

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2015-D-4803]

Public Notification of Emerging Postmarket Medical Device Signals ('Emerging Signals'); Draft Guidance for Industry and Food and Drug Administration Staff; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA or Agency) is extending the comment period for the draft guidance for Industry and Food and Drug Administration Staff entitled "Public Notification of Emerging Postmarket Medical Device Signals ('Emerging Signals')." A notice of the availability of the draft guidance and our request for comments appeared in the **Federal Register** of December 31, 2015. We initially established February 29, 2016, as the deadline for the submission of requested comments that can help improve the Agency's policy for notifying the public about medical device "emerging signals." The Agency is taking this action due to the unanticipated high-level of interest from external stakeholders and the medical device community and will allow interested persons additional time to submit comments.

DATES: FDA is extending the comment period on the "Public Notification of Emerging Postmarket Medical Device Signals ('Emerging Signals'); Draft Guidance for Industry and Food and Drug Administration Staff; Availability," which was announced in the Notice published December 31, 2015 (80 FR 81829). Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the Agency considers your comment of this draft guidance before it begins work on the final version of the guidance, submit either electronic or written comments on the draft guidance by March 29, 2016.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically,

including attachments, to <http://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <http://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA-2015-D-4803 for "Public Notification of Emerging Postmarket Medical Device Signals ('Emerging Signals')." Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <http://www.regulations.gov> or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information

redacted/blacked out, will be available for public viewing and posted on <http://www.regulations.gov>. Submit both copies to the Division of Dockets Management. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <http://www.fda.gov/regulatoryinformation/dockets/default.htm>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <http://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

An electronic copy of the draft guidance document is available for download from the Internet. See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the guidance. Submit written requests for a single hard copy of the draft guidance document entitled "Public Notification of Emerging Postmarket Medical Device Signals ('Emerging Signals')." to the Office of the Center Director, Guidance and Policy Development, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5431, Silver Spring, MD 20993-0002. Send one self-addressed adhesive label to assist that office in processing your request.

FOR FURTHER INFORMATION CONTACT:

Rebecca Nipper, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1540, Silver Spring, MD 20993-0002, 301-796-6527.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of December 31, 2015, FDA published a notice announcing the availability of a draft guidance entitled "Public Notification of Emerging Postmarket Medical Device Signals ('Emerging Signals')." with a 60-day comment period to request comments on the Agency's policy for

notifying the public about medical device “emerging signals.”

FDA is extending the comment period for the publication notification of “emerging signals” for 30 days, until March 29, 2016. The Agency believes that a 30-day extension allows adequate time for interested persons to submit comments.

II. Electronic Access

Persons interested in obtaining a copy of the draft guidance may do so by downloading an electronic copy from the Internet. A search capability for all Center for Devices and Radiological Health guidance documents is available at <http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/GuidanceDocuments/default.htm>. Guidance documents are also available at <http://www.regulations.gov>. Persons unable to download an electronic copy of “Public Notification of Emerging Postmarket Medical Device Signals (‘Emerging Signals’)” may send an email request to CDRH-Guidance@fda.hhs.gov to receive an electronic copy of the document. Please use the document number 1500027 to identify the guidance you are requesting.

Dated: January 21, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016-01610 Filed 1-26-16; 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 9¾%, as fixed by the Secretary of the Treasury, is certified for the quarter ended December 31, 2015. 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: January 13, 2016.

David C. Horn.

Director, Office of Financial Policy and Reporting.

[FR Doc. 2016-01649 Filed 1-26-16; 8:45 am]

BILLING CODE 4150-04-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[CMS-9935-N2]

HHS-Operated Risk Adjustment Methodology Meeting; March 31, 2016

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Notice of meeting.

SUMMARY: This notice announces the rescheduling of the March 25, 2016 meeting on the HHS-operated risk adjustment program, which is open to the public. The purpose of this stakeholder meeting is to solicit feedback on the HHS-operated risk adjustment methodology and to discuss potential improvements to the HHS risk adjustment methodology for the 2018 benefit year and beyond. This meeting, the “HHS-operated Risk Adjustment Methodology Conference,” will allow issuers, States, and other interested parties to discuss the contents of a White Paper to be published in advance of this meeting. This meeting will also provide an opportunity for participants to ask clarifying questions. The comments and information HHS obtains through this meeting may be used in future policy making for the HHS risk adjustment program.

DATES: *Date of Meeting:* March 31, 2016 from 9:00 a.m. to 4:30 p.m., Eastern daylight time (e.d.t.).

Deadline for Onsite Participation: March 23, 2016, 5:00 p.m., e.d.t.

Deadline for Webinar Meeting Participation: March 28, 2016, 5:00 p.m. e.d.t.

Deadline for Requesting Special Accommodations: March 23, 2016, 5:00 p.m. e.d.t.

ADDRESSES: The meeting will be held at the CMS Single Site campus, 7500

Security Boulevard, Baltimore, MD 21244.

Registration: Registration will be on a first-come, first-serve basis, limited to two (2) participants per organization for the onsite location participation, and three (3) participants per organization for the webinar participation. Each individual can only register for either the onsite location participation or webinar participation. To change a registration option from onsite to webinar participation, the registrant must cancel the existing registration (onsite or webinar) before attempting to register for the other option.

Registration Instructions: To register to attend the meeting either onsite or through webinar participation, visit the Registration for Technical Assistance Portal (REGTAP) at www.REGTAP.info. If not already a REGTAP user, register as a new user, log in and go to “My Dashboard” and select “Training Events” to register for the onsite or webinar event for the HHS-operated Risk Adjustment Methodology Meeting. Registrants can only register to attend the meeting onsite at CMS or remotely by webinar.

FOR FURTHER INFORMATION CONTACT: For further information, please send inquiries about the logistics of the meeting to registrar@REGTAP.info. Users should submit inquiries and comments pertaining to content covered during the meeting to www.REGTAP.info. To submit an inquiry in REGTAP, select “Submit an Inquiry” from “My Dashboard” then select “HHS-operated Risk Adjustment Methodology Meeting” from the Event Title dropdown menu and enter the question or comment. Users can submit their comments and upload attachments as needed. REGTAP will send the user an acknowledgement upon receipt of the comment.

The CCIIO’s Press Office at (202) 690-6145 will handle all press inquiries.

SUPPLEMENTARY INFORMATION:

I. Background

This notice announces a meeting on the HHS-operated risk adjustment program to discuss potential improvements to the HHS risk adjustment methodology for the 2018 benefit year and beyond. This meeting will focus on the permanent risk adjustment program under section 1343 of the Affordable Care Act when HHS is operating a risk adjustment program on behalf of a State (referred to as the HHS-operated risk adjustment program).

We are committed to stakeholder engagement in developing the detailed processes of the HHS-operated risk