system of records, FCC/WCB–1, Lifeline, which was published in the **Federal Register** at 89 FR 28777 (Apr. 19, 2024).

The records shared as part of this matching program reside in the ACP system of records, FCC/WCB–3, Affordable Connectivity Program, which was published in the **Federal Register** at 89 FR 28780 (Apr. 19, 2024).

Federal Communications Commission.

Marlene Dortch,

Secretary.

[FR Doc. 2026–10839 Filed 5–29–26; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

[FR ID: 348889]

Privacy Act of 1974; Matching Program

AGENCY: Federal Communications Commission.

ACTION: Notice of a new matching program.

SUMMARY: In accordance with the Privacy Act of 1974, as amended (''Privacy Act''), this document announces a new computer matching program the Federal Communications Commission (FCC or Commission or Agency) and the Universal Service Administrative Company (USAC) will conduct with the Kentucky Cabinet for Health and Family Services, Department for Community Based Services. The purpose of this matching program is to verify the eligibility of applicants to and subscribers of Lifeline, and the Affordable Connectivity Program (ACP), both of which are administered by USAC under the direction of the FCC. More information about these programs is provided in the **SUPPLEMENTARY INFORMATION** section below.

DATES: Written comments are due on or before July 1, 2026. This computer matching program will commence on July 1, 2026, and will conclude after 18 months.

ADDRESSES: Send comments to Shana Yates, FCC, 45 L Street NE, Washington, DC 20554, or to *Privacy@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: Shana Yates at (202) 418–0683 or *Privacy@fcc.gov.*

SUPPLEMENTARY INFORMATION: The Lifeline program provides support for discounted broadband and voice services to low-income consumers. Lifeline is administered by the Universal Service Administrative Company (USAC) under FCC direction. Consumers qualify for Lifeline through proof of income or participation in a qualifying program, such as Medicaid, the Supplemental Nutritional Assistance Program (SNAP), Federal Public Housing Assistance, Supplemental Security Income (SSI), Veterans and Survivors Pension Benefit, or various Tribal-specific federal assistance programs.

In the Consolidated Appropriations Act, 2021, Public Law 116–260, 134 Stat. 1182, 2129–36 (2020), Congress created the Emergency Broadband Benefit Program, and directed use of the National Verifier to determine eligibility based on various criteria, including the qualifications for Lifeline (Medicaid, SNAP, etc.). EBBP provided $3.2 billion in monthly consumer discounts for broadband service and one-time provider reimbursement for a connected device (laptop, desktop computer or tablet). In the Infrastructure Investment and Jobs Act, Public Law 117–58, 135 Stat. 429, 1238–44 (2021) (codified at 47 U.S.C. 1751–52), Congress modified and extended EBBP, provided an additional $14.2 billion, and renamed it the Affordable Connectivity Program (ACP). A household may qualify for the ACP benefit under various criteria, including an individual qualifying for the FCC's Lifeline program.

In a Report and Order adopted on March 31, 2016, (81 FR 33026, May 24, 2016) (*2016 Lifeline Modernization Order*), the Commission ordered USAC to create a National Lifeline Eligibility Verifier (''National Verifier''), including the National Lifeline Eligibility Database (LED), that would match data about Lifeline applicants and subscribers with other data sources to verify the eligibility of an applicant or subscriber. The Commission found that the National Verifier would reduce compliance costs for Lifeline service providers, improve service for Lifeline subscribers, and reduce waste, fraud, and abuse in the program.

The Consolidated Appropriations Act of 2021 directs the FCC to leverage the National Verifier to verify applicants' eligibility for ACP. The purpose of this matching program is to verify the eligibility of Lifeline and ACP applicants and subscribers by determining whether they receive SNAP and Medicaid benefits administered by the Kentucky Cabinet for Health and Family Services, Department for Community Based Services.

PARTICIPATING AGENCIES:

Kentucky Cabinet for Health and Family Services, Department for Community Based Services (source agency); Federal Communications Commission (recipient agency) and Universal Service Administrative Company.

AUTHORITY FOR CONDUCTING THE MATCHING PROGRAM:

The authority to conduct the matching program for the FCC's ACP is 47 U.S.C. 1752(a) through (b). The authority to conduct the matching program for the FCC's Lifeline program is 47 U.S.C. 254(a) through (c), and (j).

PURPOSE(S):

The purpose of this new matching agreement is to verify the eligibility of applicants and subscribers to Lifeline, as well as to ACP and other Federal programs that use qualification for Lifeline as an eligibility criterion. This new agreement will permit eligibility verification for the Lifeline program and ACP by checking an applicant's/subscriber's participation in SNAP and Medicaid in Kentucky Cabinet for Health and Family Services, Department for Community Based Services. Under FCC rules, consumers receiving these benefits qualify for Lifeline discounts and also for ACP benefits.

CATEGORIES OF INDIVIDUALS:

The categories of individuals whose information is involved in the matching program include, but are not limited to, those individuals who have applied for Lifeline and/or ACP benefits; are currently receiving Lifeline and/or ACP benefits; are individuals who enable another individual in their household to qualify for Lifeline and/or ACP benefits; are minors whose status qualifies a parent or guardian for Lifeline and/or ACP benefits; or are individuals who have received Lifeline and/or ACP benefits.

CATEGORIES OF RECORDS:

The categories of records involved in the matching program include the last four digits of the applicant's Social Security Number, date of birth, first and last name. The National Verifier will transfer these data elements to the Kentucky Cabinet for Health and Family Services, Department for Community Based Services which will respond either ''yes'' or ''no'' that the individual is enrolled in a qualifying assistance program: SNAP and Medicaid administered by the Kentucky Cabinet for Health and Family Services, Department for Community Based Services.

SYSTEM(S) OF RECORDS:

The records shared as part of this matching program reside in the Lifeline system of records, FCC/WCB–1, Lifeline, which was published in the

Federal Register at 89 FR 28777 (Apr. 19, 2024).

The records shared as part of this matching program reside in the ACP system of records, FCC/WCB–3, Affordable Connectivity Program, which was published in the **Federal Register** at 89 FR 28780 (Apr. 19, 2024).

Federal Communications Commission.

Marlene Dortch,

Secretary.

[FR Doc. 2026–10840 Filed 5–29–26; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The public portions of the applications listed below, as well as 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 received are subject to public disclosure. In general, comments received will be made available without 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 July 1, 2026.

A. Federal Reserve Bank of Cleveland (Jenni M. Frazer, Vice President) 1455 East Sixth Street, Cleveland, Ohio 44101–2566. Comments can also be sent electronically to *Comments.applications@clev.frb.org:*

1. *Peoples Bancorp Inc., Marietta, Ohio;* to acquire Citizens National Corporation, and thereby indirectly acquire Citizens Bank of Kentucky, Inc., both of Paintsville, Kentucky.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Associate Secretary of the Board.

[FR Doc. 2026–10882 Filed 5–29–26; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–D–2839]

Oncology Pharmaceuticals: Streamlined Nonclinical Safety Studies for Biologics and Conjugated Products; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the availability of a draft guidance for industry entitled ''Oncology Pharmaceuticals: Streamlined Nonclinical Safety Studies for Biologics and Conjugated Products.'' When finalized, this guidance will assist sponsors in implementing streamlined approaches for general toxicology, for nonclinical safety assessments of certain oncology pharmaceuticals. The guidance is intended to facilitate drug development for biological products and conjugated products for the treatment of cancer while avoiding unnecessary animal use. The recommendations in this draft guidance are informed by data analysis of general toxicology studies and practices developed during the COVID–19 pandemic to reduce use of non-human primates.

DATES: Submit either electronic or written comments on the draft guidance by July 31, 2026 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance 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–2026–D–2839 for ''Oncology Pharmaceuticals: Streamlined Nonclinical Safety Studies for Biologics and Conjugated Products.'' 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