

Please refer to the **SUPPLEMENTARY INFORMATION** section for additional information on the public meetings.

ADDRESSES: You may send comments, by any of the following methods:

- *Email:* Region2_MPRSA@epa.gov.

Include “NEPA” in the subject line of the message.

FOR FURTHER INFORMATION CONTACT:

Mark Reiss, EPA Region 2 Water Division, Dredging, Sediments & Oceans Section, 290 Broadway, New York, NY 10007, phone: 212-637-3799, email: Region2_MPRSA@epa.gov. Additional information is available on the EPA website at <https://www.epa.gov/marine-protection-permitting/region-2-ocean-dredged-material-management-new-york-bight-atlantic>.

SUPPLEMENTARY INFORMATION: The EPA intends to prepare an environmental assessment document consistent with NEPA to inform its options for dredged material management in the New York Bight. The environmental review document will provide the information necessary to evaluate potential adverse impacts on the environment and other ocean uses and to identify a preferred alternative that meets the EPA’s site selection criteria at 40 CFR 228.5 and 228.6.

Need for Action: Dredging is essential for maintaining safe navigation in ports and harbors in and around New York and New Jersey. Since 1997, 88 million cubic yards of dredged material from the New York/New Jersey Harbor region that meets MPRSA sediment quality standards has been placed as “remediation material” to improve ecological conditions on the seabed at the Historic Area Remediation Site (HARS), located 3.3 miles east of Highlands, New Jersey, and 7.7 miles south of Rockaway, New York. Prior to that time, the HARS bottom was adversely affected by legacy pollutants accumulated over an extended period of American history. The HARS is nearing its remediation goal of placement of one meter of remediation dredged material over the site.

Considering the HARS’s status, the U.S. Army Corps of Engineers (Army Corps) New York District compared the volume of dredged material estimated to be generated over the next 20 years to the capacity of all the nearby existing sites used to manage NY/NJ Harbor dredged material. As a result of the comparison, the Army Corps identified a shortfall in dredged material management capacity. Accordingly, the Army Corps has requested that the EPA evaluate alternatives for meeting future needs for managing HARS-suitable dredged material in the New York Bight

and to initiate any site designation or modification actions necessary.

Alternatives: Alternatives that the EPA may consider include: the continued placement of dredged material meeting current remediation standards in portions of the HARS; designation of one or more additional sites for beneficial use of HARS-suitable dredged material to remediate other areas of New York Bight seafloor adversely affected by historical contamination on the seafloor; and options for use of certain types of HARS-suitable materials to create or enhance seafloor habitat.

Scoping: This NOI commences the public scoping process for the project. The EPA is soliciting input from federal, state, and local governments, industry, non-governmental organizations, and the public on the range of alternatives considered, specific environmental issues to be evaluated in the NEPA document, and the potential impacts of alternatives for managing HARS-suitable dredged material in the New York Bight. Registration information for virtual scoping meetings regarding this project is available at the EPA website listed above. Comments received at the virtual scoping meetings will be considered as part of the record. Comments can be submitted via email to: Region2_MPRSA@epa.gov. Comments will be accepted for a period of 30 days following the date of this notice.

Estimated Date of Draft NEPA Document: September 2025.

Dated: May 29, 2025.

Michael Martucci,

Regional Administrator, EPA Region 2.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-25-24EG]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Documenting Outcomes Associated with Persistent Tic Disorders (including Tourette Syndrome) in Children, Adolescents, and Young Adults through Surveillance” to the Office of Management and Budget (OMB) for review and approval. CDC previously

published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on April 5, 2024 to obtain comments from the public and affected agencies. CDC did not receive comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Documenting outcomes associated with Persistent Tic Disorders (including Tourette Syndrome) in Children, Adolescents, and Young Adults through Surveillance—New—National Center on Birth Defects and Developmental

Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

There are an estimated 1.4 million people in the U.S. affected by persistent tic disorders (PTD) or Tourette syndrome (TS). To support people with these conditions, the impact of PTD/TS must be understood. Although some data on the impact of PTD/TS on social relationships and education are available, other potential outcomes associated with PTD/TS have not been well-documented, including associated costs, suicidality, health care transition, and the prevalence of co-occurring disorders and how co-occurring disorders modify these outcomes. Limited data are available on how these outcomes may differ among sub-populations (e.g., by sex, race/ethnicity,

age group, and geography [e.g., urban/rural]). This data collection aims to document priority outcomes including costs (e.g., education level, employment, healthcare beyond those available in claims data), prevalence of suicidality risk, transition to adult healthcare, and the prevalence of co-occurring conditions and how they modify these outcomes among children and adolescents (4–17 years) and young adults (18–26 years) with PTD/TS. Data will be collected once from a participant (i.e., individuals with PTD/TS and/or their caregiver), via a survey, and a clinical assessment of tic symptoms. We will also extract data from medical records. Most questions for the survey created for this surveillance project were selected from national surveys or previously validated measures. This will allow us to compare estimates from

this project to external prevalence estimates for the same health indicators in U.S. children, adolescents, and young adults in the general population and to previously published findings. Data will be used to inform where resources for families and healthcare providers (e.g., professional trainings) are most needed to support people with PTD/TS and their families and to address differences in health among subgroups of the population. As a result of working with awardees to finalize measures, and decisions to rely on parent-report for the majority of indicators for this age group, CDC has updated the burden estimates for this data collection. CDC requests OMB approval for an estimated 500 annual burden hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Parents of children 4–17 years with a persistent tic disorder.	Parent	300	1	45/60
Children 4–8 years with a persistent tic disorder	Child 4–8	60	1	20/60
Children 9–11 years with a persistent tic disorder	Child 9–11	100	1	30/60
Adolescents (teens) 12–17 years with a persistent tic disorder.	Adolescent	140	1	45/60
Adults (18–26 years) with a persistent tic disorder	Adult	100	1	1

Jeffrey M. Zirger,
Lead, Information Collection Review Office,
Office of Public Health Ethics and
Regulations, Office of Science, Centers for
Disease Control and Prevention.
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DEPARTMENT OF HEALTH AND
HUMAN SERVICES

Centers for Disease Control and
Prevention

[30Day–25–24EE]

Agency Forms Undergoing Paperwork
Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Division of Foodborne, Waterborne, and Environmental Diseases (DFWED) National Hypothesis Generation and Investigation Module” to the Office of Management and Budget (OMB) for review and approval. CDC previously

published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on April 5, 2024 to obtain comments from the public and affected agencies. CDC received two comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments. CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) Minimize the burden of the collection of information on those who

are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.