

change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Benjamin W. McDonough, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551-0001, not later than September 15, 2026.

A. Federal Reserve Bank of Chicago (Christopher Koopmans, Senior Vice President) 230 South LaSalle Street, Chicago, Illinois 60690-1414. Comments can also be sent electronically to

Comments.applications@chi.frb.org:

1. *William D. Mullins Revocable Trust, William D. Mullins as trustee, and the Janice M. Mullins Revocable Trust, Janice M. Mullins, as trustee, all of Algona, Iowa;* to join the Bierstedt Family Control Group, a group acting in concert, to acquire voting shares of Burt Bancshares, Inc., Burt, Iowa, and thereby indirectly acquire voting shares of Security State Bank, Algona, Iowa.

B. Federal Reserve Bank of Kansas City (Jeffrey Imgarten, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198-0001. Comments can also be sent electronically to *KCAApplicationComments@kc.frb.org:*

1. *Kimberly J. Fairbank Family Trust, Trenton Fairbank, as trustee, Trenton L. and Sara M. Fairbank Revocable Trust, Trenton and Sara Fairbank, as co-trustees, all of Cimarron, Kansas; Joe D. Butcher Family Trust-Grant Butcher Share, Cimarron, Kansas, Grant Butcher, Springdale, Arkansas, and First National Bank of Hutchinson, Hutchinson, Kansas, as co-trustees; and the Joe D. Butcher Family Trust-Robert Butcher Share, Cimarron, Kansas, Robert Butcher, DeSoto, Kansas, and First National Bank of Hutchinson, Hutchinson, Kansas, as co-trustees;* to join the Butcher Family Control Group, a group acting in concert, to retain voting shares of First National Agency, Inc., and thereby indirectly retain voting shares of First National Bank in

Cimarron, both of Cimarron, Kansas. Robert Butcher and Grant Butcher are members of the Butcher Family Control Group and were each previously permitted to acquire voting shares of First National Agency, Inc.

2. *Kimberly Jo Fairbank Family Trust, Trenton Fairbank, as trustee; Trenton L. and Sara M. Fairbank Revocable Trust, Trenton and Sara Fairbank, as co-trustees, all of Cimarron, Kansas; Grant D. Butcher GST Trust, Cimarron, Kansas, Grant Butcher, Springdale, Arkansas, and First National Bank of Hutchinson, Hutchinson, Kansas, as co-trustees; Robert L. Butcher GST Trust, Cimarron, Kansas, Robert Butcher, DeSoto, Kansas, and First National Bank of Hutchinson, Hutchinson, Kansas, as co-trustees, and Dea Ann Kreisman, Parker, Colorado;* to join the Butcher Family Control Group, a group acting in concert, to retain voting shares of Santa Fe Trail Banc Shares, Inc., and thereby indirectly retain voting shares of Centera Bank, both of Sublette, Kansas.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Associate Secretary of the Board.

[FR Doc. 2026-17753 Filed 8-28-26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for Office of Management and Budget Review; Head Start State Collaboration Office Grant Application

AGENCY: Office of Head Start, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Office of Head Start (OHS), Administration for Children and Families (ACF) is proposing a new information collection activity for the Head Start State Collaboration Office (HSCO) Grant Application. HSCO grants are authorized under the Head Start Act and support coordination between Head Start agencies and state systems serving young children and families. OHS has

developed new application instructions to align HSCO grant activities with states and federal priorities, including improving outcomes in child welfare, strengthening family formation, supporting effective service delivery, and enhancing early childhood education. This proposed information collection is necessary to support consistent implementation of program objectives and alignment across all states.

DATES: Comments due September 30, 2026.

ADDRESSES: The public may view and comment on this information collection request at: https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202608-0970-009. You can also obtain copies of the proposed collection of information by emailing infocollection@acf.hhs.gov. Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: OHS proposes collecting application materials from all states and territories receiving HSCO grants. These grants are non-competing and statutorily required. The new application instructions will guide states in aligning Collaboration Office activities with updated federal priorities and strengthening integration of Head Start within state early childhood systems.

The application will include brief narrative and structured responses describing state plans for coordination, systems integration, and program implementation. The request will ensure consistent documentation of proposed activities and enable ACF OHS to effectively administer and oversee the grants.

Existing HSCO grants will be extended through February 28, 2027, to allow time for PRA approval and transition to new grant awards. Following approval, OHS will issue new application instructions, and states will be provided with a defined application window to submit materials in advance of new 5-year grant awards.

Respondents: State governments (including the 50 states, the District of Columbia, and territories) receiving HSCO grants.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
HSCO Grant Application	52	1	10.4	540.8

Authority: Head Start Act (42 U.S.C. 9801 *et seq.*); PRA (44 U.S.C. 3501 *et seq.*)

Mary C. Jones,
ACF/OPRE Certifying Officer.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–N–6540]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Potential Tobacco Product Violations Reporting Form

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by September 30, 2026.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to <https://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. The OMB control number for this information collection is 0910–0716. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Amber Barrett, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, PRAStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Potential Tobacco Product Violations Reporting Form

OMB Control Number 0910–0716—Extension

This information collection supports the opportunity to accept consumer and other stakeholder feedback and notification of potential violations of the FD&C Act, as amended by the Tobacco Control Act. Tobacco products are generally governed by chapter IX of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (sections 900 through 920) (21 U.S.C. 387 through 21 U.S.C. 387u). The FD&C Act provides FDA authority to monitor compliance with Federal tobacco laws and regulations and take corrective action when violations occur.

As part of its enforcement strategy, FDA accepts information from the public regarding potential tobacco product violations of the FD&C Act. Potential tobacco product violations include (but are not limited to): (1) sales to underage purchasers (persons under 21); (2) flavored cigarette sales; (3) illegal marketing and advertising; (4) distribution of free samples of tobacco products except in limited circumstances; (5) placement of cigarette or smokeless tobacco product vending machines in prohibited areas (or providing access to self-service or direct access of tobacco products in prohibited areas); (6) the manufacture or sale of unauthorized tobacco products; and (7) sale of cigarettes in packages of less than 20.

FDA currently provides a form that may be used to collect this information from the public (Form FDA 3779, Potential Tobacco Product Violations Report). The Potential Tobacco Product Violations Report, Form FDA 3779, asks for the following information: (1) date potential violation occurred; (2) product type (*e.g.*, cigarette, smokeless, roll-your-own, cigar, e-cigarette, hookah, pipe tobacco); (3) tobacco brand; (4) potential violation type; (5) type of potentially violative promotional materials; (6) description of the potential violation (7) who potentially violated; (8) name and address of the potential violator (if known); (9) potential violator’s website or internet address URL (if available); (10) Optional filer contact information if additional information is needed or to receive notification the complaint was received; and (11) any additional files or information pertinent to the potential violation.

The public and interested stakeholders can report possible tobacco product violations of the FD&C Act by submitting information on Form FDA 3779 online, via email or postal mail, or by calling FDA’s Tobacco Call Center. Information on how to submit possible tobacco product violations using the options above can be found at <https://www.accessdata.fda.gov/scripts/ptvr/index.cfm>. Further details about reporting possible tobacco product violations of the FD&C Act can also be found at <https://www.fda.gov/tobacco-products/compliance-enforcement-training/report-potential-tobacco-product-violation>.

In the **Federal Register** of June 23, 2026 (91 FR 37410), FDA published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN ¹

Activity and form FDA 3779	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response in hours	Total hours
Reporting potential tobacco product violations of the FD&C Act.	3,000	2	6,000	0.25 (15 minutes) ..	1,500

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

The burden hour estimates for this collection of information were based on the type and rate of reporting submitted through the Potential Tobacco Violation Report Form and based on a review of

the information collection since our last request for OMB approval. FDA estimates that submitting the information (online, telephone, email,

or mail) will take 0.25 hours (*i.e.*, 15 minutes) per response. FDA estimates the number of annual respondents to this collection of information will be 3,000, who will each submit 2 reports. Each report is