

## FEDLINE 2022 FEE SCHEDULE—Continued

[Effective January 3, 2022. **Bold indicates changes from 2021 prices.**]

	Fee
Software Certification .....	\$0.00 to \$8,000.00.
Vendor Pass-Through Fee .....	various.
Electronic Access Credit Adjustment .....	various.
Electronic Access Debit Adjustment .....	various.

<sup>80</sup> FedComplete packages are all-electronic service options that bundle payment services with an access solution for one monthly fee.

<sup>81</sup> FedComplete customers that use the email service would be charged the FedMail Email a la carte fee and for all FedMail-FedLine Exchange Subscriber 5-packs.

<sup>82</sup> Packages with an "A" include the FedLine Advantage channel, and packages with "C" include the FedLine Command channel.

<sup>83</sup> Per-item surcharges are in addition to the standard fees listed in the applicable priced services fee schedules.

<sup>84</sup> FedComplete customers will be charged \$4 for each FedForward cash letter over the monthly package threshold. This activity will appear under billing code 51998 in Service Area 1521 on a month-lagged basis.

<sup>85</sup> FedMail and FedLine Exchange packages do not include user credentials, which are required to access priced services and certain informational services. Credentials are sold separately in packs of five via the FedMail-FedLine Exchange Subscriber 5-pack.

<sup>86</sup> FedLine Web and Advantage packages do not include user credentials, which are required to access priced services and certain informational services. Credentials are sold separately in packs of five via the FedLine Subscriber 5-pack.

<sup>87</sup> Early termination fees and/or expedited order fees may apply to all FedLine Direct packages and FedLine Direct a la carte options.

<sup>88</sup> These add-on services can be purchased only with a FedLine Solution.

<sup>89</sup> Additional VPNs are available for FedLine Advantage, FedLine Command, and FedLine Direct packages only.

<sup>90</sup> Fee is in addition to the FedLine Direct package fees or additional 2Mbps WAN fees.

<sup>91</sup> The FedLine Custom Implementation Fee is \$2,500 or \$5,000 based on the complexity of the setup.

<sup>92</sup> Available only to customers with a priced FedLine package.

<sup>93</sup> Limited to installed base only.

<sup>94</sup> Five download codes are included at no cost in all Plus and Premier packages.

<sup>95</sup> Cash Management Service options are limited to Plus and Premier packages.

<sup>96</sup> The End of Day Financial Institution Reconciliation Data (FRD) File option is available for FedLine Web Plus, FedLine Advantage Plus, and Premier packages. It is available for no extra fee in FedLine Command Plus and Direct packages.

<sup>97</sup> The Statement of Account Spreadsheet File (SASF) option is available for FedLine Web Plus, FedLine Advantage Plus, and Premier packages. It is available for no extra fee in FedLine Command Plus and Direct packages.

<sup>98</sup> The Intra-day Download Search Results in Spreadsheet Format option is available for the FedLine Web Plus package. It is available for no extra fee in FedLine Advantage and higher packages.

By order of the Board of Governors of the Federal Reserve System.

**Ann Misback,**

*Secretary of the Board.*

[FR Doc. 2021-26395 Filed 12-3-21; 8:45 am]

**BILLING CODE 6210-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[60Day-22-22BC; Docket No. CDC-2021-0128]**

#### 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 and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled *Enhancing Data-drive Disease Detection in Newborns (ED3N)*. CDC is developing this new national newborn screening (NBS) data platform to serve as a secure, central, and national data sharing resource for the U.S. state and territorial NBS community.

**DATES:** CDC must receive written comments on or before February 4, 2022.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2021-0128 by any of the following methods:

- **Federal eRulemaking Portal:** *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 *Regulations.gov*.

**Please note:** Submit all comments through the Federal eRulemaking portal (*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; phone: 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**

Enhancing Data-driven Disease Detection in Newborns (ED3N)—New—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

#### *Background and Brief Description*

The Newborn Screening and Molecular Biology Branch (NSMBB), in the National Center for Environmental Health (NCEH) Division of Laboratory Science (DLS), has the only laboratory in the world devoted to ensuring the accuracy of newborn screening (NBS) tests in every state and more than 78 countries. NSMBB supports NBS programs by conducting research, developing methods, and performing analyses by using complex, state-of-the-art molecular and biochemical techniques for identifying risk factors for diseases of public health importance.

Both NSMBB and state NBS programs are experiencing increased data analytic challenges associated with continued expansion of the number of newborn screening diseases, increased complexity of disease detection, and difficulties in correlating disease markers with disease risk. Further, the addition of late-onset diseases to NBS

panels necessitates a better way to routinely capture clinical information and outcomes so that NBS programs can fully appreciate the spectrum of disease they are detecting.

The NSMBB is requesting a three-year Paperwork Reduction Act (PRA) clearance for Enhancing Data-driven Disease Detection in Newborns (ED3N), a new national NBS data platform, that will address these analytic and post-analytic challenges, and promote sharing of molecular, biochemical, and clinical information amongst NBS partners. The information shared will help NSMBB and newborn screening partners be better equipped to assess disease risk and will help harmonize approaches for disease detection in newborns. Given the rarity of newborn screening diseases, it is imperative that data be collected and analyzed at a national level in order to glean useful insights and to analyze trends. The NSMBB is best suited to oversee this work given its role in providing technical assistance to NBS programs nationally.

Numerous studies along with presentations by NBS programs suggest that gaps in programmatic resources and expertise are hampering the ability to perform more complex data analytics resulting in low positive predictive values for a number of conditions (which subsequently results in higher false positive and negative rates and downstream burden to families and the medical system). Smaller-scale work on the use of post-analytical tools such as machine learning algorithms have shown that incorporation of these

elements into newborn screening can improve detection rates, while reducing false positives. These studies, however, have been limited to single sites and have not been integrated into the daily workflow of high-throughput NBS programs. Without this project, NBS programs will continue to be unable to keep up with the increasing complexity and future demands of screening, perpetuating inequities in screening across the nation.

The estimated annualized burden hours were determined as follows. There are 53 domestic NBS programs in the United States. A “respondent” refers to a single NBS program. Given that data submission will ultimately be accomplished through automatic electronic data transfer, each respondent’s burden hours were split into two estimates: (1) The one-time need to set-up, test, and implement the electronic data transfer mechanism, and 2) the ongoing automatic electronic data transfer occurring after initial set-up. Initial set-up time burden was estimated based on analysis of similar data transfer projects embarked upon by NBS programs as well as brief discussions with NBS Program Laboratory Information Management System vendors. The one-time burden to set up the data transfer interface was estimated to be 40 hours total, annualized to 14 hours per year. Ongoing daily data submission burden for NBS programs was estimated assuming one minute per automatic transfer thereafter. CDC has estimated the total annualized burden for this project to be 1,064 hours per year.

## ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)	Total burden (in hr)
Newborn Screening Programs .....	Set-up and initial submission of ED3N Data Elements.	53	1	14	742
	Ongoing transfer of ED3N Data Elements.	53	364	1/60	322
Total .....	.....	53	.....	.....	1,064

**Jeffrey M. Zirger,**

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

[FR Doc. 2021-26400 Filed 12-3-21; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day-22-22BB; Docket No. CDC-2021-0127]

#### 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 and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled “Building Resilience Against Climate Effects (BRACE) Performance Measures.” The National Center for Environmental Health’s Climate and Health Program (CHP) supports U.S. cities and states to build and enhance resilience to the health impacts of climate change.

**DATES:** CDC must receive written comments on or before February 4, 2022.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2021-0127 by any of the following methods:

- *Federal eRulemaking Portal: 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 *Regulations.gov*.

*Please note: Submit all comments through the Federal eRulemaking portal (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; phone: 404-639-7118; Email: *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

Building Resilience Against Climate Effects (BRACE) Performance Measures—New—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

The purpose of this information collection request (ICR) is to continue the Climate and Health Program (CHP) monitoring of recipient programs’ planning and delivery of public health activities and adaptation strategies under a new cooperative agreement Building Resilience Against Climate Effects: Implementing and Evaluating Adaptation Strategies that Protect and Promote Human Health (CDC-RFA-EH21-2101). CDC collects information related to each recipient’s strategies and activities through performance measures (PMs) outlined by the cooperative agreement. A new PM electronic reporting tool has been developed, which will allow recipients to report PM information in a streamlined way that will also enhance CHP’s ability to analyze and use the information quickly to help support the program. Since its inception, the National Center for Environmental Health’s (NCEH) CHP has funded state and local health departments or their agents as they prepare for and respond to the health effects that a changing climate will bring to the communities they serve. The