

will receive written comments relating to the settlement. The Agency will consider all comments received and may modify or withdraw its consent to the settlement if comments received disclose facts or considerations which indicate that the settlement is inappropriate, improper, or inadequate. The proposed settlement is available for public inspection at <https://www.regulations.gov>. The Agency's response to any comments received will be available for public inspection at the EPA, Region 5, Records Center, 77 W. Jackson Blvd., 7th Fl., Chicago, Illinois 60604. Commenters may request an opportunity for a public hearing in the affected area, in accordance with Section 7003(d) of RCRA.

DATES: Comments must be submitted on or before August 18, 2023.

ADDRESSES: You may send comments, identified by Docket ID No. EPA-R05-INSERT, by any of the following methods:

- **Federal eRulemaking Portal:** <https://www.regulations.gov/> (our preferred method). Follow the online instructions for submitting comments.
- **Mail:** U.S. Environmental Protection Agency, ATTN: Mark Koller, Associate Regional Counsel, Office of Regional Counsel (C-14J), 77 W. Jackson Blvd., Chicago, Illinois 60604.

Instructions: All submissions received must include the Docket ID No. for this rulemaking. Comments received may be posted without change to <https://www.regulations.gov/>, including any personal information provided. For detailed instructions on sending comments and additional information on the rulemaking process, see the "Public Participation" heading of the **SUPPLEMENTARY INFORMATION** section of this document.

FOR FURTHER INFORMATION CONTACT: Mark Koller, Office of Regional Counsel, Environmental Protection Agency, telephone number: (312) 353-2591; email address: koller.mark@epa.gov.

SUPPLEMENTARY INFORMATION:

I. Public Participation

A. Written Comments

Submit your comments, identified by Docket ID No. EPA-R05-INSERT, at <https://www.regulations.gov> (our preferred method), or the other methods identified in the **ADDRESSES** section. Once submitted, comments cannot be edited or removed from the docket. The EPA may publish any comment received to its public docket. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information

whose disclosure is restricted by statute. Multimedia submissions (audio, video, etc.) must be accompanied by a written comment. The written comment is considered the official comment and should include discussion of all points you wish to make. The EPA will generally not consider comments or comment contents located outside of the primary submission (*i.e.* on the web, cloud, or other file sharing system). For additional submission methods, the full EPA public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit <https://www.epa.gov/dockets/commenting-epa-dockets>.

II. Background Information

The Settling Party proposes to acquire ownership of a portion of the former General Motors Corporation North American operation, at 2915 Doctor Martin Luther King Junior Boulevard. The Site is one of the 89 sites that were placed into an Environmental Response Trust (the "Trust") as a result of the resolution of the 2009 GM bankruptcy. The Trust is administrated by Revitalizing Auto Communities Environmental Response.

Douglas Ballotti,

Director, Superfund & Emergency Management Division, Region 5.

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ENVIRONMENTAL PROTECTION AGENCY

[FRL-10978-01-R3]

Clean Water Act: Identification of Water Quality-Limited Segments To Be Added to West Virginia's Section 303(d) List

AGENCY: Environmental Protection Agency (EPA).

ACTION: Public notice.

SUMMARY: The Clean Water Act (CWA) requires that states periodically submit, and the Environmental Protection Agency (EPA) approve or disapprove, lists of waters (called "Section 303(d) lists") for which existing technology-based pollution controls are not stringent enough to attain or maintain State water quality standards and for which total maximum daily loads (TMDLs) must be prepared. Waters identified on Section 303(d) lists are called "water quality-limited segments." This notice announces the EPA's identification of certain additional water quality-limited segments for West

Virginia's Combined 2018-2020-2022 Section 303(d) list and requests public comment on those additions.

DATES: Comments must be received on or before August 18, 2023.

ADDRESSES: You may send written comments to Mr. Gregory Voigt by the following methods:

- **Electronic mail:** voigt.gregory@epa.gov. Include 'FRL-10978-01-R3 comment' in the subject line of the message.
- **Mail:** Mr. Gregory Voigt, Mail Code 3WD42, U.S. Environmental Protection Agency Region 3 Water Division, Four Penn Center, 1600 John F. Kennedy Blvd., Philadelphia, PA 19103-2029.

FOR FURTHER INFORMATION CONTACT: Mr. Gregory Voigt, Water Division Standards and TMDL Section, 3WD42, Environmental Protection Agency at voigt.gregory@epa.gov or (215) 814-5737. Additional information regarding the basis for this EPA action is available at <https://www.epa.gov/tmdl/wv-303d-list-public-notice>.

SUPPLEMENTARY INFORMATION: Section 303(d) of the Clean Water Act requires that each state identify those water quality-limited segments for which existing technology-based pollution controls are not stringent enough to attain or maintain state water quality standards and for which total maximum daily loads (TMDLs) must be prepared. For each water quality-limited segment on the list, the state identifies the pollutant causing the impairment, when known. In addition, the state assigns a priority ranking for development of TMDLs based on the severity of the pollution and the uses to be made of the waters, among other factors (40 CFR 130.7(b)(4)).

The EPA's Water Quality Planning and Management regulations include requirements related to the implementation of Section 303(d) of the CWA (40 CFR 130.7). The regulations require states to assemble and evaluate all existing and readily-available water quality data and to use that data to identify water quality-limited segments still requiring TMDLs every two years. Where a state does not use certain data, it must provide a rationale. The list of waters still needing TMDL development must also include priority rankings and must identify the waters targeted for TMDL development during the next two years (40 CFR 130.7).

The EPA received West Virginia's submittal of its listing decisions under Section 303(d)(2) on May 5, 2023. On June 1, 2023, the EPA partially approved and partially disapproved West Virginia's Combined 2018-2020-2022 Section 303(d) list of water

quality-limited segments based upon the Agency's finding that West Virginia did not use certain water quality information and therefore did not identify certain water quality-limited segments based upon existing data and public input. The EPA analyzed the information and identified three-hundred forty-six (346) additional water quality-limited segments for inclusion on West Virginia's Combined 2018–2020–2022 Section 303(d) list. These water quality-limited segments are identified in Enclosure 3 of the June 1, 2023 decision document and displayed on an interactive webmap, both of which are available at the website link provided.

The EPA is providing the public an opportunity to review and comment on its identification of these water quality-limited segments for West Virginia's Combined 2018–2020–2022 Section 303(d) list as required by 40 CFR 130.7(d)(2). The EPA will consider public comments and make any appropriate revisions before transmitting its final list of water quality-limited segments to the State.

Catherine Libertz,

Director, Water Division, Region III.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPPT–2023–0061; FRL–10581–06–OCSPP]

Certain New Chemicals; Receipt and Status Information for June 2023

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA is required under the Toxic Substances Control Act (TSCA), as amended by the Frank R. Lautenberg Chemical Safety for the 21st Century Act, to make information publicly available and to publish information in the **Federal Register** pertaining to submissions under TSCA Section 5, including notice of receipt of a Premanufacture notice (PMN), Significant New Use Notice (SNUN) or Microbial Commercial Activity Notice (MCAN), including an amended notice or test information; an exemption application (Biotech exemption); an application for a test marketing exemption (TME), both pending and/or concluded; a notice of commencement (NOC) of manufacture (including import) for new chemical substances; and a periodic status report on new

chemical substances that are currently under EPA review or have recently concluded review. This document covers the period from 6/1/2023 to 6/30/2023.

DATES: Comments identified by the specific case number provided in this document must be received on or before August 18, 2023.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPPT–2023–0061, through the *Federal eRulemaking Portal* at <https://www.regulations.gov>. Follow the online instructions for submitting comments. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Additional instructions on commenting and visiting the docket, along with more information about dockets generally, is available at <https://www.epa.gov/dockets>.

FOR FURTHER INFORMATION CONTACT:

For technical information contact: Jim Rahai, Project Management and Operations Division (MC 7407M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 564–8593; email address: rahai.jim@epa.gov.

For general information contact: The TSCA–Hotline, ABVI–Goodwill, 422 South Clinton Ave., Rochester, NY 14620; telephone number: (202) 554–1404; email address: TSCA-Hotline@epa.gov.

SUPPLEMENTARY INFORMATION:

I. Executive Summary

A. What action is the Agency taking?

This document provides the receipt and status reports for the period from 6/01/2023 to 6/30/2023. The Agency is providing notice of receipt of PMNs, SNUNs, and MCANs (including amended notices and test information); an exemption application under 40 CFR part 725 (Biotech exemption); TMEs, both pending and/or concluded; NOCs to manufacture a new chemical substance; and a periodic status report on new chemical substances that are currently under EPA review or have recently concluded review.

EPA is also providing information on its website about cases reviewed under the amended TSCA, including the section 5 PMN/SNUN/MCAN and exemption notices received, the date of receipt, the final EPA determination on the notice, and the effective date of EPA's determination for PMN/SNUN/MCAN notices on its website at: [https://](https://www.epa.gov/reviewing-new-chemicals-under-toxic-substances-control-act-tsca/status-pre-manufacture-notices)

www.epa.gov/reviewing-new-chemicals-under-toxic-substances-control-act-tsca/status-pre-manufacture-notices. This information is updated on a weekly basis.

B. What is the Agency's authority for taking this action?

Under the Toxic Substances Control Act (TSCA), 15 U.S.C. 2601 *et seq.*, a chemical substance may be either an “existing” chemical substance or a “new” chemical substance. Any chemical substance that is not on EPA's TSCA Inventory of Chemical Substances (TSCA Inventory) is classified as a “new chemical substance,” while a chemical substance that is listed on the TSCA Inventory is classified as an “existing chemical substance.” (See TSCA section 3(11).) For more information about the TSCA Inventory please go to: <https://www.epa.gov/tsca-inventory>.

Any person who intends to manufacture (including import) a new chemical substance for a non-exempt commercial purpose, or to manufacture or process a chemical substance in a non-exempt manner for a use that EPA has determined is a significant new use, is required by TSCA section 5 to provide EPA with a PMN, MCAN, or SNUN, as appropriate, before initiating the activity. EPA will review the notice, make a risk determination on the chemical substance or significant new use, and take appropriate action as described in TSCA section 5(a)(3).

TSCA section 5(h)(1) authorizes EPA to allow persons, upon application and under appropriate restrictions, to manufacture or process a new chemical substance, or a chemical substance subject to a significant new use rule (SNUR) issued under TSCA section 5(a)(2), for “test marketing” purposes, upon a showing that the manufacture, processing, distribution in commerce, use, and disposal of the chemical will not present an unreasonable risk of injury to health or the environment. This is referred to as a test marketing exemption, or TME. For more information about the requirements applicable to a new chemical go to: <https://www.epa.gov/chemicals-under-tsca>.

Under TSCA sections 5 and 8 and EPA regulations, EPA is required to publish in the **Federal Register** certain information, including notice of receipt of a PMN/SNUN/MCAN (including amended notices and test information); an exemption application under 40 CFR part 725 (biotech exemption); an application for a TME, both pending and concluded; NOCs to manufacture a new chemical substance; and a periodic status report on the new chemical