

the Tariff Act of 1930; this notice is published pursuant to § 207.21 of the Commission's rules.

By order of the Commission.

Issued: June 6, 2024.

**Lisa Barton,**

*Secretary to the Commission.*

[FR Doc. 2024–12776 Filed 6–11–24; 8:45 am]

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## INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–721 and 731–TA–1689 (Preliminary)]

### Alkyl Phosphate Esters From China; Determinations

On the basis of the record<sup>1</sup> developed in the subject investigations, the United States International Trade Commission (“Commission”) determines, pursuant to the Tariff Act of 1930 (“the Act”), that there is a reasonable indication that an industry in the United States is materially injured by reason of imports of alkyl phosphate esters from China, provided for in subheading 2919.90.50 of the Harmonized Tariff Schedule of the United States, that are alleged to be sold in the United States at less than fair value (“LTFV”) and imports of the subject merchandise from China that are alleged to be subsidized by the government of China.<sup>2</sup>

### Commencement of Final Phase Investigations

Pursuant to section 207.18 of the Commission's rules, the Commission also gives notice of the commencement of the final phase of its investigations. The Commission will issue a final phase notice of scheduling, which will be published in the **Federal Register** as provided in § 207.21 of the Commission's rules, upon notice from the U.S. Department of Commerce (“Commerce”) of affirmative preliminary determinations in the investigations under §§ 703(b) or 733(b) of the Act, or, if the preliminary determinations are negative, upon notice of affirmative final determinations in those investigations under §§ 705(a) or 735(a) of the Act. Parties that filed entries of appearance in the preliminary phase of the investigations need not enter a separate appearance for the final phase of the investigations. Any other party may file an entry of appearance for the final phase of the investigations after

publication of the final phase notice of scheduling. Industrial users, and, if the merchandise under investigation is sold at the retail level, representative consumer organizations have the right to appear as parties in Commission antidumping and countervailing duty investigations. The Secretary will prepare a public service list containing the names and addresses of all persons, or their representatives, who are parties to the investigations. As provided in section 207.20 of the Commission's rules, the Director of the Office of Investigations will circulate draft questionnaires for the final phase of the investigations to parties to the investigations, placing copies on the Commission's Electronic Document Information System (EDIS, <https://edis.usitc.gov>), for comment.

### Background

On April 23, 2024, ICL–IP America, Inc., St. Louis, Missouri filed petitions with the Commission and Commerce, alleging that an industry in the United States is materially injured or threatened with material injury by reason of subsidized imports of alkyl phosphate esters from China and LTFV imports of alkyl phosphate esters from China. Accordingly, effective April 23, 2024, the Commission instituted countervailing duty investigation No. 701–TA–721 and antidumping duty investigation No. 731–TA–1689 (Preliminary).

Notice of the institution of the Commission's investigations and of a public conference to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the **Federal Register** of April 30, 2024 (89 FR 34270). The Commission conducted its conference on May 14, 2024. All persons who requested the opportunity were permitted to participate.

The Commission made these determinations pursuant to §§ 703(a) and 733(a) of the Act (19 U.S.C. 1671b(a) and 1673b(a)). It completed and filed its determinations in these investigations on June 7, 2024. The views of the Commission are contained in USITC Publication 5516 (June 2024), entitled *Alkyl Phosphate Esters from China: Investigation Nos. 701–TA–721 and 731–TA–1689 (Preliminary)*.

By order of the Commission.

Issued: June 7, 2024.

**Lisa Barton,**

*Secretary to the Commission.*

[FR Doc. 2024–12876 Filed 6–11–24; 8:45 am]

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## DEPARTMENT OF JUSTICE

[OMB Number 1105–NEW]

### Agency Information Collection Activities; Proposed eCollection eComments Requested; New Collection; SSO–012, Request To Reevaluate Special Security Officer's Medical Qualification

**AGENCY:** U.S. Marshals Service, Department of Justice.

**ACTION:** 60-Day notice.

**SUMMARY:** The U.S. Marshals Service, Department of Justice (DOJ), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995.

**DATES:** Comments are encouraged and will be accepted for 60 days until August 12, 2024.

**FOR FURTHER INFORMATION CONTACT:** If you have additional comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact [Benjamin Cho/Business Component/Management Support Division, Address U.S. Marshals Service Headquarters, 1215 S Clark St., Ste. 10022B, Arlington, VA 22202–4837, Phone 240–401–0008 AND [benjamin.cho@usdoj.gov](mailto:benjamin.cho@usdoj.gov)].

**SUPPLEMENTARY INFORMATION:** Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Bureau of Justice Statistics, including whether the information will have practical utility;
- Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Evaluate whether and if so how the quality, utility, and clarity of the information to be collected can be enhanced; and
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g.,

<sup>1</sup> The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

<sup>2</sup> 89 FR 43801 and 89 FR 43821 (May 20, 2024).

permitting electronic submission of responses.

**Abstract:** Form SSO-012 must be completed by a Special Security Officer (SSO)'s attending physician when an SSO is returning to perform security services for the U.S. Marshals Service after recovering from an injury, extended illness, and/or outpatient or inpatient surgery to ensure the SSO is medically qualified to return to duty.

**Overview of This Information Collection**

1. *Type of Information Collection:* New collection

2. *The Title of the Form/Collection:* SSO-012, Request to Reevaluate Special Security Officer's Medical Qualification

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* SSO-012

4. *Affected public who will be asked or required to respond, as well as the obligation to respond:* Individuals or households. The obligation to respond is voluntary.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* The total or estimated number

of respondents for SSO-012, Request to Reevaluate Special Security Officer's Medical Qualification is 27. The time per response is 30 minutes to complete SSO-012, Request to Reevaluate Special Security Officer's Medical Qualification.

6. An estimate of the total annual burden (in hours) associated with the collection: Ex: The total annual burden hours for this collection is 13.5 hours (27 respondents \* 0.5 hours per response).

7. An estimate of the total annual cost burden associated with the collection, if applicable:

**TOTAL BURDEN HOURS**

Activity	Number of respondents	Frequency	Total annual responses	Time per response	Total annual burden (hours)
Survey (individuals or households) .....	27	1/annually .....	27	30 min .....	13.5 hrs.

*If additional information is required contact:* Darwin Arceo, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, 4W-218, Washington, DC.

Dated: June 7, 2024.

**Darwin Arceo,**

*Department Clearance Officer for PRA, U.S. Department of Justice.*

[FR Doc. 2024-12828 Filed 6-11-24; 8:45 am]

**BILLING CODE 4410-04-P**

**DEPARTMENT OF JUSTICE**

[CPCLO Order No. 003-2024]

**Privacy Act of 1974; Systems of Records**

**AGENCY:** Federal Bureau of Prisons, United States Department of Justice.

**ACTION:** Notice of a modified system of records.

**SUMMARY:** Pursuant to the Privacy Act of 1974 and Office of Management and Budget (OMB) Circular No. A-108, notice is hereby given that the Federal Bureau of Prisons (hereinafter the Bureau or FBOP), a component within the United States Department of Justice (DOJ or Department), proposes to modify a system of records notice titled, "Inmate Physical and Mental Health Record System," JUSTICE/BOP-007, last modified on May 25, 2017, in order to consolidate previously published modifications of the system of records into one document to promote transparency. The modifications also

incorporate OMB guidance, technological advancements, and changes to four (4) routine uses.

**DATES:** In accordance with 5 U.S.C. 552a(e)(4) and (11), this notice is applicable upon publication, subject to a 30-day period in which to comment on the routine uses, described below. Therefore, please submit any comments by July 12, 2024.

**ADDRESSES:** The public, OMB, and Congress are invited to submit any comments: by mail to the United States Department of Justice, Office of Privacy and Civil Liberties, ATTN: Privacy Analyst, Two Constitution Square (2CON), 145 N Street NE, Suite 8W.300, Washington, DC 20002; by facsimile at 202-307-0693; or by email at [privacy.compliance@usdoj.gov](mailto:privacy.compliance@usdoj.gov). To ensure proper handling, please reference the above CPCLO Order No. on your correspondence.

**FOR FURTHER INFORMATION CONTACT:** Eugene Baime, Supervisory Attorney, Freedom of Information Act and Privacy Act Section, Office of General Counsel, Federal Bureau of Prisons, 320 First Street NW, Suite 924A, Washington, DC 20534, [BOP-OGC-EFOIA-S@BOP.GOV](mailto:BOP-OGC-EFOIA-S@BOP.GOV), 202-616-7750.

**SUPPLEMENTARY INFORMATION:** FBOP is modifying this system of records to consolidate earlier modifications of the system of records into one document to promote transparency. For a detailed list of previously published modifications, please review the "History" section below. Additionally, modifications to the system of records have been made to incorporate OMB guidance, technological advancements, and the

modification of four (4) routine uses. Pursuant to OMB Circular No. A-108, various sections were rearranged, and various section titles were edited.

FBOP also has changed the "System Location" section by adding to the list of locations where records may be maintained contractor-operated correctional facilities, National Archive Centers, secure cloud computing environments, and contracted storage facilities. FBOP has updated the **ADDRESSES** section to reflect administrative changes. FBOP has made a slight change to the Purpose(s) of the System to add the Attorney General as one of the parties that this system assists. FBOP has updated the "Categories of Individuals Covered by the System" to clarify that it includes those individuals under custody for criminal and civil commitments. FBOP has updated the "Categories of Records" in the system to include additional identifying particulars and information regarding specific health conditions.

FBOP has added modified routine uses that: allow medical health care professionals to access former inmates' medical records for continuity of care purposes, and eliminating the requirement the records only be provided for pre-existing condition (see RU b); include specific mention of the United States Probation Office (see RU c); permit Members of Congress to request and receive records at the bequest of the authorized next-of-kin for inmates who are mentally or physically incapacitated (see RU g); and account for authorized disclosures related to Coronavirus-19 (SARS-CoV-2) and other unknown infectious diseases (see