SUMMARY: In this document, the Commission released a public notice announcing the meeting in accordance with the Federal Advisory Committee Act, this notice advises interested persons that the Federal Communications Commission's (FCC) Technological Advisory Council will hold its first meeting of the Technological Advisory Council for 2015.

DATES: Wednesday, April 1, 2015, from 1:00 p.m. to 4:00 p.m.

ADDRESSES: Federal Communications Commission, 445 12th Street SW., Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: Walter Johnston, Chief, Electromagnetic Compatibility Division, 202–418–0807; *Walter.Johnston@FCC.gov.*

SUPPLEMENTARY INFORMATION: This is a summary of the Commission's Public Notice, DA 15–184 released February 10, 2015, announcing the first meeting of the Technological Advisory Council for 2015. At its prior meeting on December 4, 2014, the Council had discussed possible work initiatives for 2015. These initiatives have been discussed in the interim within the FCC, with the TAC chairman, as well as with individual TAC members. At the April meeting, the FCC Technological Advisory Council will discuss its proposed work program for 2015. The FCC will attempt to accommodate as many people as possible. However, admittance will be limited to seating availability. Meetings are also broadcast live with open captioning over the Internet from the FCC Live Web page at *http://www.fcc.gov/live/.* The public may submit written comments before the meeting to: Walter Johnston, the FCC's Designated Federal Officer for Technological Advisory Council by email: *Walter.Johnston@fcc.gov* or U.S. Postal Service Mail (Walter Johnston, Federal Communications Commission, Room 7–A224, 445 12th Street SW., Washington, DC 20554). Open captioning will be provided for this event. Other reasonable accommodations for people with disabilities are available upon request. Requests for such accommodations should be submitted via email to *fcc504@fcc.gov* or by calling the Office of Engineering and Technology at 202–418–2470 (voice), (202) 418–1944 (fax). Such requests should include a detailed description of the accommodation needed. In addition, please include your contact information. Please allow at least five days advance notice; last minute requests will be accepted, but may be impossible to fill.

Federal Communications Commission.

Julius P. Knapp,

Chief, Office of Engineering and Technology.

[FR Doc. 2015–04202 Filed 2–26–15; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Notice to All Interested Parties of the Termination of the Receivership of 10201, American National Bank, Parma, Ohio

Notice is hereby given that the Federal Deposit Insurance Corporation (''FDIC'') as Receiver for American National Bank, Parma, Ohio (''the Receiver'') intends to terminate its receivership for said institution. The FDIC was appointed receiver of American National Bank on March 19, 2010. The liquidation of the receivership assets has been completed. To the extent permitted by available funds and in accordance with law, the Receiver will be making a final dividend payment to proven creditors.

Based upon the foregoing, the Receiver has determined that the continued existence of the receivership will serve no useful purpose. Consequently, notice is given that the receivership shall be terminated, to be effective no sooner than thirty days after the date of this Notice. If any person wishes to comment concerning the termination of the receivership, such comment must be made in writing and sent within thirty days of the date of this Notice to: Federal Deposit Insurance Corporation, Division of Resolutions and Receiverships, Attention: Receivership Oversight Department 34.1, 1601 Bryan Street, Dallas, TX 75201.

No comments concerning the termination of this receivership will be considered which are not sent within this time frame.

Dated: February 23, 2015.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2015–04043 Filed 2–26–15; 8:45 am]

BILLING CODE 6714–01–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0080]; [Docket 2015–0001; Sequence 2]

General Services Administration Acquisition Regulation; Information Collection; Contract Financing Final Payment (GSA Form 1142 Release of Claims)

AGENCY: Office of Acquisition Policy, GSA.

ACTION: Notice of request for comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement and the reinstatement of GSA Form 1142, Release of Claims, regarding final payment under construction and building services contract. GSA Form 1142 was inadvertently deleted as part of the rewrite of GSAR regulations on Contract Financing. GSA Contracting Officers have used this form to achieve uniformity and consistency in the release of claims process.

DATES: Submit comments on or before: April 28, 2015.

FOR FURTHER INFORMATION CONTACT: Ms. Dana Munson, General Services Acquisition Policy Division, GSA, (202) 357–9652 or email *Dana.Munson@gsa.gov.*

ADDRESSES: Submit comments identified by Information Collection 3090–0080, Contract Financing Final Payment; (GSA Form 1142, Release of Claims) by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching the OMB Control number 3090–0080. Select the link ''Comment Now'' that corresponds with ''Information Collection 3090–0080, Contract Financing Final Payment; (GSA Form 1142, Release of Claims).'' Follow the instructions on the screen. Please include your name, company name (if any), and ''Information Collection 3090–0080, Contract Financing Final Payment; (GSA Form 1142, Release of Claims),'' on your attached document.
- *Fax:* 202–501–4067.
- *Mail:* General Services Administration, Regulatory Secretariat (MVCB), 1800 F Street NW.,

Washington, DC 20405. ATTN: Ms. Hada Flowers/IC 3090–0080, Contract Financing Final Payment; (GSA Form 1142, Release of Claims).

Instructions: Please submit comments only and cite Information Collection 3090–0080, Contract Financing Final Payment; (GSA Form 1142, Release of Claims), in all correspondence related to this collection. All comments received will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided.

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration Acquisition Regulation (GSAR) clause 552.232–72 requires construction and building services contractors to submit a release of claims before final payment is made to ensure contractors are paid in accordance with their contract requirements and for work performed. GSA Form 1142, Release of Claims is used to achieve uniformity and consistency in the release of claims process.

B. Annual Reporting Burden

Respondents: 2000.

Responses per Respondent: 1.

Hours per Response: .10.

Total Burden Hours: 200.

C. Public Comment

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVCB), 1800 F Street NW., Washington, DC 20405, telephone (202) 501–4755. Please cite OMB Control No. 3090–0080, Contract Financing Final Payment; (GSA Form 1142, Release of Claims), in all correspondence.

Dated: February 24, 2015.

Jeffrey A. Koses,

Director, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2015–04116 Filed 2–26–15; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Determination and Declaration Regarding Emergency Use of New In Vitro Diagnostics for Detection of Enterovirus D68

AGENCY: Office of the Secretary, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: The Secretary of Health and Human Services (HHS) is issuing this notice pursuant to section 564 of the Federal Food, Drug, and Cosmetic (FD&C) Act, 21 U.S.C. 360bbb–3. On February 6, 2015, the Secretary determined that there is a significant potential for a public health emergency that has a significant potential to affect national security or the health and security of United States citizens living abroad and that involves enterovirus D68 (EV–D68). On the basis of this determination, she also declared that circumstances exist justifying the authorization of emergency use of new in vitro diagnostics for detection of EV–D68 pursuant to section 564 of the FD&C Act, subject to the terms of any authorization issued under that section.

DATES: The determination and declaration are effective February 6, 2015.

FOR FURTHER INFORMATION CONTACT: Karen Mason, Centers for Disease Control and Prevention, 1600 Clifton Road MS–A34, Atlanta, GA 30333, Telephone (404) 639–1297 (this is not a toll free number).

SUPPLEMENTARY INFORMATION:

I. Background

Under Section 564 of the FD&C Act, the Commissioner of the Food and Drug Administration (FDA), acting under delegated authority from the Secretary of HHS, may issue an Emergency Use Authorization (EUA) authorizing (1) the emergency use of an unapproved drug, an unapproved or uncleared device, or an unlicensed biological product; or (2) an unapproved use of an approved drug, approved or cleared device, or licensed biological product. Before an EUA may be issued, the Secretary of HHS must declare that circumstances exist justifying the authorization based on one of four determinations: (1) A determination by the Secretary of Homeland Security that there is a domestic emergency, or a significant potential for a domestic emergency, involving a heightened risk of attack with a chemical, biological, radiological, or nuclear (''CBRN'') agent or agents; (2) the identification of a material threat by the Secretary of Homeland Security pursuant to section 319F–2 of the Public Health Service (PHS) Act [1] sufficient to affect national security or the health and security of United States citizens living abroad; (3) a determination by the Secretary of Defense that there is a military emergency, or a significant potential for a military emergency, involving a heightened risk to United States military forces of attack with a CBRN agent or agents; or (4) a determination by the Secretary that there is a public health emergency, or a significant potential for a public health emergency, that affects, or has a significant potential to affect national security or the health and security of United States citizens living abroad, and that involves a CBRN agent or agents, or a disease or condition that may be attributable to such agent or agents.[2]

Based on any of these four determinations, the Secretary of HHS may then declare that circumstances exist that justify the EUA, at which point the FDA Commissioner may issue an EUA if the criteria for issuance of an authorization under section 564 of the FD&C Act are met. The Centers for Disease Control and Prevention (CDC), HHS, requested that the FDA, HHS, issue an EUA for new in vitro diagnostics for detection of EV–D68 to allow the Department to take preparedness measures based on information currently available about the EV–D68.

The determination of a significant potential for a public health emergency, and the declaration that circumstances exist justifying emergency use of new in vitro diagnostics for detection of EV–D68 by the Secretary of HHS, as described below, enable the FDA Commissioner to issue an EUA for in vitro diagnostics for detection of EV–D68 for emergency use under section 564 of the FD&C Act.

II. Determination by the Secretary of Health and Human Services

On February 6, 2015, pursuant to section 564 of the FD&C Act, I determined that there is a significant potential for a public health emergency that has a significant potential to affect national security or the health and

[1] 42 U.S.C. 247d–6b

[2] As amended by the Pandemic and All-Hazards Preparedness Reauthorization Act, Public Law 113–5, the Secretary may make determination of a public health emergency, or a significant potential for a public health emergency, under section 564 of the FD&C Act. The Secretary is no longer required to make a determination of a public health emergency in accordance with section 319 of the PHS Act, 42 U.S.C. 247d, to support a determination or declaration made under section 564 of the FD&C Act.