TABLE 1 TO PARAGRAPH (f)(1)—FACILITY-WIDE 30-DAY ROLLING AVERAGE NO_X EMISSION RATE LIMITS—Continued

Facility	Facility-wide 30-day rolling average NO_X emission rate limit (lb/MMBtu)
Homer City	0.088
Keystone	0.074
Montour	0.069

(2) The Facility shall achieve and maintain their Unit-specific Daily NO_X Mass Emissions to not exceed the Unit-specific limit in Table 2 to this paragraph (f)(2).

TABLE 2 TO PARAGRAPH (f)(2)—UNIT-SPECIFIC DAILY NO_X MASS EMISSIONS LIMITS

Facility	Unit	Unit-specific daily NO_X Mass emissions limit (lb/day)
Cheswick	1	14,256
Conemaugh	1	18,084
Conemaugh	2	18,084
Homer City	1	14,345
Homer City	2	14,345
Homer City	3	15,333
Keystone	1	15,481
Keystone	2	15,481
Montour	1	12,117
Montour	2	11,988

(g) *Monitoring of NO_X emissions.* (1) In determining the Facility-wide 30-Day Rolling Average NO_X Emission Rate, the Facility shall use CEMS in accordance with the procedures of 40 CFR part 60 and 40 CFR part 75, appendix F, Procedure 1.

(2) For purposes of calculating the Unit-specific Daily NO_X Mass Emissions Limits, the Facility shall use CEMS in accordance with the procedures at 40 CFR part 75. Emissions rates, mass emissions, and other quantitative standards set by or under this section must be met to the number of significant digits in which the standard or limit is expressed. For example, an emission rate of 0.100 is not met if the actual emission rate is 0.101. The Facility shall round the fourth significant digit to the nearest third significant digit, or the sixth significant digit to the nearest fifth significant digit, depending upon whether the limit is expressed to three or five significant digits. For example, if an actual emission rate is 0.1004, that shall be reported as 0.100, and shall be in compliance with an emission rate of 0.100, and if an actual emission rate is 0.1005, that shall be reported as 0.101, and shall not be in compliance with an emission rate of 0.100. The Facility shall report data to the number of significant digits in which the standard or limit is expressed.

(h) *Recordkeeping and periodic reporting.* (1) The Facility shall electronically submit to EPA a periodic report, within thirty (30) days after the end of each six-month reporting period (January through June, July through December in each calendar year). The portion of the periodic report containing the data required to be reported by this paragraph (h) shall be in an unlocked electronic spreadsheet format, such as Excel or other widely-used software, and contain data for each Operating Day during the reporting period, including, but not limited to: Facility ID (ORISPL); Facility name; Unit ID; Date; Unit-specific total Daily Operating Time (hours); Unit-specific Daily NO_X Mass Emissions (lbs); Unit-specific total Daily Heat Input (MMBtu); Unit-specific Daily NO_X Emission Rate (lb/MMBtu); Facility-wide 30-Day Rolling Average NO_X Emission Rate (lb/MMBtu); Owner; Operator; Representative (Primary); and Representative (Secondary). In addition, the Facility shall maintain the following information for 5 years from the date of creation of the data and make such information available to EPA if requested: Unit-specific hourly heat input, Unit-specific hourly ammonia injection amounts, and Unit-specific hourly NO_X emission rate.

(2) In any periodic report submitted pursuant to this section, the Facility may incorporate by reference information previously submitted to EPA under its Title V permitting requirements in this chapter, so long as that information is adequate to determine compliance with the emission limits and in the same electronic format as required for the periodic report, and provided that the Facility attaches the Title V Permit report (or the pertinent portions of such report) and provides a specific reference to the provisions of the Title V Permit report that are responsive to the information required in the periodic report.

(3) In addition to the reports required pursuant to this section, if the Facility exceeds the Facility-wide 30-day rolling average NO_X emission limit on three or more days during any 30-day period, or exceeds the Unit-specific daily mass emission limit for any Unit on three or more days during any 30-day period, the Facility shall electronically submit to EPA a report on the exceedances within ten (10) business days after the Facility knew or should have known of the event. In the report, the Facility shall explain the cause or causes of the exceedances and any measures taken or to be taken to cure the reported exceedances or to prevent such exceedances in the future. If at any time, the provisions of this section are included in Title V Permits, consistent with the requirements for such inclusion in this section, then the deviation reports required under applicable Title V regulations in this chapter shall be deemed to satisfy all the requirements of this paragraph (h)(3).

(4) Each report shall be signed by the Responsible Official as defined in Title V of the Clean Air Act, or his or her equivalent or designee of at least the rank of Vice President. The signatory shall also electronically submit the following certification, which may be contained in a separate document:

> This information was prepared either by me or under my direction or supervision in accordance with a system designed to assure that qualified personnel properly gather and evaluate the information submitted. Based on my evaluation, or the direction and my inquiry of the person(s) who manage the system, or the person(s) directly responsible for gathering the information, I hereby certify under penalty of law that, to the best of my knowledge and belief, this information is true, accurate, and complete. I understand that there are significant penalties for submitting false, inaccurate, or incomplete information to the United States.

(5) Whenever notifications, submissions, or communications are required by this section, they shall be made electronically to the attention of the Air Enforcement Manager via email to the following address: *R3_ORC_mailbox@epa.gov.*

[FR Doc. 2022–10765 Filed 5–24–22; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 751

[EPA–HQ–OPPT–2021–0057; FRL–8332–03–OCSPP]

RIN 2070–AK86

Asbestos Part 1: Chrysotile Asbestos; Regulation of Certain Conditions of Use Under Section 6(a) of the Toxic Substances Control Act (TSCA); Extension of Comment Period

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule; extension of comment period.

SUMMARY: EPA proposed a rule under the Toxic Substances Control Act

(TSCA) to address the unreasonable risk of injury to health it has identified for conditions of use of chrysotile asbestos following completion of the TSCA Risk Evaluation for Asbestos, Part 1: Chrysotile Asbestos.

DATES: The comment period for the proposed rule published April 12, 2022, 87 FR 21706, is extended. Comments must be received on or before July 13, 2022.

ADDRESSES: Submit your comments, identified by ID number EPA–HQ–OPPT–2021–0057, 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 or visiting the docket, along with more information about dockets generally, is available at *https://www.epa.gov/dockets/about-epa-dockets.*

FOR FURTHER INFORMATION CONTACT: *For technical information contact:* Peter Gimlin, Existing Chemicals Risk Management Division (Mail Code 7404T), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 566–0515; email address: *gimlin.peter@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: This document extends the public comment period established in the **Federal Register** of April 12, 2022 (87 FR 21706) (FRL–8332–02–OCSPP) for 30 days, from June 13, 2022 to July 13, 2022. In that document, EPA proposed a rule under TSCA to address the unreasonable risk of injury to health it has identified for conditions of use of chrysotile asbestos following completion of the TSCA Risk Evaluation for Asbestos, Part 1: Chrysotile Asbestos, and solicited public comment on the proposed rule. More information on EPA's proposed regulation and solicitation of comment can be found in the **Federal Register** of April 12, 2022.

EPA received requests to extend the comment period and believes it is appropriate to do so in order to give stakeholders additional time to review the proposed regulation and prepare comments.

If you have questions, consult the technical person listed under **FOR FURTHER INFORMATION CONTACT**.

List of Subjects in 40 CFR Part 751

Environmental protection, Chemicals, Export certification, Hazardous substances, Import certification, Recordkeeping.

Dated: May 16, 2022.

Michal Freedhoff,

Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

[FR Doc. 2022–10854 Filed 5–24–22; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

42 CFR Part 600

[CMS–2441–P]

RIN 0938–AU89

Basic Health Program; Federal Funding Methodology for Program Year 2023 and Proposed Changes to Basic Health Program Regulations

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Proposed rule.

SUMMARY: This document proposes the methodology and data sources necessary to determine Federal payment amounts to be made for program year 2023 to States that elect to establish a Basic Health Program under the Patient Protection and Affordable Care Act to offer health benefits coverage to low-income individuals otherwise eligible to purchase coverage through Health Insurance Exchanges.

DATES: To be assured consideration, comments must be received at one of the addresses provided below, no later than 5 p.m. on June 24, 2022.

ADDRESSES: In commenting, refer to file code CMS–2441–P.

Comments, including mass comment submissions, must be submitted in one of the following three ways (please choose only one of the ways listed):

1. *Electronically.* You may submit electronic comments on this regulation to *http://www.regulations.gov.* Follow the ''Submit a comment'' instructions.

2. *By regular mail.* You may mail written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–2441–P, P.O. Box 8016, Baltimore, MD 21244–8016.

Please allow sufficient time for mailed comments to be received before the close of the comment period.

3. *By express or overnight mail.* You may send written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–2441–P, Mail Stop C4–26–05, 7500 Security Boulevard, Baltimore, MD 21244–1850.

For information on viewing public comments, see the beginning of the **SUPPLEMENTARY INFORMATION** section.

FOR FURTHER INFORMATION CONTACT: Christopher Truffer, (410) 786–1264; or Cassandra Lagorio, (410) 786–4554.

SUPPLEMENTARY INFORMATION:

Inspection of Public Comments: All comments received before the close of the comment period are available for viewing by the public, including any personally identifiable or confidential business information that is included in a comment. We post all comments received before the close of the comment period on the following website as soon as possible after they have been received: *http://www.regulations.gov.* Follow the search instructions on that website to view public comments. CMS will not post on *Regulations.gov* public comments that make threats to individuals or institutions or suggest that the commenter will take actions to harm another individual. CMS continues to encourage individuals not to submit duplicative comments. We will post acceptable comments from multiple unique commenters even if the content is identical or nearly identical to other comments.

I. Background

A. Overview of the Basic Health Program

Section 1331 of the Patient Protection and Affordable Care Act (Pub. L. 111–148, enacted on March 23, 2010), as amended by the Health Care and Education Reconciliation Act of 2010 (Pub. L. 111–152, enacted on March 30, 2010) (collectively referred to as the Affordable Care Act or ACA) provides States with an option to establish a Basic Health Program (BHP). In the States that elect to operate a BHP, the BHP makes affordable health benefits coverage available for individuals under age 65 with household incomes between 133 percent and 200 percent of the Federal poverty level (FPL) who are not otherwise eligible for Medicaid, the Children's Health Insurance Program (CHIP), or affordable employer-sponsored coverage, or for individuals whose income is below these levels but