

other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at <https://www.federalreserve.gov/foia/request.htm>. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)).

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551-0001, not later than January 24, 2022.

A. *Federal Reserve Bank of Dallas* (Karen Smith, Director, Applications) 2200 North Pearl Street, Dallas, Texas 75201-2272:

1. *Texas State Bankshares, Inc., Harlingen, Texas*; to merge with Access Bancorp, Inc., and therefore indirectly acquire AccessBank Texas, both of Denton, Texas.

Board of Governors of the Federal Reserve System, December 20, 2021.

**Maragaret M. Shanks,**  
*Deputy Secretary of the Board.*

[FR Doc. 2021-27884 Filed 12-22-21; 8:45 am]

**BILLING CODE 6210-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Proposed Information Collection Activity; ACF Uniform Project Description**

**AGENCY:** Office of Administration, Office of Grants Policy, Administration for Children and Families, HHS.

**ACTION:** Request for public comment.

**SUMMARY:** The Administration for Children and Families (ACF) is requesting a 3-year extension of the ACF Uniform Project Description (UPD) (OMB #0970-0139, expiration 2/28/2022). There are no changes requested to the form.

**DATES:** *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork

Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

**ADDRESSES:** Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). Identify all requests by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* The proposed information collection would renew the ACF UPD. The UPD provides a uniform format for applicants to submit project information in response to ACF discretionary Notices of Funding Opportunities. The UPD requires applicants to describe how program objectives will be achieved and provide a rationale for the project's budgeted costs. All ACF discretionary grant programs are required to use the UPD.

ACF uses this information, along with other OMB-approved information collections (Standard Forms), to evaluate and rank applications. Use of the UPD protects the integrity of the ACF award selection process.

*Respondents:* Applicants responding to ACF Discretionary Notices of Funding Opportunities.

**ANNUAL BURDEN ESTIMATES**

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
ACF Uniform Project Description .....	3,218	1	60	193,080	64,360

*Estimated Total Annual Burden Hours:* 64,360.

*Comments:* The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

*Authority:* 45 CFR 75.203-75.204 and 45 CFR part 75, Appendix I.

**Mary B. Jones,**  
*ACF/OPRE Certifying Officer.*

[FR Doc. 2021-27837 Filed 12-22-21; 8:45 am]

**BILLING CODE 4184-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2021-D-0980]

**Assessing the Credibility of Computational Modeling and Simulation in Medical Device Submissions; Draft 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 the draft guidance entitled "Assessing the Credibility of Computational Modeling and Simulation in Medical Device Submissions." Computational modeling and simulation (CM&S) can be used in a variety of ways in medical device applications, including to perform "in silico" device testing or as part of software embedded in a device. This guidance provides a risk-based framework that can be used in the credibility assessment of computational modeling and simulation (CM&S) used in medical device regulatory submissions. The draft guidance is intended to improve the consistency and transparency of the review of computational modeling evidence. This draft guidance is not final nor is it in effect at this time.

**DATES:** Submit either electronic or written comments on the draft guidance