operation of a nationwide interoperable public safety broadband network. The FirstNet Authority Board is responsible for making strategic decisions regarding the operations of the FirstNet Authority.

*Matters to be Considered:* The FirstNet Authority will post a detailed agenda for the Combined Board and Board Committees Meeting on *FirstNet.gov* prior to the meeting. The agenda topics are subject to change. Please note that the subjects discussed by the Board and Board Committees may involve commercial or financial information that is privileged or confidential, or other legal matters affecting the FirstNet Authority. As such, the Board may, by majority vote, close the meeting only for the time necessary to preserve the confidentiality of such information, pursuant to 47 U.S.C. 1424(e)(2).

*Other Information:* The public Combined Board and Board Committees Meeting is accessible to people with disabilities. Individuals requiring accommodations, such as sign language interpretation or other ancillary aids, are asked to notify Jennifer Watts at (571) 665–6178 or email: *Jennifer.Watts@FirstNet.gov* before the meeting.

*Records:* The FirstNet Authority maintains records of all Board proceedings. Minutes of the Combined Board and Board Committees Meeting will be available on *FirstNet.gov.*

Dated: June 14, 2024.

**Jennifer Watts,**

*Board Secretary, First Responder Network Authority.*

[FR Doc. 2024–13499 Filed 6–18–24; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF COMMERCE

### Foreign-Trade Zones Board

**[B–33–2024]**

### Foreign-Trade Zone 75; Application for Expansion of Subzone 75C; Intel Corporation; Phoenix, Arizona

An application has been submitted to the Foreign-Trade Zones (FTZ) Board by the City of Phoenix, grantee of FTZ 75, requesting an expansion of Subzone 75C on behalf of Intel Corporation. The application was submitted pursuant to the provisions of the Foreign-Trade Zones Act, as amended (19 U.S.C. 81a–81u), and the regulations of the FTZ Board (15 CFR part 400). It was formally docketed on June 13, 2024.

The applicant is requesting authority to expand the subzone to include a new site located at 3929 West Lower Buckeye Road, Phoenix (Site 6—1.7 acres).

In accordance with the FTZ Board's regulations, Qahira El-Amin of the FTZ Staff is designated examiner to review the application and make recommendations to the FTZ Board.

Public comment is invited from interested parties. Submissions shall be addressed to the FTZ Board's Executive Secretary and sent to: *ftz@trade.gov.* The closing period for their receipt is July 30, 2024. Rebuttal comments in response to material submitted during the foregoing period may be submitted during the subsequent 15-day period to August 14, 2024.

A copy of the application will be available for public inspection in the ''Online FTZ Information Section'' section of the FTZ Board's website, which is accessible via *www.trade.gov/ftz.*

For further information, contact Qahira El-Amin at *Qahira.El-Amin@trade.gov.*

Dated: June 14, 2024.

**Elizabeth Whiteman,**

*Executive Secretary.*

[FR Doc. 2024–13451 Filed 6–18–24; 8:45 am]

**BILLING CODE 3510–DS–P**

## DEPARTMENT OF COMMERCE

### Foreign-Trade Zones Board

**[B–31–2024]**

### Foreign-Trade Zone (FTZ) 230, Notification of Proposed Production Activity; Patheon Softgels; (Pharmaceutical Products); High Point, North Carolina

The Piedmont Triad Partnership, grantee of FTZ 230, submitted a notification of proposed production activity to the FTZ Board (the Board) on behalf of Patheon Softgels (Patheon), for Patheon's facility in High Point, North Carolina within Subzone 230C. The notification conforming to the requirements of the Board's regulations (15 CFR 400.22) was received on June 12, 2024.

Pursuant to 15 CFR 400.14(b), FTZ production activity would be limited to the specific foreign-status material(s)/component(s) and specific finished product(s) described in the submitted notification (summarized below) and subsequently authorized by the Board. The benefits that may stem from conducting production activity under FTZ procedures are explained in the background section of the Board's website—accessible via *www.trade.gov/ftz.* The proposed finished product(s) and material(s)/component(s) would be added to the production authority that the Board previously approved for the operation, as reflected on the Board's website.

The proposed finished products include CU06–1004 and northenindrone acetate (duty-free).

The proposed foreign-status materials/components include: CU06–1004(active pharmaceutical ingredient); titanium dioxide (35%); mono diglycerides; surfactants; glycerol monolinoneate; red iron oxide; yellow iron oxide; medium chain triglycerides; vitamin E; sesame oil nf; linoleoyl polyoxyl-6 glycerides; ethinylestradiol micronized; glycerin and northenindrone acetate (active pharmaceutical ingredient) (duty rate ranges from duty-free to 6.5%).

Public comment is invited from interested parties. Submissions shall be addressed to the Board's Executive Secretary and sent to: *ftz@trade.gov.* The closing period for their receipt is July 30, 2024.

A copy of the notification will be available for public inspection in the ''Online FTZ Information System'' section of the Board's website.

For further information, contact Kolade Osho at *Kolade.Osho@trade.gov.*

Dated: June 13, 2024.

**Elizabeth Whiteman,**

*Executive Secretary.*

[FR Doc. 2024–13449 Filed 6–18–24; 8:45 am]

**BILLING CODE 3510–DS–P**

## DEPARTMENT OF DEFENSE

### Office of the Secretary

**[Docket ID: DoD–2024–HA–0068]**

### Proposed Collection; Comment Request

**AGENCY:** The Office of the Assistant Secretary of Defense for Health Affairs (OASD(HA)), Department of Defense (DoD).

**ACTION:** 60-Day information collection notice.

**SUMMARY:** In compliance with the *Paperwork Reduction Act of 1995,* the Defense Health Agency (DHA) announces a proposed public information collection and seeks public comment on the provisions thereof. Comments are invited on: whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; the accuracy of the agency's estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and

clarity of the information to be collected; and ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology.

**DATES:** Consideration will be given to all comments received by August 19, 2024.

**ADDRESSES:** You may submit comments, identified by docket number and title, by any of the following methods:

*Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.

*Mail:* Department of Defense, Office of the Assistant to the Secretary of Defense for Privacy, Civil Liberties, and Transparency, Regulatory Directorate, 4800 Mark Center Drive, Mailbox #24, Suite 08D09, Alexandria, VA 22350–1700.

*Instructions:* All submissions received must include the agency name, docket number and title for this **Federal Register** document. The general policy for comments and other submissions from members of the public is to make these submissions available for public viewing on the internet at *http://www.regulations.gov* as they are received without change, including any personal identifiers or contact information.

**FOR FURTHER INFORMATION CONTACT:** To request more information on this proposed information collection or to obtain a copy of the proposal and associated collection instruments, please write to Defense Health Agency, 7700 Arlington Blvd., Falls Church, VA 22042, Amanda Grifka, 703–681–1771.

**SUPPLEMENTARY INFORMATION:**

*Title; Associated Form; and OMB Number:* Department of Defense (DoD) Patient Safety Culture Survey; OMB Control Number 0720–0034.

*Needs and Uses:* The 2001 National Defense Authorization Act contains specific sections addressing patient safety in military and veterans' health care systems. This legislation states that the Secretary of Defense shall establish a patient care error reporting and management system to study occurrences of errors in patient care and that one purpose of the system should be to ''identify systemic factors that are associated with such occurrences'' and ''to provide for action to be taken to correct the identified systemic factors'' (sec. 754, items b2 and b3). In addition, the legislation states that the Secretary shall ''continue research and development investments to improve communication, coordination, and teamwork in the provision of health care'' (Sec. 754, item d4). In its ongoing response to this legislation and in support of its mission to ''promote a culture of safety to eliminate preventable patient harm by engaging, educating and equipping patient-care teams to institutionalize evidence-based safe practices,'' the DoD Patient Safety Program plans to field the DoD Patient Safety Culture Survey. The Culture Survey is based on the Department of Health and Human Services' Agency for Healthcare Research and Quality's validated survey instrument. The survey obtains Military Health System staff opinions on patient safety issues such as teamwork, communications, medical error occurrence and response, error reporting, and overall perceptions of patient safety.

*Affected Public:* Federal government; individuals or households.

*Annual Burden Hours:* 1,228.

*Number of Respondents:* 7,370.

*Responses per Respondent:* 1.

*Annual Responses:* 7,370.

*Average Burden per Response:* 10 minutes.

*Frequency:* As required.

Dated: June 12, 2024.

**Aaron T. Siegel,**

*Alternate OSD Federal Register Liaison Officer, Department of Defense.*

[FR Doc. 2024–13462 Filed 6–18–24; 8:45 am]

**BILLING CODE 6001–FR–P**

## DEPARTMENT OF DEFENSE

### Office of the Secretary

**[Docket ID: DoD–2024–OS–0069]**

### Proposed Collection; Comment Request

**AGENCY:** Office of the Under Secretary of Defense for Personnel and Readiness (OUSD(P&R)), Department of Defense (DoD).

**ACTION:** 60-Day information collection notice.

**SUMMARY:** In compliance with the *Paperwork Reduction Act of 1995,* the Under Secretary of Defense for Personnel and Readiness announces the proposed public information collections and seeks public comment on the provisions thereof. Comments are invited on: whether the proposed collections of information are necessary for the proper performance of the functions of the agencies, including whether the information shall have practical utility; the accuracy of the agencies' estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; and ways to minimize the burden of the information collections on respondents, including through the use of automated collection techniques or other forms of information technology.

**DATES:** Consideration will be given to all comments received by August 19, 2024.

**ADDRESSES:** You may submit comments, identified by docket number and title, by any of the following methods:

*Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.

*Mail:* Department of Defense, Office of the Assistant to the Secretary of Defense for Privacy, Civil Liberties, and Transparency, Regulatory Directorate, 4800 Mark Center Drive, Mailbox #24, Suite 08D09, Alexandria, VA 22350–1700.

*Instructions:* All submissions received must include the agency name, docket number and title for this **Federal Register** document. The general policy for comments and other submissions from members of the public is to make these submissions available for public viewing on the internet at *http://www.regulations.gov* as they are received without change, including any personal identifiers or contact information.

**FOR FURTHER INFORMATION CONTACT:** To request more information on these proposed information collections or to obtain a copy of the proposal and associated collection instruments, please write to DoD, Physical Disability Board of Review (PDBR), 3351 Celmers Lane, Joint Base Andrews 20762, Phyllis M. Joyner, 240–612–4392.

**SUPPLEMENTARY INFORMATION:**

*Title; Associated Form; and OMB Number:* Application for Review by the PDBR; DD Form 294; OMB Control Number 0704–0453.

*Needs and Uses:* The Fiscal Year 2008 National Defense Authorization Act amended Title 10, United States Code by adding Section 1554a. That provision of law directs the Secretary of Defense to establish a board of review to review the disability determinations of covered individuals by Physical Evaluation Boards. Covered individuals are members and former members of the armed forces who, during the period beginning on September 11, 2001, and ending on December 31, 2009, were (1) separated from the armed forces due to unfitness for duty due to a medical condition with a disability rating of 20 percent disabled or less; and (2) are found to be not eligible for retirement. On June 27, 2008, The Department of Defense published DoDI 6040.44, which provides the guidance for this process.

The DD Form 294, ''Application for Review by the PDBR of the Rating Awarded Accompanying a Medical