

# Minnesota Pollution Control Agency

## Resource Management and Assistance Division

### REQUEST FOR COMMENTS

#### Planned New Rules Governing Reporting by Manufacturers Upon Submission of Required Information about Products Containing Per-and polyfluoroalkyl substances (PFAS), Revisor's ID Number R-4828

**NOTICE IS HEREBY GIVEN** that the Minnesota Pollution Control Agency (MPCA) is requesting comments on planned new rules for submission of required information about products containing PFAS. This rulemaking is referred to as the **PFAS in Products Reporting Rule**. The main purpose of this rulemaking is to establish a program for the MPCA to collect information about products containing PFAS intentionally added to products sold, offered for sale, or distributed in Minnesota as required by [Minnesota Session Law – 2023, chapter 60, article 3, section 21, \(Minnesota Statutes 116.943\) subdivision 2](#). Comments are requested from affected or interested parties. Comments should be submitted in writing as described in the [Comments](#) section below.

This Request for Comments is the MPCA's legal notice of its intent to begin rulemaking. This is an opportunity to provide information or comment on any relevant issues related to this rulemaking that we need to consider. For example, we recognize that the cost of compliance for regulated parties can be a concern. If you have cost information or data related to collecting information about PFAS in products reporting that you wish to share with us to inform our decisions, please submit that information. Draft rule language is not available at this time. We want your written comments on the ideas described under the [Subject of Rules](#) section.

Submitting your ideas and information at this early stage in rulemaking allows us more time to address issues that may come up and helps to ensure informed decision-making on our part. If the planned rules affect you in any way, the MPCA encourages you to participate in the rulemaking process.

**Alternative Format/Accommodation.** Upon request, this information can be made available in an alternative format, such as large print, braille, or audio. To make such a request, please contact the [MPCA Contact Person](#).

**Statutory Authority.** Minnesota Session Law – 2023, chapter 60, article 3, section 21, ([Minnesota Statutes 116.943](#)); and [Minnesota Statutes, section 116.943, subdivision 9](#).

**Subject of Rules.** The MPCA requests comments on planned new rules governing PFAS in products reporting. Information must be submitted by “manufacturers” of a product containing intentionally added PFAS about the product, the PFAS, and the submitter, as authorized by Minnesota Session Law – 2023, chapter 60, article 3, section 21, subdivision 2, which states:

“(a) On or before January 1, 2026, a manufacturer of a product sold, offered for sale, or distributed in the state that contains intentionally added PFAS must submit to the commissioner information that includes:

- (1) a brief description of the product, including a universal product code (UPC), stock keeping unit (SKU), or other numeric code assigned to the product;
- (2) the purpose for which PFAS are used in the product, including in any product components;

(3) the amount of each PFAS, identified by its chemical abstracts service registry number, in the product, reported as an exact quantity determined using commercially available analytical methods or as falling within a range approved for reporting purposes by the commissioner;

(4) the name and address of the manufacturer and the name, address, and phone number of a contact person for the manufacturer; and

(5) any additional information requested by the commissioner as necessary to implement the requirements of this section.

(b) With the approval of the commissioner, a manufacturer may supply the information required in paragraph (a) for a category or type of product rather than for each individual product.

(c) A manufacturer must submit the information required under this subdivision whenever a new product that contains intentionally added PFAS is sold, offered for sale, or distributed in the state and update and revise the information whenever there is significant change in the information or when requested to do so by the commissioner.

(d) A person may not sell, offer for sale, or distribute for sale in the state a product containing intentionally added PFAS if the manufacturer has failed to provide the information required under this subdivision and the person has received notification under subdivision 4.”

In addition, other subdivisions in the law may directly relate to or have implications for reporting, such as the subdivisions referring to definitions, waivers and extensions, and exemptions.

In developing the reporting rule, the MPCA would appreciate comments on the following questions:

- 1) Are there definitions in subdivision 1 for which clarification would be useful to understanding reporting responsibilities?
- 2) Are there terms or processes in subdivision 2 for which clarifications will help reporting entities determine reporting status or data-gathering process?
- 3) How should the MPCA balance public availability of data and trade secrecy as part of the reporting requirements?
- 4) Are there any terms used in subdivision 3 that should be further defined or where examples would be helpful?
- 5) Are there specific portions of the reporting process that should not be defined through guidance or the development of an application form?
- 6) Other questions or comments relating to reporting or the process of reporting.

The MPCA is interpreting these information submittals (“reporting”) to occur once, on or before January 1, 2026, and to not involve resubmittal of the same information. However, updates to reported products when PFAS are added or subtracted (“whenever there is a significant change”) are required.

**Parties Affected.** The new rule would affect any manufacturer of a product sold, offered for sale, or distributed in the state that contains intentionally added PFAS. Definitions of pertinent terms are provided in Minnesota Session Law – 2023, chapter 60, article 3, section 21, subdivision 1, including:

- "Manufacturer" means the person that creates or produces a product or whose brand name is affixed to the product. In the case of a product imported into the United States, manufacturer includes the importer or first domestic distributor of the product if the person that manufactured or assembled the product or whose brand name is affixed to the product does not have a presence in the United States.

- "Intentionally added" means PFAS deliberately added during the manufacture of a product where the continued presence of PFAS is desired in the final product or one of the product's components to perform a specific function.
- "Perfluoroalkyl and polyfluoroalkyl substances" or "PFAS" means a class of fluorinated organic chemicals containing at least one fully fluorinated carbon atom.

"Manufacturers" can be located in Minnesota or anywhere outside the State, as long as they are selling a product, offering a product for sale, or distributing a product (or component) in the state that contains intentionally added PFAS.

Note that in some cases "manufacturer" may include people or entities not typically thought of as manufacturers, such as retailers which have their brand name or a private label brand name they own affixed to products. "Manufacturer" does not include landfill or wastewater treatment operators or any person who sells, offers for sale, or distributes in Minnesota:

- Products for which federal law governs the presence of PFAS in the product in a manner that preempts state authority;
- A product regulated under section 325F.072 or 325F.075;
- A used product; or
- Products which contain a pesticidal ingredient regulated by and reported to the Minnesota Department of Agriculture.

While subdivision 8 of the law exempts them from prohibitions and testing and certificate of compliance requirements, "manufacturers" of prosthetic or orthotic devices or any products that are medical devices or drugs or that are otherwise used in a medical setting or in medical applications regulated by the United States Food and Drug Administration are not exempted from information submittal requirements.

**Where to Get More Information.** Information about this rulemaking is available on the rulemaking website at <https://www.pca.state.mn.us/get-engaged/pfas-in-products-reporting>. As stated above, we do not yet have draft rule language. If you are interested in being notified of opportunities for public comment, when the draft rules are available for review, and of other activities relating to this (or other MPCA rulemakings) register for GovDelivery bulletins at <https://public.govdelivery.com/accounts/MNPCA/subscriber/new>.

**Comments.** Interested parties may submit written comments or information on these possible rules until **4:30 p.m. on Tuesday, November 28, 2023**. Submit written comments or information to the Office of Administrative Hearings (OAH) Rulemaking eComments website at <https://minnesotaoah.granicusideas.com/>. Any questions about submitting comments via the Rulemaking eComments website should be directed to William Moore, OAH, telephone 651-361-7893, email [William.T.Moore@state.mn.us](mailto:William.T.Moore@state.mn.us). You may view frequently asked questions about the OAH eComments website at [https://mn.gov/oah/assets/ecommments-faq\\_tcm19-82012.pdf](https://mn.gov/oah/assets/ecommments-faq_tcm19-82012.pdf). Comments received are public and will be available for review at the OAH Rulemaking eComments website at <https://minnesotaoah.granicusideas.com/discussions> and at the OAH, 600 North Robert Street, P.O. Box 64620, St. Paul, Minnesota 55164-0620.

The MPCA will not publish a Notice of Intent to Adopt the rules until more than 60 days have elapsed from the date of this RFC.

The MPCA does not anticipate that the rule amendments will require a local government to adopt or amend an ordinance or other regulation under *Minnesota Statutes*, section 14.128. Local governments may submit written information to the contrary.

The MPCA requests any information pertaining to the cumulative effect of the rule amendments with other federal and state regulations related to the specific purpose of the rule. Cumulative effect means the impact that results from incremental impact of the proposed rule in addition to other rules, regardless of what state or federal agency has adopted the other rules.

NOTE: The MPCA will carefully consider all comments received in response to this RFC. However, these comments will not necessarily be included in the formal rulemaking record submitted to the Administrative Law Judge (ALJ) if and when a proceeding to adopt rules is started. The MPCA is required to submit to the ALJ only the written comments received in response to the draft rules after they are proposed with a Notice of Intent to Adopt Rules. If you submit comments during the RFC stage of rule development and want to ensure that the ALJ reviews them, you should resubmit your comments after the rules are formally proposed with a Notice of Intent to Adopt the rules.

**MPCA Contact Person.** The MPCA contact person is Mary H. Lynn at the MPCA, 520 Lafayette Road North, St. Paul, Minnesota 55155-4194; telephone 651-757-2439, email [mary.lynn@state.mn.us](mailto:mary.lynn@state.mn.us). Technical questions on the planned rules should be submitted to Al Innes, telephone 651-757-2457, email [alister.innes@state.mn.us](mailto:alister.innes@state.mn.us). You may also call the MPCA at 651-296-6300 or 1-800-657-3864; use your preferred relay service.



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Katrina Kessler, Commissioner  
Minnesota Pollution Control Agency

September 11, 2023  
Date