Network (CDC TRAIN) since 2010. There are now more than two million CDC TRAIN learner accounts. PHF also manages the Council on Linkages Between Academic and Public Health Practice, which is the steward of the Core Competencies for Public Health Professionals, the Academic Health Department (AHD) Learning Community, and the Retention and Recruitment Learning Community. Finally, PHF is a field-recognized lead in unique public health tools and training for performance management and quality improvement provided to state, tribal, local, and territorial health departments.

Summary of the Award

Recipient: Public Health Foundation.

Purpose of the award: The purpose of this award is to ensure that state, tribal, local, and territorial (STLT) public health and healthcare professionals have the knowledge and skills to best serve their communities. This will be achieved by the following: (1) enhancing and expanding the TRAIN Learning Network, an existing learning management system (LMS) focused on upskilling the public health and healthcare workforce and (2) strengthening public health workforce services through support for advancing performance management and quality improvement, public health workforce competencies, academic health departments, and other cross-sector collaborations.

Amount of award: The approximate year 1 funding amount will be $1.5 million in Federal Fiscal Year 2026 funds, subject to the availability of funds. Funding amounts for years 2–5 will be set at continuation.

Authority: This program is authorized under the Public Health Service Act: Section 1703 (a)(2) and (3) (42 U.S.C. 300u–2(a)(2) and (3)); Section 1704 (42 U.S.C. 300u–3); and Section 317(k)(2) (42 U.S.C. 247(b)(k)(2)).

Period of performance: September 1, 2026, through August 31, 2031.

Jamie Legier,

Director, Office of Grants Services, Centers for Disease Control and Prevention.

[FR Doc. 2026–15315 Filed 7–28–26; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

Privacy Act of 1974; Matching Program

AGENCY: Centers for Medicare & Medicaid Services (CMS), Department of Health and Human Services (HHS).

ACTION: Notice of a new matching program.

SUMMARY: In accordance with the Privacy Act of 1974, as amended, the Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS) is providing notice of the re-establishment of a matching program between CMS and the Office of Personnel Management, ''Verification of Eligibility for Minimum Essential Coverage Under the Patient Protection and Affordable Care Act through an Office of Personnel Management Health Benefit Plan''. The matching program provides CMS and State Administering Entities with the Office of Personnel Management (OPM) data to use in determining individuals' eligibility to enroll in a qualified health plan through an exchange established under the ACA; for insurance affordability programs, certifications of exemption, and to make eligibility redeterminations and renewals, including appeal determinations.

DATES: Comments on the matching program must be received by August 28, 2026. The matching program will be effective, and matching activity may commence, 30 days after this notice is published in the **Federal Register**. The matching program conducted for an initial term of 18 months, and within 3 months of expiration may be renewed for up to one additional year if the parties make no change to the matching program and certify that the program has been conducted in compliance with the matching agreement.

ADDRESSES: Interested parties may submit comments on this notice to the CMS Privacy Act Officer by mail at: Division of Security, Privacy Policy & Governance, Information Security & Privacy Group, Office of Information Technology, Centers for Medicare & Medicaid Services, Location: N1–14–56, 7500 Security Blvd., Baltimore, MD 21244–1850 or by email at *Barbara.Demopulos@cms.hhs.gov*.

FOR FURTHER INFORMATION CONTACT: If you have questions about the matching program, you may contact Terrence Kane, Director, Division of Automated Verifications and SEP Policy, Marketplace Eligibility and Enrollment Group, Center for Consumer Information and Insurance Oversight, Centers for Medicare & Medicaid Services, at (301) 492–4449, by email at *Terrence.kane@cms.hhs.gov,* or by mail at 7501 Wisconsin Avenue, Bethesda, MD 20814.

SUPPLEMENTARY INFORMATION: The Privacy Act of 1974, as amended (5 U.S.C. 552a) provides certain protections for individuals applying for and receiving federal benefits. The law governs the use of computer matching by federal agencies when records in a system of records (meaning, federal agency records about individuals retrieved by name or other personal identifier) are matched with records of other federal or non-federal agencies. The Privacy Act requires agencies involved in a matching program to:

1. Enter into a written agreement, which must be prepared in accordance with the Privacy Act, approved by the Data Integrity Board of each source and recipient Federal agency, provided to Congress and the Office of Management and Budget (OMB), and made available to the public, as required by 5 U.S.C. 552a(o), (u)(3)(A), and (u)(4).
2. Notify the individuals whose information will be used in the matching program that the information they provide is subject to verification through matching, as required by 5 U.S.C. 552a(o)(1)(D).
3. Verify match findings before suspending, terminating, reducing, or making a final denial of an individual's benefits or payments or taking other adverse action against the individual, as required by 5 U.S.C. 552a(p).
4. Report the matching program to Congress and the OMB, in advance and annually, as required by 5 U.S.C. 552a(o) (2)(A)(i), (r), and (u)(3)(D).
5. Publish advance notice of the matching program in the **Federal Register** as required by 5 U.S.C. 552a(e)(12).

This matching program meets these requirements.

Barbara Demopulos,

Privacy Act Officer, Division of Security, Privacy Policy and Governance, Office of Information Technology, Centers for Medicare & Medicaid Services.

Participating Agencies: Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS), and the Office of Personnel Management (OPM).

Authority for Conducting the Matching Program: The principal authority for the matching program is 42 U.S.C. 18001, *et seq.*

Purpose(s): The matching program will enable CMS to make initial

Eligibility Determinations for eligibility to enroll in a Qualified Health Plan (QHP) through an Exchange established under ACA; eligibility for Insurance Affordability Programs and for certifications of Exemption; and eligibility Redeterminations and Renewal decisions, including appeal determinations, for enrollment in a QHP through an Exchange and Insurance Affordability Programs and for certifications of Exemption.

Categories of Individuals: The categories of individuals whose information is involved in the matching program are: (1) active Federal Employees, identified in data CMS receives from OPM, and (2) consumers who apply for or are enrolled in private insurance coverage under a qualified health plan through a federally-facilitated or state-based health insurance exchange (and other relevant individuals, such as applicants' and enrollees' household members), whose records are matched against the data CMS receives from OPM.

Categories of Records: OPM will send a monthly, full refreshed Status File that contains a list of and data for active Federal employees. The Status File will include the following specified data elements: Record type; Record number; Unique person ID; Social Security Number; Last name; Middle name; First name; Last name suffix; Sex; Date of birth; and Health Plan Code.

OPM will send to CMS, on an annual basis, a Premium Spread Index File that identifies the lowest premium available to a Federal employee in each of the 32 premium localities.

System(s) of Records:

A. System of Records Maintained by CMS: The CMS SOR that supports this matching program is the ''CMS Health Insurance Exchanges System (HIX)'', CMS System No. 09–70–0560, last published in full at 78 FR 63211 (October 23, 2013), as amended at 83 FR 6591 (February 14, 2018).

B. System of Records Maintained by the Office of Personnel Management: The OPM SOR for this matching program is titled ''General Personnel Records'' (OPM/GOVT–1), published at 77 FR, 73694 (December 11, 2012).

[FR Doc. 2026–15282 Filed 7–28–26; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2007–D–0369]

Product-Specific Guidances; Revised Draft Guidances for Industry; 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 additional revised draft product-specific guidances. The draft guidances provide product-specific recommendations on, among other things, the design of bioequivalence (BE) studies to support abbreviated new drug applications (ANDAs). In the **Federal Register** of June 11, 2010, FDA announced the availability of a guidance for industry entitled ''Bioequivalence Recommendations for Specific Products'' that explained the process that would be used to make product-specific guidances available to the public on FDA's website. The draft guidances identified in this notice were developed using the process described in that guidance.

DATES: Submit either electronic or written comments on the draft guidance by September 28, 2026 to ensure that the Agency considers your comment on these draft guidances before it begins work on the final version of these guidances.

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–2007–D–0369 for ''Product-Specific Guidances; Revised Draft Guidances for Industry.'' 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 with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://*