

DATES: Comments must be submitted on or before October 21, 2022.

ADDRESSES: Written comments and recommendations for the proposed information collection should be submitted within 30 days of this notice's publication to OIRA, at <https://www.reginfo.gov/public/do/PRAMain>. Please find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the website's search function. Comments can be entered electronically by clicking on the "comment" button next to the information collection on the "OIRA Information Collections Under Review" page, or the "View ICR—Agency Submission" page. A copy of the supporting statement for the collection of information discussed herein may be obtained by visiting <https://www.reginfo.gov/public/do/PRAMain>.

In addition to the submission of comments to <https://Reginfo.gov> as indicated above, a copy of all comments submitted to OIRA may also be submitted to the Commodity Futures Trading Commission (the "Commission" or "CFTC") by clicking on the "Submit Comment" box next to the descriptive entry for OMB Control No. 3038–0017, at <https://comments.cftc.gov/FederalRegister/PublicInfo.aspx>.

Or by either of the following methods:

- **Mail:** Christopher Kirkpatrick, Secretary of the Commission, Commodity Futures Trading Commission, Three Lafayette Centre, 1155 21st Street NW, Washington, DC 20581.

- **Hand Delivery/Courier:** Same as Mail above.

All comments must be submitted in English, or if not, accompanied by an English translation. Comments submitted to the Commission should include only information that you wish to make available publicly. If you wish the Commission to consider information that you believe is exempt from disclosure under the Freedom of Information Act, a petition for confidential treatment of the exempt information may be submitted according to the procedures established in § 145.9 of the Commission's regulations.¹ The Commission reserves the right, but shall have no obligation, to review, pre-screen, filter, redact, refuse or remove any or all of your submission from <https://www.cftc.gov> that it may deem to be inappropriate for publication, such as obscene language. All submissions that have been redacted or removed that contain comments on the merits of the

ICR will be retained in the public comment file and will be considered as required under the Administrative Procedure Act and other applicable laws, and may be accessible under the Freedom of Information Act.

FOR FURTHER INFORMATION CONTACT:

Adam Charnisky, Market Analyst, Division of Market Oversight, Commodity Futures Trading Commission, (312) 596–0630; email: acharnisky@cftc.gov; and refer to "OMB Control No. 3038–0017."

SUPPLEMENTARY INFORMATION:

Title: Market Surveys (OMB Control No. 3038–0017). This is a request for extension of a currently approved information collection.

Abstract: Under Commission Rule 21.02, 17 CFR 21.02, upon call by the Commission, information must be furnished related to futures or options positions held or introduced by futures commission merchants, members of contract markets, introducing brokers, and foreign brokers and, for options positions, by each reporting market. This rule is designed to assist the Commission in prevention of market manipulation and is promulgated pursuant to the Commission's rulemaking authority contained in section 8a of the Commodity Exchange Act, 7 U.S.C. 12a (2010).

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. On July 12, 2022, the Commission published in the **Federal Register** notice of the proposed extension of this information collection and provided 60 days for public comment on the proposed extension, 87 FR 41293 ("60-Day Notice"). The Commission did not receive any relevant comments on the 60-Day Notice.

Burden Statement: The Commission is updating its estimate of the burden for this collection for Market Surveys (OMB Control No. 3038–0017). The Commission estimates the burden of this collection of information as follows:

Estimated Number of Respondents: 100.

Estimated Average Burden Hours per Respondent: 1.75 hours.

Estimated Total Annual Burden Hours: 175 hours.

Frequency of Collection: Annually.

There are no capital costs or operating and maintenance costs associated with this collection.

(Authority: 44 U.S.C. 3501 *et seq.*)

Dated: September 16, 2022.

Robert Sidman,

Deputy Secretary of the Commission.

[FR Doc. 2022–20466 Filed 9–20–22; 8:45 am]

BILLING CODE 6351–01–P

COMMODITY FUTURES TRADING COMMISSION

Agency Information Collection Activities Under OMB Review

AGENCY: Commodity Futures Trading Commission.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995 (PRA), this notice announces that the Information Collection Request (ICR) abstracted below has been forwarded to the Office of Information and Regulatory Affairs (OIRA) of the Office of Management and Budget (OMB) for review and comment. The ICR describes the nature of the information collection and its expected costs and burden.

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In addition to the submission of comments to <https://Reginfo.gov> as indicated above, a copy of all comments submitted to OIRA may also be submitted to the Commodity Futures Trading Commission (the Commission or CFTC) by clicking on the "Submit Comment" box next to the descriptive entry for OMB Control No. 3038–0099, at <https://comments.cftc.gov/FederalRegister/PublicInfo.aspx>, or by any of the following methods:

- **Mail:** Christopher Kirkpatrick, Secretary of the Commission, Commodity Futures Trading Commission, Three Lafayette Centre, 1155 21st Street NW, Washington, DC 20581.

¹ 17 CFR 145.9.

• *Hand Delivery/Courier*: Same as Mail above.

All comments must be submitted in English, or if not, accompanied by an English translation. Comments submitted to the Commission should include only information that you wish to make available publicly. If you wish the Commission to consider information that you believe is exempt from disclosure under the Freedom of Information Act, a petition for confidential treatment of the exempt information may be submitted according to the procedures established in § 145.9 of the Commission's regulations.¹ The Commission reserves the right, but shall have no obligation, to review, pre-screen, filter, redact, refuse or remove any or all of your submission from <https://www.cftc.gov> that it may deem to be inappropriate for publication, such as obscene language. All submissions that have been redacted or removed that contain comments on the merits of the ICR will be retained in the public comment file and will be considered as required under the Administrative Procedure Act and other applicable laws, and may be accessible under the Freedom of Information Act.

FOR FURTHER INFORMATION CONTACT:

Roger Smith, Associate Chief Counsel, Division of Market Oversight, Commodity Futures Trading Commission, (202) 418-5344; email: RSmith@CFTC.gov, or Rebecca Mersand, Paralegal Specialist, Division of Market Oversight, Commodity Futures Trading Commission, 202-941-8910; email: RMersand@CFTC.gov, and refer to OMB Control No. 3038-0099.

SUPPLEMENTARY INFORMATION:

Title: Process for a Swap Execution Facility or Designated Contract Market to Make a Swap Available to Trade (OMB Control No. 3038-0099). This is a request for extension of a currently approved information collection.

Abstract: The collection of information is needed to help determine which swaps should be subject to the trade execution requirement under section 2(h)(8) of the Commodity Exchange Act pursuant to section 723 of the Dodd-Frank Wall Street Reform and Consumer Protection Act. A SEF or DCM that submits a determination that a swap is available to trade must address at least one of several factors to demonstrate that the swap is suitable for trading pursuant to the trade execution requirement. The Commission uses the collection of information to facilitate the application of the trade execution requirement and the requirements

associated with methods of execution under parts 37 and 38 of the Commission's regulations.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. On July 13, 2022, the Commission published in the **Federal Register** notice of the proposed extension of this information collection and provided 60 days for public comment on the proposed extension, 87 FR 41674 (60-Day Notice). The Commission did not receive any substantive comments on the 60-Day Notice.

Burden Statement: The Commission estimates the burden of reviewing the prescribed factors and data to make a determination for this collection to be 16 hours per response.

Respondents/Affected Entities: SEFs, DCMs.

Estimated Number of Respondents: 5.

Estimated Average Burden Hours per Respondent: 16.

Estimated Total Annual Burden Hours: 80.

Frequency of Collection: On occasion.

There are no capital costs or operating and maintenance costs associated with this collection.

(Authority: 44 U.S.C. 3501 *et seq.*)

Dated: September 16, 2022.

Robert Sidman,

Deputy Secretary of the Commission.

[FR Doc. 2022-20465 Filed 9-20-22; 8:45 am]

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DEPARTMENT OF DEFENSE

Office of the Secretary

Uniform Formulary Beneficiary Advisory Panel; Notice of Federal Advisory Committee

AGENCY: Under Secretary of Defense for Personnel and Readiness, Department of Defense (DoD).

ACTION: Notice of Federal Advisory Committee meeting.

SUMMARY: The DoD is publishing this notice to announce that the following Federal Advisory Committee meeting of the Uniform Formulary Beneficiary Advisory Panel (UF BAP) will take place.

DATES: Open to the public Thursday, September 29, 2022, 10 a.m.–1 p.m. (eastern standard time).

ADDRESSES: The meeting will be held telephonically or via conference call. The phone number for the remote access on September 29, 2022 is: CONUS: 1–

800-369-2046; OCONUS: 1-203-827-7030; Participant Code: 8546285.

These numbers and the dial-in instructions will also be posted on the UF BAP website at: <https://www.health.mil/Military-Health-Topics/Access-Cost-Quality-and-Safety/Pharmacy-Operations/BAP>.

FOR FURTHER INFORMATION CONTACT:

Designated Federal Official (DFO) Colonel Paul J. Hoerner, USAF, 703-681-2890 (voice), dha.ncr.j-6.mbx.baprequests@mail.mil (email). Mailing address is 7700 Arlington Boulevard, Suite 5101, Falls Church, VA 22042-5101. Website: <https://www.health.mil/Military-Health-Topics/Access-Cost-Quality-and-Safety/Pharmacy-Operations/BAP>. The most up-to-date changes to the meeting agenda can be found on the website.

SUPPLEMENTARY INFORMATION: This meeting is being held under the provisions of the Federal Advisory Committee Act (FACA) of 1972 (5 U.S.C., Appendix, as amended), the Government in the Sunshine Act of 1976 (5 U.S.C. 552b, as amended), and 41 CFR 102-3.140 and 102-3.150.

Due to circumstances beyond the control of the Designated Federal Officer, the Uniform Formulary Beneficiary Advisory Panel was unable to provide public notification required by 41 CFR 102-3.150(a) concerning its September 29, 2022 meeting. Accordingly, the Advisory Committee Management Officer for the Department of Defense, pursuant to 41 CFR 102-3.150(b), waives the 15-calendar day notification requirement.

Purpose of the Meeting: The Panel will review and comment on recommendations made to the Director, Defense Health Agency, by the Pharmacy and Therapeutics Committee, regarding the Uniform Formulary.

Agenda:

1. 10:00 a.m.–10:10 a.m. Sign In for UF BAP members.
2. 10:10 a.m.–10:40 a.m. Welcome and Opening Remarks.
 - a. Welcome, Opening Remarks, and Introduction of UF BAP Members by Col Paul J. Hoerner, DFO, UF BAP.
 - b. Opening Remarks by UF BAP Co-Chair Senior Chief Petty Officer Jon R. Ostrowski, Non-Commissioned Officers Association.
 - c. Introductory Remarks by Dr Edward Vonberg, Chief, Formulary Management Branch.
 - d. Public Written Comments by Dr Edward Vonberg, Chief, Formulary Management Branch.
3. 10:40 a.m.–11:45 a.m. Scheduled Therapeutic Class Reviews.
 - a. Overactive Bladder Agents—Beta 3 Adrenergic Agonists Subclass.

¹ 17 CFR 145.9.