3722, Legal Division. Board of Governors of the Federal Reserve System, 20th and C Streets NW, Washington, DC 20551. For the hearing impaired only, Telecommunications Device for the Deaf (TDD) users may contact (202) 263–4869.

SUPPLEMENTARY INFORMATION: The Board's GSIB surcharge rule establishes a methodology to identify global systemically important bank holding companies in the United States (GSIBs) based on indicators that are correlated with systemic importance.[1] Under the GSIB surcharge rule, a firm must calculate its GSIB score using a specific formula (Method 1). Method 1 uses five equally weighted categories that are correlated with systemic importance—size, interconnectedness, cross-jurisdictional activity, substitutability, and complexity—and subdivided into twelve systemic indicators. For each indicator, a firm divides its own measure of each systemic indicator by an aggregate global indicator amount. A firm's Method 1 score is the sum of its weighted systemic indicator scores expressed in basis points. The GSIB surcharge for a firm is the higher of the GSIB surcharge determined under Method 1 and a second method, Method 2, which weights size, interconnectedness, cross-jurisdictional activity, complexity, and a measure of a firm's reliance on wholesale funding (instead of substitutability).[2]

The aggregate global indicator amounts used in the score calculation under Method 1 are based on data collected by the Basel Committee on Banking Supervision (BCBS). The BCBS amounts are determined based on the sum of the systemic indicator scores of the 75 largest U.S. and foreign banking organizations as measured by the BCBS, and any other banking organization that the BCBS includes in its sample total for that year. The BCBS publicly releases these values, denominated in euros, each year. Pursuant to the GSIB surcharge rule, the Board publishes the aggregate global indicator amounts each year as denominated in U.S. dollars using the euro-dollar exchange rate provided by the BCBS.[3] Specifically, the Board multiplied each of the euro-denominated indicator amounts made publicly available by the BCBS by 1.1450, which was the daily euro to U.S. dollar spot rate on December 31, 2018, provided by the BCBS (as published by the European Central Bank, available at *http://www.ecb.europa.eu/stats/eurofxref/index.en.html*).

The aggregate global indicator amounts for purposes of the 2019 Method 1 score calculation under § 217.404(b)(1)(i)(B) of the GSIB surcharge rule are:

AGGREGATE GLOBAL INDICATOR AMOUNTS IN U.S. DOLLARS (USD) FOR 2019

Category	Systemic indicator	Aggregate global indicator amount (in USD)
Size	Total exposures	86,929,981,510,715
Interconnectedness	Intra-financial system assets	8,378,699,821,090
	Intra-financial system liabilities	9,423,444,832,391
	Securities outstanding	14,980,796,701,622
Substitutability	Payments activity	2,451,526,935,926,810
	Assets under custody	162,964,740,953,671
	Underwritten transactions in debt and equity markets	6,508,969,472,114
Complexity	Notional amount of over-the-counter (OTC) derivatives	606,648,652,426,571
	Trading and available-for-sale (AFS) securities	3,572,783,522,209
	Level 3 assets	530,724,384,529
Cross-jurisdictional activity	Cross-jurisdictional claims	21,901,114,980,308
	Cross-jurisdictional liabilities	18,341,219,019,191

Authority: 12 U.S.C. 248(a), 321–338a, 481–486, 1462a, 1467a, 1818, 1828, 1831n, 1831o, 1831p–l, 1831w, 1835, 1844(b), 1851, 3904, 3906–3909, 4808, 5365, 5368, 5371.

Board of Governors of the Federal Reserve System, December 16, 2019.

Ann Misback,

Secretary of the Board.

[FR Doc. 2019–27414 Filed 12–18–19; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10302]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

[1] *See* 12 CFR 217.402, 217.404.

[2] Method 2 uses similar inputs to those used in Method 1, but replaces the substitutability category with a measure of a firm's use of short-term wholesale funding. In addition, Method 2 is calibrated differently from Method 1.

[3] 12 CFR 217.404(b)(1)(i)(B); 80 FR 49082, 49086–87 (August 14, 2015). In addition, the Board maintains the GSIB Framework Denominators on its website, available at *https://www.federalreserve.gov/bankinforeg/basel/denominators.htm.*

DATES: Comments must be received by February 18, 2020.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number ____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10302 Collection Requirements for Compendia for Determination of Medically-Accepted Indications for Off-label Uses of Drugs and Biologicals in an Anti-Cancer Chemotherapeutic Regimen

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Collection Requirements for Compendia for Determination of Medically-accepted Indications for Off-label Uses of Drugs and Biologicals in an Anti-cancer Chemotherapeutic Regimen; *Use:* Section 182(b) of the Medicare Improvement of Patients and Providers Act (MIPPA) amended section 1861(t)(2)(B) of the Social Security Act (42 U.S.C. 1395x(t)(2)(B)) by adding at the end the following new sentence: 'On and after January 1, 2010, no compendia may be included on the list of compendia under this subparagraph unless the compendia has a publicly transparent process for evaluating therapies and for identifying potential conflicts of interest.' We believe that the implementation of this statutory provision that compendia have a ''publicly transparent process for evaluating therapies and for identifying potential conflicts of interests'' is best accomplished by amending 42 CFR 414.930 to include the MIPPA requirements and by defining the key components of publicly transparent processes for evaluating therapies and for identifying potential conflicts of interests.

All currently listed compendia will be required to comply with these provisions, as of January 1, 2010, to remain on the list of recognized compendia. In addition, any compendium that is the subject of a future request for inclusion on the list of recognized compendia will be required to comply with these provisions. No compendium can be on the list if it does not fully meet the standard described in section 1861(t)(2)(B) of the Act, as revised by section 182(b) of the MIPPA. *Form Number:* CMS–10302 (OMB control number: 0938–1078); *Frequency:* Annually; *Affected Public:* Business and other for-profits and Not-for-profit institutions; *Number of Respondents:* 845; *Total Annual Responses:* 900; *Total Annual Hours:* 5,135. (For policy questions regarding this collection contact Sarah Fulton at 410–786–2749.)

Dated: December 16, 2019.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2019–27385 Filed 12–18–19; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; OCSE–75 Tribal Child Support Enforcement Program Annual Data Report (OMB #0970–0320)

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

ACTION: Request for public comment.

SUMMARY: The Office of Child Support Enforcement (OCSE), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is requesting a three-year extension of the form OCSE–75—Tribal Child Support Enforcement Annual Data Report (OMB # 0970–0320, expiration 03/31/2020). There are no changes requested to the form.

DATES: *Comments due within 30 days of publication.* OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **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 directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV.* Attn: Desk Officer for the Administration for Children and Families.

Copies of the proposed collection may be obtained by emailing *infocollection@acf.hhs.gov.* Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: ACF Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION: