

## ESTIMATED ANNUALIZED BURDEN HOURS

| Type of survey                                                               | Respondent                                               | Number of respondents | Number of responses/ respondent | Average burden/ response (in hours) | Response burden (in hours) |
|------------------------------------------------------------------------------|----------------------------------------------------------|-----------------------|---------------------------------|-------------------------------------|----------------------------|
| Questionnaires/Applications .....                                            | Student/Youth .....                                      | 800                   | 1                               | 30/60                               | 400                        |
| Applicants Questionnaire/Application .....                                   | Parents/Guardians of Applicants .....                    | 800                   | 1                               | 30/60                               | 400                        |
| Applications, Recommendations, and Other applicant-supporting documentation. | School Officials/Community Representatives.              | 1200                  | 1                               | 30/60                               | 600                        |
| Focus Groups .....                                                           | Student/Youth; Parent/Guardian; School Officials; Other. | 50                    | 1                               | 60/60                               | 50                         |
| Other Program Surveys .....                                                  | Student/Youth; Parent/Guardian; School Officials; Other. | 600                   | 1                               | 30/60                               | 300                        |
| Total .....                                                                  | .....                                                    | .....                 | .....                           | .....                               | 1,750                      |

**Jeffrey M. Zirger,**

*Lead, Information Collection Review Office,  
Office of Scientific Integrity, 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

[60Day-22-22IV; Docket No. CDC-2022-0112]

#### Proposed Data Collection Submitted for Public Comment and Recommendations

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice with comment period.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other federal agencies the opportunity to comment on a proposed information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled The Muscular Dystrophy Surveillance, Tracking, and Research Network (MD STARnet) Muscular Dystrophy Questionnaire. The purpose of the proposed study is to describe the epidemiology of COVID-19 and flu and the experience with pain, fatigue, pregnancy, and infertility for adults living with muscular dystrophy (MD) who are identified through MD STARnet. Information will be used to develop interventions that improve the lives of people with MD and their families.

**DATES:** CDC must receive written comments on or before November 22, 2022.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2022-0112 by either of the following methods:

- **Federal eRulemaking Portal:** [www.regulations.gov](http://www.regulations.gov). Follow the instructions for submitting comments.
- **Mail:** Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21-8, Atlanta, Georgia 30329.

**Instructions:** All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to [www.regulations.gov](http://www.regulations.gov).

**Please note:** Submit all comments through the Federal eRulemaking portal ([www.regulations.gov](http://www.regulations.gov)) or by U.S. mail to the address listed above.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21-8, Atlanta, Georgia 30329; Telephone: 404-639-7118; Email: [omb@cdc.gov](mailto:omb@cdc.gov).

**SUPPLEMENTARY INFORMATION:** Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the

collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. 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;
2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected;
4. 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 submissions of responses; and
5. Assess information collection costs.

#### Proposed Project

The Muscular Dystrophy Surveillance, Tracking, and Research Network (MD STARnet) Muscular Dystrophy Questionnaire—New—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

Since its establishment in 2002, the MD STARnet has been a population-based surveillance system that aims to identify and collect clinical data on individuals with muscular dystrophy (MD) in select surveillance areas. MD STARnet identifies and collects data on cases at sources, including healthcare facilities where patients with MD

receive care and administrative datasets such as vital records and hospital discharge data. While MDs are rare genetic diseases with an estimated prevalence of 16.1/100,000, they have a high impact on affected individuals, their families, and society. MDs can be classified into nine major groups: Duchenne muscular dystrophy (DMD), Becker muscular dystrophy (BMD), myotonic dystrophy (DM), facioscapulohumeral muscular dystrophy (FSHD), limb-girdle muscular dystrophy (LGMD), congenital muscular dystrophy (CMD), Emery-Dreifuss muscular dystrophy (EDMD), and distal MD. A recent MD STARnet study has estimated the combined prevalence for DMD and BMD to be 1.92–2.48/10,000

males age 5–9 years old. MD STARnet aims to improve understanding of MDs and ultimately the quality of life of people and their families living with MD. Individuals with MDs frequently report pain and fatigue, but studies have largely been conducted in clinic-based populations and included the three most common MDs. Population-based studies are needed to describe the frequency and management of pain and fatigue and their impact on the lives of individuals with various types of MD.

The purpose of the proposed study is to describe the epidemiology of COVID-19 and flu and the experience with pain, fatigue, pregnancy, and infertility for adults living with MD who are identified through MD STARnet. Results

generated from the study will provide a better understanding of: (1) the occurrence, testing, treatment and severity of COVID-19 in relation to MD; (2) vaccination status and reasons for not receiving COVID-19 and flu vaccinations; (3) the frequency, intensity, and management of pain and fatigue; and (4) the effect of having MD on pregnancy and fertility on adults living with MD. Ultimately, this information can be used to develop interventions that improve the lives of people with MD and their families.

CDC requests OMB approval for an estimated 974 annual burden hours. There are no costs 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) | Total burden hours |
|---------------------------------|-----------------------------------|-----------------------|------------------------------------|----------------------------------------|--------------------|
| Adult males 18 and over .....   | MD STARnet Male Questionnaire ... | 1,794                 | 1                                  | 15/60                                  | 449                |
| Adult females 18 and over ..... | MD STARnet Female Questionnaire   | 1,574                 | 1                                  | 20/60                                  | 525                |
| Total .....                     |                                   |                       |                                    |                                        | 974                |

**Jeffrey M. Zirger,**

*Lead, Information Collection Review Office,  
Office of Scientific Integrity, Office of Science,  
Centers for Disease Control and Prevention.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10146]

#### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are

invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments must be received by November 22, 2022.

**ADDRESSES:** When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for “Comment or Submission” or “More Search Options” to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number: \_\_\_\_, Room C4–26–05,

7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing>.

**FOR FURTHER INFORMATION CONTACT:** William N. Parham at (410) 786–4669.

#### SUPPLEMENTARY INFORMATION:

##### Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

**CMS–10146 Notice of Denial of Medicare Prescription Drug Coverage**  
Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party.