213, WC Docket No. 20–89 (FCC 20–44), establishing two programs designed to assist health care providers in providing connected care services to consumers—the COVID–19 Telehealth Program and the Connected Care Pilot Program (collectively, Programs). June 2021, the Commission adopted a Second Report and Order, WC Docket No. 18–213 (FCC 21–74), that provided guidance on eligible services, competitive bidding, invoicing, and data reporting for Pilot Program participants. The information collected herein is necessary to meet the specific requirements for information that must be submitted as part of the annual and final reports to the Commission as outlined in the *Second Connected Care Report and Order,* and for the Commission to receive and evaluate data for the selected projects and ensure compliance with the Commission's rules and procedures applicable to the Connected Care Pilot Program. This submission does not make any changes to the previously approved information collections for the COVID–19 Telehealth Program and some of the previously approved requirements for the Pilot Program.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2022–07631 Filed 4–8–22; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

[OMB No. 3064–0207]

Agency Information Collection Activities: Proposed Collection Renewal; Comment Request

AGENCY: Federal Deposit Insurance Corporation (FDIC).

ACTION: Notice and request for comment.

SUMMARY: The FDIC, as part of its obligations under the Paperwork Reduction Act of 1995 (PRA), invites the general public and other Federal agencies to take this opportunity to comment on the renewal of the existing information collection described below (OMB Control No. 3064–0207).

DATES: Comments must be submitted on or before June 10, 2022.

ADDRESSES: Interested parties are invited to submit written comments to the FDIC by any of the following methods:

- *Agency Website: https://www.fdic.gov/resources/regulations/federal-register-publications/.*
- *Email: comments@fdic.gov.* Include the name and number of the collection in the subject line of the message.
- *Mail:* Manny Cabeza (202–898–3767), Regulatory Counsel, MB–3128, Federal Deposit Insurance Corporation, 550 17th Street NW, Washington, DC 20429.
- *Hand Delivery:* Comments may be hand-delivered to the guard station at the rear of the 17th Street NW building (located on F Street NW), on business days between 7:00 a.m. and 5:00 p.m.

All comments should refer to the relevant OMB control number. A copy of the comments may also be submitted to the OMB desk officer for the FDIC: Office of Information and Regulatory Affairs, Office of Management and Budget, New Executive Office Building, Washington, DC 20503.

FOR FURTHER INFORMATION CONTACT: Manny Cabeza, Regulatory Counsel, 202–898–3767, *mcabeza@fdic.gov,* MB–3128, Federal Deposit Insurance Corporation, 550 17th Street NW, Washington, DC 20429.

SUPPLEMENTARY INFORMATION:

Proposal to renew the following currently approved collection of information:

1. *Title:* Loans in Areas Having Special Flood Hazards.

OMB Number: 3064–0207.

Form Number: None.

Affected Public: Private Sector.

Burden Estimate:

BURDEN CALCULATION

[OMB No. 3064–0207]

Description	Estimated annual number of respondents	Estimated annual number of responses per respondent	Estimated hours per response	Total hours
Recordkeeping:				
Private flood Insurance (Required to obtain benefits)	3,106	1	0.500	1,553.00
Standard flood hazard determination form (Mandatory)	3,106	313	0.042	40,831.48
Retention of notice of special flood hazards and availability of Federal disaster relief assistance (Mandatory)	3,106	36	0.250	27,954.00
Disclosure:				
Notice of requirement to escrow flood insurance payments and fees (Mandatory)	470	82	0.083	3,198.82
Change in status (Mandatory)	30	2	40	2,400.00
Notice of option to escrow flood insurance payments and fees (Mandatory)	30	22	0.083	54.78
Notice to borrower to purchase flood insurance (Mandatory)	3,106	10	0.083	2,577.98
Notification to terminate flood insurance purchased on behalf of a borrower (Mandatory)	3,106	1	0.250	776.50
Notice of special flood hazards and availability of Federal disaster relief assistance (Mandatory)	3,106	36	0.250	27,954.00
Notice to Administrator of FEMA of servicer's identity (Mandatory)	3,106	18	0.083	4,640.36
Notice to Administrator of FEMA of a change in loan servicer (Mandatory)	3,106	22	0.083	5,671.56

Total Estimated Burden Hours: $117,612.48.

General Description of Collection: Each supervised lending institution is required to provide a notice of special flood hazards to a borrower acquiring a loan secured by a building on real property located in an area identified by FEMA as subject to special flood hazards, and various other notices to borrowers, servicers and FEMA. The Riegle Community Development Act

requires that each institution also provide a copy of the notice to the servicer of the loan (if different from the originating lender). Section 100239 of the Biggert-Waters Flood Insurance Reform Act of 2012 requires each federal banking agency (including the FDIC), and the Farm Credit Administration, to adopt implementing regulations to direct regulated lending institutions to accept ''private flood insurance,'' as defined by the Biggert-Waters Act. A lending institution would be required to implement policies and procedures to comply with the Biggert-Waters Act provision and verify in writing that a private insurance policy satisfies the criteria included in the definition or document findings that separate required criteria have been met when accepting a private flood insurance policy in satisfaction of the mandatory flood insurance purchase requirement of the Flood Disaster Protection Act. The institution must also maintain records to permit examination staff to ascertain how the institution has met the requirements of the regulation.

The FDIC has reviewed its previous submission related to the PRA and has updated its methodology to align with the Office of the Comptroller of the Currency's corresponding information collection (1557–0326). The decrease in the estimated annual burden of 409,935 hours is the result of this change in methodology.

Request for Comment

Comments are invited on: (a) Whether the collection of information is necessary for the proper performance of the FDIC's functions, including whether the information has practical utility; (b) the accuracy of the estimates of the burden of the information collection, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. All comments will become a matter of public record.

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on April 5, 2022.

James P. Sheesley,

Assistant Executive Secretary.

[FR Doc. 2022–07639 Filed 4–8–22; 8:45 am]

BILLING CODE 6714–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–D–0552]

Safety and Performance Based Pathway Device-Specific Guidances; Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of two final device-specific guidance documents for the Safety and Performance Based Pathway—specifically, ''Surgical Sutures—Performance Criteria for Safety and Performance Based Pathway'' and ''Orthopedic Fracture Fixation Plates—Performance Criteria for Safety and Performance Based Pathway.'' The device-specific guidances identified in this notice were developed in accordance with the finalized guidance entitled ''Safety and Performance Based Pathway.''

DATES: The announcement of the guidances is published in the **Federal Register** on April 11, 2022.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time 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–2022–D–0552 for ''Surgical Sutures—Performance Criteria for Safety and Performance Based Pathway'' or ''Orthopedic Fracture Fixation Plates—Performance Criteria for Safety and Performance Based Pathway.'' Received comments 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.'' 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 *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 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: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*