

Board's monetary policy and supervisory mandates by providing greater insight into funding market conditions in periods where conditions change rapidly, potentially affecting policy measures taken by the Federal Reserve. The proposed revisions to FR 2420 would be effective with the January 1, 2022, as of date.

Reporting Form Revisions

The Board proposes to add a data item to specify the day-count convention used for all interest rates reported on FR 2420. The Federal Reserve has identified limited instances of reporting institutions using multiple day-count conventions in calculating reported interest rates, specifically found in the reporting of Part C interest rates. The proposed revision would improve the accuracy of reported data, benefiting the Federal Reserve's monitoring of funding market conditions and strengthening the production of the EFFR and OBFR. The proposed data item would provide the following day-count conventions as options: Actual/360, actual/365, 30/360, 30/365, actual/actual, and other.

Instruction Revisions

Additional Reference Rate Options for Floating-Rate Time Deposits and CDs (Part C)

The Board proposes to include additional reference rates to which floating-rate time deposits and CDs are tied. The additional rates include the Secured Overnight Financing Rate (SOFR), other SOFR-based rates, and OBFR, all of which are published daily by the Federal Reserve Bank of New York (FRBNY). Other SOFR-based rates include the SOFR Index and the SOFR Averages over 30, 90, and 180 days. This revision would improve the ability of the Federal Reserve to monitor the progress of the transition from LIBOR to SOFR with respect to floating-rate money market instruments.²

Earlier Deadline for Submission of Time Deposit and CD Data (General Instructions)

The Board proposes to change the deadline for submission of time deposit and CD data in Part C to 2 p.m. ET one business day (T+1) after the report date, rather than two business days (T+2) after the report date. This proposed change would provide more timely data and improve the Federal Reserve's monitoring of funding market

conditions. The change would be particularly beneficial on occasions when market conditions change quickly, such as when a deterioration in time deposit and CD markets may produce spillovers to other markets.

Earlier Deadline for Submission of Federal Funds Purchased, Eurodollar, and Selected Deposits Data (General Instructions)

The Board proposes to change the deadline for submission of Federal Funds Purchased, Eurodollars, and Selected Deposits data in Parts A, B, and D to 7 p.m. ET the same day (T+0) as the transaction date, rather than 7 a.m. ET one business day (T+1) after the transaction date. The proposed earlier reporting deadline would allow for more opportunity for data review and validation, reducing operational risk associated with the publication of the EFFR and OBFR.

Clarifications To Prevent Errors (Parts C and D)

The Board proposes other minor additions to the FR 2420 instructions to prevent confusion and errors on the part of reporting institutions. Guidance would be added for certain reciprocal deposits, including insured deposit cash sweeps and Certificate of Deposit Account Registry Service deposits (Part C). Additional guidance would be included on the correct reporting of brokered deposits (Part C) and certain securities lending transactions (Part D).

Legal authorization and confidentiality: The FR 2420 is authorized by section 11 of the Federal Reserve Act (FRA) and section 7 of the International Banking Act of 1978 (IBA). Section 11 of the FRA authorizes the Board to require reports from depository institutions as it may deem necessary and authorizes the Board to prescribe reports of liabilities and assets from insured depository institutions to enable the Board to discharge its responsibility to monitor and control monetary and credit aggregates (12 U.S.C. 248(a)). Section 7 of the IBA provides that federal branches and agencies of foreign banks are subject to section 11 of the FRA as if they were state member banks (12 U.S.C. 3105(c)). The obligation to respond to the FR 2420 is mandatory.

The FRBNY uses aggregate data from the FR 2420 to publish the EFFR, OBFR, and associated statistics daily. The information provided by individual respondents to the FR 2420 is nonpublic commercial or financial information, which is both customarily and actually treated as private by the respondents. Responses to the FR 2420 are therefore accorded confidential treatment

pursuant to exemption 4 of the Freedom of Information Act (5 U.S.C. 552(b)(4)).

Consultation outside the agency: A group of large FR 2420 respondents (less than 10) were consulted in November 2020 regarding the feasibility of reporting timestamps for FR 2420 transactions, shifting reporting deadlines, and the day-count conventions used when reporting interest rates on FR 2420 transactions. Outreach results suggest that timestamps for transactions are not recorded in a consistent fashion across respondents, and thus the current proposals do not call for the reporting of timestamps. Outreach also suggests that most respondents currently report Parts A, B, and D of the FR 2420 report on a T+0 basis, and no respondents consulted suggested that a T+0 reporting deadline for Parts A, B, and D was not feasible. Most respondents consulted noted that they should be able to report Part C transactions on a T+1 basis. Feedback also showed that most transactions are reported using the actual/360 day-count convention for interest rates, but other day-count conventions are used for some reported transactions.

Board of Governors of the Federal Reserve System, April 29, 2021.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2021-09424 Filed 5-4-21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Grants to States for Access and Visitation, OMB #0970-0204

AGENCY: Division of Program Innovation, Office of Child Support Enforcement, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Division of Program Innovation (DPI), Office of Child Support Enforcement (OCSE), Administration for Children and Families (ACF) is requesting a 3-year extension of the Access and Visitation Survey: Annual Report (OMB #0970-0204, expiration 10/31/2021). There are no changes requested to the form.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the

² The Alternative Reference Rates Committee is a group of private-market participants convened by the Board and the FRBNY to help ensure a successful transition from U.S. dollar LIBOR to a more robust reference rate, its recommended alternative, the SOFR.

Federal Register. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

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 30-day Review—Open for Public Comments” or by using the search function.

SUPPLEMENTARY INFORMATION:
Description: The grantee and/or sub-grantee submits the spreadsheet and

survey yearly. Information is used by OCSE as the primary means for adhering to the statutory (Sec. 469B. [42 U.S.C. 669b]) and regulatory (45 CFR part 303) requirements for recipients of “*Grants to States for Access and Visitation.*”

Respondents: State Child Access and Visitation Programs and state and/or local service providers.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Online Portal Survey by States and Jurisdictions	54	1	16	864
Survey of local service grantees	296	1	16	4,736

Estimated Total Annual Burden Hours: 5,600.

Authority: Sec.469B [42 U.S.C.669b]; 45 CFR part 303.

Mary B. Jones,
ACF/OPRE Certifying Officer.

[FR Doc. 2021-09452 Filed 5-4-21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Biomedical Imaging and Bioengineering; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the National Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel; P41 NCBIB Review F-SEP 2.

Date: June 21–23, 2021.

Time: 9:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy

Boulevard, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Dennis Hlasta, Ph.D., Scientific Review Officer, National Institute of Biomedical Imaging and Bioengineering, National Institutes of Health, 6707 Democracy Blvd., Bethesda, MD 20892, (301) 451-4794, dennis.hlasta@mail.nih.gov.

Name of Committee: National Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel; P41 NCBIB Review F-SEP 1.

Date: June 30, 2021.

Time: 9:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Plaza, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Dennis Hlasta, Ph.D., Scientific Review Officer, National Institute of Biomedical Imaging and Bioengineering, National Institutes of Health, 6707 Democracy Blvd., Bethesda, MD 20892, (301) 451-4794, dennis.hlasta@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.866, National Institute of Biomedical Imaging and Bioengineering, National Institutes of Health, HHS)

Dated: April 30, 2021.

Ronald J. Livingston, Jr.,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021-09491 Filed 5-4-21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Generic Clearance To Collect Stakeholder Feedback on the Research Domain Criteria (RDoC) Initiative, (NIMH)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Andrew Hooper, Ph.D., NIMH Project Clearance Liaison, Science Policy and Evaluation Branch, Office of Science Policy, Planning and Communications, NIMH, Neuroscience Center, 6001 Executive Boulevard, MSC 9667, Bethesda, Maryland 20892, call (301) 480-8433, or email your request, including your mailing address, to nimhprapubliccomments@mail.nih.gov. Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: Written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency’s estimate of the burden of the proposed collection of information,