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Lead, Information Collection Review Office,  
Office of Public Health Ethics and  
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Disease Control and Prevention.

[FR Doc. 2024-05386 Filed 3-13-24; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Notice of Closed Meeting

Pursuant to 5 U.S.C. 1009(d), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92-463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—PAR 18-812, NIOSH Member Conflict Review.

*Date:* June 13, 2024.

*Time:* 1 p.m.–4 p.m., EDT.

*Place:* Teleconference.

*Agenda:* To review and evaluate grant applications.

*For Further Information Contact:* Michael Goldcamp, Ph.D., Scientific Review Officer, Office of Extramural Programs, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 1095 Willowdale Road, Morgantown, West Virginia 26505. Telephone: (304) 285-5951; Email: [MGoldcamp@cdc.gov](mailto:MGoldcamp@cdc.gov).

The Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Office of Strategic Business  
Initiatives, Office of the Chief Operating  
Officer, Centers for Disease Control and  
Prevention.

[FR Doc. 2024-05458 Filed 3-13-24; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2023-N-5323]

#### Hoffmann-La Roche, Inc., et al.; Withdrawal of Approval of Two New Drug Applications; Correction

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; correction.

**SUMMARY:** The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** on December 18, 2023. The document announced the withdrawal of approval of two new drug applications (NDA), withdrawn as of January 17, 2024. The document indicated that FDA was withdrawing approval of NDA 022424, FLOWTUSS (guaifenesin 200 milligrams (mg)/5 milliliters (mL) and hydrocodone bitartrate 2.5 mg/5 mL) Oral Solution held by Chartwell RX Sciences, LLC, 77 Brenner Dr., Congers, NY 10920. Before FDA withdrew the approval of this NDA, Chartwell RX Sciences, LLC informed FDA that it did not want the approval of the NDA withdrawn. Because Chartwell RX Sciences, LLC timely requested that approval of NDA 022424 not be withdrawn, the approval is still in effect. This notice corrects that error.

**FOR FURTHER INFORMATION CONTACT:** Kimberly Lehrfeld, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6226, Silver Spring, MD 20993-0002, 301-796-3137, [Kimberly.Lehrfeld@fda.hhs.gov](mailto:Kimberly.Lehrfeld@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of Monday, December 18, 2023 (88 FR 87433), appearing on page 87433 in FR Doc. 2023-27661, the following correction is made:

On page 87433, in the table, the entry for NDA 022424 is removed.

Dated: March 11, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024-05426 Filed 3-13-24; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket Nos. FDA-2023-E-1620 and FDA-2023-E-1597]

#### Determination of Regulatory Review Period for Purposes of Patent Extension; PYRUKYND

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or the Agency) has determined the regulatory review period for PYRUKYND and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of applications to the Director of the U.S. Patent and Trademark Office (USPTO), Department of Commerce, for the extension of patents which claim that human drug product.

**DATES:** Anyone with knowledge that any of the dates as published (see **SUPPLEMENTARY INFORMATION**) are incorrect must submit either electronic or written comments and ask for a redetermination by May 13, 2024. Furthermore, any interested person may petition FDA for a determination regarding whether the applicant for extension acted with due diligence during the regulatory review period by September 10, 2024. See “Petitions” in the **SUPPLEMENTARY INFORMATION** section for more information.

**ADDRESSES:** You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of May 13, 2024. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

#### Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any