

circumstances or conditions related to the subject accounts; or failure to satisfy conditions applicable to each. The following exception was granted by the FDIC as of February 4, 2021.

I. Mortgage Servicing Accounts for Which the Covered Institution’s System of Record Cannot Calculate Principal and Interest at an Account Level at a Given Point in Time

The FDIC granted time-limited exception relief to covered institutions up to March 31, 2022, from the information technology system requirements of 12 CFR 370.3 and the recordkeeping requirements of 12 CFR 370.4 for principal and interest payments held in mortgage servicing accounts for which the covered institutions act as servicers or sub-servicers. The recommended relief will provide the covered institutions additional time to remediate their servicing platforms and internal processing capabilities pending further direction from the FDIC.

Pursuant to 12 CFR 330.7(d), mortgage principal and interest payments are insured for the cumulative balance paid into the account by the mortgagors, up to the limit of the standard maximum deposit insurance amount per mortgagor. If a covered institution does not maintain deposit records that enable it to calculate deposit insurance, the covered institution must maintain, at a minimum, the following in its deposit account records: (i) The unique identifier of the account holder; and (ii) the corresponding “pending reason”

code listed in pending file format set forth in Appendix B to Part 370.

The covered institutions service the mortgage loans using platforms hosted by third party vendors. Principal and interest payments from mortgagors are placed into the mortgage servicing accounts with the funds held in custody for the investors that own the underlying mortgages. Because the loans are tracked and managed as a group by pool, the servicing platforms do not have a mechanism to allocate the mortgage servicing accounts balances to specific mortgagors. As a result, the covered institutions do not have a process to input mortgagor principal and interest data into their information technology systems to calculate deposit insurance coverage for the mortgage servicing accounts.

Remediation efforts are underway and include the development of a business requirements document, system updates, implementation, and testing. However, a number of the covered institutions have asked the FDIC for additional clarification of the part 370 recordkeeping rule with respect to the mortgage servicing accounts to determine how to produce borrower account level principal and interest data on a date of failure. Given the complexities of payments to investors under the agreements with the covered institutions, additional information from the FDIC is needed to finalize programming logic and various business requirements documents between the Banks and their service providers.

The FDIC’s grant of relief is subject to the condition that each covered

institution must submit within 60 days, upon receipt of additional information from the FDIC with respect to the part 370 processing for the mortgage servicing account ownership right and capacity code, a status report setting forth the project plan and timeline for integrating the mortgage servicing account ownership right and capacity code processing capabilities into the covered institution’s information technology system.

The FDIC reserves the right to rescind or modify the grant of relief upon any material change of circumstances or conditions related to the accounts subject to this request.

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on February 5, 2021.

James P. Sheesley,

Assistant Executive Secretary.

[FR Doc. 2021–02781 Filed 2–10–21; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Notice of Termination of Receiverships

The Federal Deposit Insurance Corporation (FDIC or Receiver), as Receiver for each of the following insured depository institutions, was charged with the duty of winding up the affairs of the former institutions and liquidating all related assets. The Receiver has fulfilled its obligations and made all dividend distributions required by law.

NOTICE OF TERMINATION OF RECEIVERSHIPS

Fund	Receivership name	City	State	Termination date
10152	The Buckhead Community Bank	Atlanta	GA	02/01/2021
10245	Sun West Bank	Las Vegas	NV	02/01/2021
10277	Palos Bank And Trust Company	Palos Heights	IL	02/01/2021
10280	Imperial Savings & Loan Association	Martinsville	VA	02/01/2021
10502	Valley Bank	Moline	IL	02/01/2021

The Receiver has further irrevocably authorized and appointed FDIC-Corporate as its attorney-in-fact to execute and file any and all documents that may be required to be executed by the Receiver which FDIC-Corporate, in its sole discretion, deems necessary, including but not limited to releases, discharges, satisfactions, endorsements, assignments, and deeds. Effective on the termination dates listed above, the Receiverships have been terminated, the Receiver has been discharged, and the

Receiverships have ceased to exist as legal entities.

(Authority: 12 U.S.C. 1819)

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on February 5, 2021.

James P. Sheesley,

Assistant Executive Secretary.

[FR Doc. 2021–02783 Filed 2–10–21; 8:45 am]

BILLING CODE 6714–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0173]

Vaccines and Related Biological Products Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA or Agency) announces a forthcoming public advisory committee meeting of the Vaccines and Related Biological Products Advisory Committee (the committee). The general function of the committee is to provide advice and recommendations to FDA on regulatory issues. The meeting will be open to the public. FDA is establishing a docket for public comment on this document. Consistent with FDA's regulations, this notice is being published with less than 15 days prior to the date of the meeting based on a determination that convening a meeting of the Vaccines and Related Biological Products Advisory Committee as soon as possible is warranted. This **Federal Register** notice could not be published 15 days prior to the date of the meeting due to a recent submission by Janssen Biotech Inc. of a request for emergency use authorization (EUA) for an investigational vaccine to prevent Coronavirus Disease 2019 (COVID-19) and the need for prompt discussion of such submission, given the COVID-19 pandemic.

DATES: The meeting will be held on February 26, 2021, from 9 a.m. Eastern Time to 5:30 p.m. Eastern Time.

ADDRESSES: Please note that due to the impact of this COVID-19 pandemic, all meeting participants will be joining this advisory committee meeting via an online teleconferencing platform. Answers to commonly asked questions including information regarding special accommodations due to a disability may be accessed at: <https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm>. The online web conference meeting will be available at the following link on the day of the meeting: <https://youtu.be/Qd7mlCD-rEA>.

FDA is establishing a docket for public comment on this meeting. The docket number is FDA-2021-N-0173. The docket will close on February 25, 2021. Submit either electronic or written comments on this public meeting by February 25, 2021. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before February 25, 2021. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of February 25, 2021. Comments received by mail/hand delivery/courier (for

written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Comments received on or before February 18, 2021, will be provided to the committee. Comments received after February 18, 2021, and by February 25, 2021, will be taken into consideration by FDA. In the event that the meeting is cancelled, FDA will continue to evaluate any relevant applications, submissions, or information, and consider any comments submitted to the docket, as appropriate.

You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://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 <https://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):** Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, 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-2021-N-0173 for "Vaccines and Related

Biological Products; Notice of Meeting; Establishment of a Public Docket; Request for Comments." Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <https://www.regulations.gov> or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240-402-7500.

- **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." FDA 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 <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. 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 the 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: <https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <https://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 Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240-402-7500.

FOR FURTHER INFORMATION CONTACT: Prabhakara Atreya or Kathleen Hayes, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6306, Silver Spring, MD 20993-0002, CBERVRBPAC@fda.hhs.gov; or FDA Advisory Committee Information Line, 1-800-

741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's website at <https://www.fda.gov/advisory-committees> and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before joining the meeting.

SUPPLEMENTARY INFORMATION:

Agenda: The meeting presentations will be heard, viewed, captioned, and recorded through an online teleconferencing platform. The committee will meet in open session to discuss EUA of the Janssen Biotech Inc. COVID–19 Vaccine for active immunization to prevent COVID–19 caused by SARS–CoV–2 in individuals 18 years and older. EUA authority allows FDA to help strengthen the nation's public health protections against chemical, biological, radiological, nuclear (CBRN) threats by facilitating the availability and use of Medical Countermeasures (MCMs) needed during public health emergencies. Under section 564 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360bbb–3), FDA may allow unapproved medical products or unapproved uses of approved medical products to be used in an emergency to diagnose, treat, or prevent serious or life-threatening diseases or conditions caused by CBRN threat agents when certain statutory criteria have been met, including that there are no adequate, approved, and available alternatives. Additional information about EUAs can be found at <https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization>.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its website prior to the meeting, background material will be made publicly available on FDA's website at the time of the advisory committee meeting. Background material and the link to the online teleconference meeting room will be available at <https://www.fda.gov/advisory-committees/advisory-committee-calendar>. Scroll down to the appropriate advisory committee meeting link. The meeting will include slide presentations with audio components to

allow the presentation of materials in a manner that most closely resembles an in-person advisory committee meeting.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. All electronic and written submissions submitted to the Docket (see **ADDRESSES**) on or before February 18, 2021, will be provided to the committee. Comments received after February 18, 2021, and by February 25, 2021, will be taken into consideration by FDA. Oral presentations from the public will be scheduled between approximately 1:25 p.m. Eastern Time and 2:25 p.m. Eastern Time. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before February 17, 2021. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by February 18, 2021.

For press inquiries, please contact the Office of Media Affairs at fdaoma@fda.hhs.gov or 301–796–4540.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with disabilities. If you require accommodations due to a disability, please contact Prabhakara Atreya or Kathleen Hayes (CBERVBPAC@fda.hhs.gov) at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our website at: <https://www.fda.gov/advisory-committees/about-advisory-committees/public-conduct-during-fda-advisory-committee-meetings> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: February 8, 2021.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2021–02845 Filed 2–10–21; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Public Comment Request; Information Collection Request Title: Health Professions Student Loan Program, Loans for Disadvantaged Students, Primary Care Loan Program, and Nursing Student Loan Program Administrative Requirements. OMB No. 0915–0047—Revision

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, HRSA submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period. OMB may act on HRSA's ICR only after the 30 day comment period for this notice has closed.

DATES: Comments on this ICR should be received no later than March 15, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting "Currently under Review—Open for Public Comments" or by using the search function.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email Lisa Wright-Solomon, the HRSA Information Collection Clearance Officer at paperwork@hrsa.gov or call (301) 443–1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Health Professions Student Loan (HPSL) Program, Loans for Disadvantaged Students, Primary Care Loan Program (PCL), and Nursing Student Loan Program Administrative Requirements.

OMB No. 0915–0047—Revision

Abstract: This clearance request is for approval of the Health Professions Student Loan (HPSL) Program, Loans for Disadvantaged Students (LDS), Primary Care Loan Program (PCL), and Nursing Student Loan (NSL) Program