

indirectly acquire American Bank of Commerce, both of Provo, Utah.

Board of Governors of the Federal Reserve System, September 28, 2018.

Yao-Chin Chao,

Assistant Secretary of the Board.

[FR Doc. 2018–21544 Filed 10–2–18; 8:45 am]

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FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than October 15, 2018.

A. Federal Reserve Bank of Kansas City (Dennis Denney, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198–0001:

1. *Breck C. Collingsworth, Lincoln, Nebraska, individually and as part of a group acting in concert with Susan Chrastil, Crete, Nebraska;* to acquire voting shares of TCM Company, Crete, Nebraska, and thereby indirectly acquire shares of City Bank & Trust Co., Lincoln, Nebraska.

Board of Governors of the Federal Reserve System, September 28, 2018.

Yao-Chin Chao,

Assistant Secretary of the Board.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2018–0091]

Proposed Revised Vaccine Information Materials for Meningococcal ACWY and DTaP (Diphtheria, Tetanus, Pertussis) Vaccines

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: Under the National Childhood Vaccine Injury Act (NCVIA), the Centers for Disease Control and Prevention (CDC) within the Department of Health and Human Services (HHS) develops vaccine information materials that all health care providers are required to give to patients/parents prior to administration of specific vaccines. HHS/CDC seeks written comment on the proposed updated vaccine information statements for meningococcal ACWY and DTaP (diphtheria, tetanus, acellular pertussis) vaccines.

DATES: Written comments must be received on or before December 3, 2018.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2018–0091, by any of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments.
- *Mail:* Written comments should be addressed to Suzanne Johnson-DeLeon (VISComments@cdc.gov), National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, Mailstop A–19, 1600 Clifton Road NE, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and docket number. All relevant comments received will be posted without change to <http://www.regulations.gov>, including any personal information provided. For access to the docket to read background documents or comments received, go to <http://www.regulations.gov>.

FOR FURTHER INFORMATION CONTACT: Skip Wolfe, National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, Mailstop A–19, 1600 Clifton Road NE, Atlanta, Georgia 30329; VISComments@cdc.gov.

SUPPLEMENTARY INFORMATION: The National Childhood Vaccine Injury Act of 1986 (Pub. L. 99–660), as amended by section 708 of Public Law 103–183,

added section 2126 to the Public Health Service Act. Section 2126, codified at 42 U.S.C. 300aa–26, requires the Secretary of Health and Human Services to develop and disseminate vaccine information materials for distribution by all health care providers in the United States to any patient (or to the patient's parent or legal representative in the case the patient is a child) receiving vaccines covered under the National Vaccine Injury Compensation Program (VICP).

Development and revision of the vaccine information materials, also known as Vaccine Information Statements (VIS), have been delegated by the Secretary to the Centers for Disease Control and Prevention (CDC). Section 2126 requires that the materials be developed, or revised, after notice to the public, with a 60-day comment period, and in consultation with the Advisory Commission on Childhood Vaccines, appropriate health care provider and parent organizations, and the Food and Drug Administration. The law also requires that the information contained in the materials be based on available data and information, be presented in understandable terms, and include:

- (1) A concise description of the benefits of the vaccine,
- (2) A concise description of the risks associated with the vaccine,
- (3) A statement of the availability of the National Vaccine Injury Compensation Program, and
- (4) Such other relevant information as may be determined by the Secretary.

The vaccines initially covered under the National Vaccine Injury Compensation Program were diphtheria, tetanus, pertussis, measles, mumps, rubella and poliomyelitis vaccines. Since April 15, 1992, any health care provider in the United States who intends to administer one of these covered vaccines is required to provide copies of the relevant vaccine information materials prior to administration of any of these vaccines. Since then, the following vaccines have been added to the National Vaccine Injury Compensation Program, requiring provision of vaccine information materials before vaccine administration for them as well: hepatitis B, *Haemophilus influenzae* type b (Hib), varicella (chickenpox), pneumococcal conjugate, rotavirus, hepatitis A, meningococcal, human papillomavirus (HPV), and seasonal influenza vaccines. Instructions for use of the vaccine information materials are found on the CDC website at: <https://www.cdc.gov/vaccines/hcp/vis/index.html>.

CDC is proposing updated versions of the meningococcal ACWY and DTaP

(diphtheria, tetanus, acellular pertussis) vaccine information statements.

Changes to the meningococcal ACWY VIS are minimal. Reference to the MPSV4 vaccine, no longer available in the United States, is removed. HIV infection is added as an indication for vaccination, and wording related to meningococcal ACWY vaccination during pregnancy is updated.

Proposed revisions to the DTaP VIS reflect new recommendations of the Advisory Committee on Immunization Practices (ACIP), including updated information about contraindications and precautions. Minor changes are proposed to simplify and streamline the sections about what to do if there is a reaction and finding additional information about the vaccine and the Vaccine Injury Compensation Program. The most recent previous final version of the DTaP VIS was published in 2007; proposed revisions to this document will help to bring it in line with the structure and general approach of more recently-published VISs for other vaccines.

The vaccine information materials referenced in this notice are being developed in consultation with the Advisory Commission on Childhood Vaccines, the Food and Drug Administration, and parent and health care provider groups.

We invite written comment on the proposed vaccine information materials entitled “Meningococcal ACWY Vaccine: What You Need to Know” and “DTaP (Diphtheria, Tetanus, Pertussis) Vaccine: What You Need to Know.” Copies of the proposed vaccine information materials are available at <http://www.regulations.gov> (see Docket Number CDC-2018-0091). Comments submitted will be considered in finalizing these materials. When the final materials are published in the **Federal Register**, the notice will include an effective date for their mandatory use.

Dated: September 27, 2018.

Sandra Cashman,

Executive Secretary, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

Privacy Act of 1974; Matching Program

AGENCY: Centers for Medicare & Medicaid Services, Department of Health and Human Services.

ACTION: Notice of a new matching program.

SUMMARY: In accordance with subsection (e)(12) of 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 a new matching program between CMS and the Department of Veterans Affairs (VA), Veterans Health Administration (VHA), “Verification of Eligibility for Minimum Essential Coverage Under the Patient Protection and Affordable Care Act Through a Veterans Health Administration Plan.”

DATES: The deadline for comments on this notice is November 2, 2018. The re-established matching program will commence not sooner than 30 days after publication of this notice, provided no comments are received that warrant a change to this notice. The matching program will be conducted for an initial term of 18 months (from approximately October 2018 to April 2020) and within 3 months of expiration may be renewed for 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 written comments to: CMS Privacy Act Officer, Division of Security, Privacy Policy & Governance, Information Security & Privacy Group, Office of Information Technology, CMS, 7500 Security Blvd., Baltimore, MD 21244-1870, Mailstop: N3-15-25, or by email to: walter.stone@cms.hhs.gov. Comments received will be available for review at this location, by appointment, during regular business hours, Monday through Friday from 9:00 a.m. to 3:00 p.m.

FOR FURTHER INFORMATION CONTACT: If you have questions about the matching program, you may contact Jack Lavelle, Senior Advisor, Marketplace Eligibility and Enrollment Group, Center for Consumer Information and Insurance Oversight, CMS, 7501 Wisconsin Ave., Bethesda, MD 20814, (410) 786-0639, or by email at Jack.Lavelle1@cms.hhs.gov.

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 Demopolos,

CMS Privacy Advisor, Division of Security, Privacy Policy and Governance, Information Security and Privacy Group, Office of Information Technology, Centers for Medicare & Medicaid Services.

Participating Agencies

The Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS) is the recipient agency, and the Department of Veterans Affairs (VA), Veterans Health Administration (VHA) is the source agency.

Authority for Conducting the Matching Program

The statutory authority for the matching program is 42 U.S.C. 18001.

Purpose(s)

The purpose of the matching program is to assist CMS in determining individuals' eligibility for financial assistance in paying for private health