Office of Management will use the information collected to initiate the administrative screening of new animal drug applications and supplements to determine whether the payment has been received.

*Description of Respondents:* Respondents to this collection of information are new animal drug applicants.

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

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

| FD&C Act section; description | FDA form No. | Number of respondents | Number of responses per respondent | Total annual responses | Average burden per response | Total hours |
|---|---|---|---|---|---|---|
| 740(a)(1); Animal Drug User Fee cover sheet. | FDA 3546 | 21 | 1 | 21 | 1 | 21 |

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

The estimates in table 1 are based on our experience with new animal drug applications and supplemental animal drug applications and the average number of Animal Drug User Fee cover sheets submitted during fiscal years 2013–2015. We estimate 21 respondents will each submit a cover sheet (Form FDA 3546), for a total of 21 responses. We calculate a reporting burden of 1 hour per response, for a total of 21 hours.

Dated: October 17, 2016.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2016–25482 Filed 10–20–16; 8:45 am]

**BILLING CODE 4164–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Office of the Secretary

### Notice of Interest Rate on Overdue Debts

Section 30.18 of the Department of Health and Human Services' claims collection regulations (45 CFR part 30) provides that the Secretary shall charge an annual rate of interest, which is determined and fixed by the Secretary of the Treasury after considering private consumer rates of interest on the date that the Department of Health and Human Services becomes entitled to recovery. The rate cannot be lower than the Department of Treasury's current value of funds rate or the applicable rate determined from the ''Schedule of Certified Interest Rates with Range of Maturities'' unless the Secretary waives interest in whole or part, or a different rate is prescribed by statute, contract, or repayment agreement. The Secretary of the Treasury may revise this rate quarterly. The Department of Health and Human Services publishes this rate in the **Federal Register**.

The current rate of 9⅝%, as fixed by the Secretary of the Treasury, is certified for the quarter ended September 30, 2016. This rate is based on the Interest Rates for Specific Legislation, ''National Health Services Corps Scholarship Program (42 U.S.C. 254o(b)(1)(A))'' and ''National Research Service Award Program (42 U.S.C. 288(c)(4)(B)).'' This interest rate will be applied to overdue debt until the Department of Health and Human Services publishes a revision.

Dated: October 13, 2016.

**David C. Horn,**

*Director, Office of Financial Policy and Reporting.*

[FR Doc. 2016–25459 Filed 10–20–16; 8:45 am]

**BILLING CODE 4150–04–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Meeting Announcement for the Technical Advisory Panel on Medicare Trustee Reports

**AGENCY:** Department of Health and Human Services (HHS).

**ACTION:** Notice of public meeting.

**SUMMARY:** This notice announces the meeting dates for the Technical Advisory Panel on Medicare Trustee Reports on Monday, October 31, 2016 and Tuesday, November 1, 2016 in Washington, DC.

**DATES:** The meeting will be held on Monday, October 31, 2016 from 9:30 a.m. to 5:00 p.m. and Tuesday, November 1, 2016, from 9:00 a.m. to 3:30 p.m. Eastern Daylight Time (EDT) and it is open to the public.

**ADDRESSES:** The meeting will be held at the Hubert Humphrey Building, 200 Independence Ave. SW., Washington, DC, 20201 Room 738G.3.

**FOR FURTHER INFORMATION CONTACT:** Dr. Donald Oellerich, Designated Federal Officer, at the Office of Human Services Policy, Assistant Secretary for Planning and Evaluation, U.S. Department of Health and Human Services, 200 Independence Ave. SW., Washington, DC 20201, (202) 690–8410.

**SUPPLEMENTARY INFORMATION:**