information collection should be submitted on or before April 12, 2023.

ADDRESSES: Comments should be sent to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. Your comment must be submitted into *www.reginfo.gov* per the above instructions for it to be considered. In addition to submitting in *www.reginfo.gov* also send a copy of your comment on the proposed information collection to Cathy Williams, FCC, via email to *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.* Include in the comments the OMB control number as shown in the **SUPPLEMENTARY INFORMATION** below.

FOR FURTHER INFORMATION CONTACT: For additional information or copies of the information collection, contact Cathy Williams at (202) 418–2918. To view a copy of this information collection request (ICR) submitted to OMB: (1) go to the web page *http://www.reginfo.gov/public/do/PRAMain,* (2) look for the section of the web page called ''Currently Under Review,'' (3) click on the downward-pointing arrow in the ''Select Agency'' box below the ''Currently Under Review'' heading, (4) select ''Federal Communications Commission'' from the list of agencies presented in the ''Select Agency'' box, (5) click the ''Submit'' button to the right of the ''Select Agency'' box, (6) when the list of FCC ICRs currently under review appears, look for the Title of this ICR and then click on the ICR Reference Number. A copy of the FCC submission to OMB will be displayed.

SUPPLEMENTARY INFORMATION: The Commission may not conduct or sponsor a collection of information unless it displays a currently valid Office of Management and Budget (OMB) control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB control number.

As part of its continuing effort to reduce paperwork burdens, as required by the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501–3520), the FCC invited the general public and other Federal Agencies to take this opportunity to comment on the following information collection. Comments are requested concerning: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission's burden estimates; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology. Pursuant to the Small Business Paperwork Relief Act of 2002, Public Law 107–198, see 44 U.S.C. 3506(c)(4), the FCC seeks specific comment on how it might ''further reduce the information collection burden for small business concerns with fewer than 25 employees.''

OMB Control Number: 3060–0548.

Title: Sections 76.1709 and 76.1620, Availability of Signals; Section 76.1614, Identification of Must-Carry Signals.

Type of Review: Extension without change of a currently approved collection.

Respondents: Businesses or other for-profit.

Number of Respondents and Responses: 4,103 respondents; 49,236 responses.

Estimated Time per Response: 0.5–1.0 hour.

Frequency of Response: Recordkeeping requirement, Third party disclosure requirement, On occasion reporting requirement.

Obligation to Respond: Voluntary.

Total Annual Burden: 24,618 hours.

Total Annual Cost: No cost.

Needs and Uses: 47 CFR 76.1709(a) states that the operator of every cable television system shall maintain for public inspection a file containing a list of all broadcast television stations carried by its system in fulfillment of the must-carry requirements. Such list shall include the call sign; community of license, broadcast channel number, cable channel number, and in the case of a noncommercial educational broadcast station, whether that station was carried by the cable system on March 29, 1990. 47 CFR 76.1614 and 47 CFR 76.1709(c) each state that a cable operator shall respond in writing within 30 days to any written request by any person for the identification of the signals carried on its system in fulfillment of the must-carry requirements. In addition, 47 CFR 76.1614 states that the required written response may be delivered by email, if the consumer used email to make the request or complaint directly to the cable operator, or if the consumer specifies email as the preferred delivery method in the request or complaint.

47 CFR 76.1620, pursuant to 47 U.S.C. 614(b)(7), states that if a cable operator authorizes subscribers to install additional receiver connections, but does not provide the subscriber with such connections, or with the equipment and materials for such connections, the operator shall notify such subscribers of all broadcast stations carried on the cable system which cannot be viewed via cable without a converter box and shall offer to sell or lease such a converter box to such subscribers. Such notification must be provided by June 2, 1993, and annually thereafter and to each new subscriber upon initial installation. The notice, which may be included in routine billing statements, shall identify the signals that are unavailable without an additional connection, the manner for obtaining such additional connection and instructions for installation. 47 CFR 76.1600(a) provides that written information provided by cable operators to subscribers or customers pursuant to § 76.1620 may be delivered electronically by email to any subscriber who has not opted out of electronic delivery if the entity: (1) Sends the notice to the subscriber's or customer's verified email address; (2) Provides either the entirety of the written information or a weblink to the written information in the notice; and (3) Includes, in the body of the notice, a telephone number that is clearly and prominently presented to subscribers so that it is readily identifiable as an opt-out mechanism that will allow subscribers to continue to receive paper copies of the written material.

Note: These recordkeeping and notification requirements ensure that subscribers are aware of the broadcast stations carried in compliance with the Commission's cable must-carry rules, see 47 CFR 76.56.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2023–05048 Filed 3–10–23; 8:45 am]

BILLING CODE 6712–01–P

GENERAL SERVICES ADMINISTRATION

[Notice–MA–2023–01; Docket No. 2023–0002; Sequence No. 1]

Revision to Foreign Gift Minimal Value

AGENCY: Office of Government-wide Policy (OGP), General Services Administration (GSA).

ACTION: Notice of GSA Bulletin FMR B–52, Foreign Gift and Decoration Minimal Value.

SUMMARY: GSA, in consultation with the U.S. Department of State, must redefine the minimal value of foreign gift items to reflect changes in the Consumer Price

Index (CPI) for the preceding 3-year period, as specified under the law concerning the Receipt and Disposition of Foreign Gifts and Decorations. The minimal value was last defined effective January 1, 2020, and must be redefined effective as of January 1, 2023. This bulletin cancels FMR Bulletin B–50, ''Foreign Gift and Decoration Minimal Value,'' issued March 10, 2020, as this bulletin provides updated information on the same topic.

DATES: *Applicability Date:* January 1, 2023.

This notice applies to foreign gifts and decorations received on or after January 1, 2023.

FOR FURTHER INFORMATION CONTACT: For clarification of content, contact Mr. William Garrett, Director, Personal Property Policy, Office of Government-wide Policy, Office of Asset and Transportation Management, at 202–368–8163, or by email at *william.garrett@gsa.gov.* Please cite Notice of GSA Bulletin FMR B–52.

SUPPLEMENTARY INFORMATION:

Background

Foreign gifts and decorations above the GSA-defined minimal value are handled differently than lesser-valued foreign gifts and decorations under the provisions of 5 U.S.C. 7342 and FMR 102–42.

Foreign gifts and decorations above the minimal value become the property of the Federal Government and must be reported to GSA for disposal if not immediately needed by the agency for official purposes. Additionally, those items initially retained by the agencies for official use are reported to GSA upon termination of official use.

The foreign gifts and decorations minimal value was last redefined effective January 1, 2020, at $415, and therefore, must be redefined as of January 1, 2023, to reflect the CPI increase of 15.33 percent for the preceding three years.

Pursuant to FMR 102–42.10, the approved revised minimal value will be published in an FMR Bulletin posted on OGP's website (*www.gsa.gov/foreigngifts*).

Calculations using the consumer prices over the past three years show that the minimal value must increase 15.33 percent from its current $415, which yields an amount of $478.62. As in previous years, GSA is rounding the amount to the nearest five dollar increments.

Therefore, GSA is adjusting the new minimal value to $480.00. Per FMR 102–42.10, an agency may, by regulation, specify a lower value than this Government-wide value for its agency employees.

FMR Bulletin 52 is available at *https://www.gsa.gov/policy-regulations/regulations/federal-management-regulation/federal-management-regulation-fmr-related-files#PersonalPropertyManagement.*

Krystal J. Brumfield,

Associate Administrator, Office of Government-wide Policy.

[FR Doc. 2023–05093 Filed 3–10–23; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–0577]

Authorization of Emergency Use of a Drug Product During the COVID–19 Pandemic; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the issuance of an Emergency Use Authorization (EUA) (the Authorization) under the Federal Food, Drug, and Cosmetic Act (FD&C Act) for use during the COVID–19 pandemic. FDA has issued one Authorization for the drug product KINERET (anakinra) as requested by Swedish Orphan Biovitrum AB (Sobi). The Authorization contains, among other things, conditions on the emergency use of the authorized product. The Authorization follows the February 4, 2020, determination by the Secretary of Health and Human Services (HHS) that there is a public health emergency that has a significant potential to affect national security or the health and security of U.S. citizens living abroad and that involves a novel (new) coronavirus. The virus, now named SARS–CoV–2, causes the illness COVID–19. On the basis of such determination, the Secretary of HHS declared on March 27, 2020, that circumstances exist justifying the authorization of emergency use of drugs and biological products during the COVID–19 pandemic, pursuant to the FD&C Act, subject to the terms of any authorization issued under that section. The Authorization, which includes an explanation of the reasons for issuance, is reprinted in this document.

DATES: The Authorization is effective as of November 8, 2022.

ADDRESSES: Submit written requests for a single copy of the EUA to the Office of Executive Programs, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, 6th Floor, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your request or include a Fax number to which the Authorization may be sent. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the Authorization.

FOR FURTHER INFORMATION CONTACT: Johanna McLatchy, Office of Executive Programs, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, 6th Floor, Silver Spring, MD 20993–0002, 301–796–3200 (this is not a toll free number).

SUPPLEMENTARY INFORMATION:

I. Background

Section 564 of the FD&C Act (21 U.S.C. 360bbb–3) allows FDA to strengthen public health protections against biological, chemical, nuclear, and radiological agents. Among other things, section 564 of the FD&C Act allows FDA to authorize the use of an unapproved medical product or an unapproved use of an approved medical product in certain situations. With this EUA authority, FDA can help ensure that medical countermeasures may be used in emergencies to diagnose, treat, or prevent serious or life-threatening diseases or conditions caused by biological, chemical, nuclear, or radiological agents when there are no adequate, approved, and available alternatives (among other criteria).

II. Criteria for EUA Authorization

Section 564(b)(1) of the FD&C Act provides that, before an EUA may be issued, the Secretary of HHS must declare that circumstances exist justifying the authorization based on one of the following grounds: (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 biological, chemical, radiological, or nuclear agent or agents; (2) 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 U.S. military forces, including personnel operating under the authority of title 10 or title 50, U.S. Code, of attack with (A) a biological, chemical, radiological, or nuclear agent or agents; or (B) an agent or agents that may cause, or are otherwise associated