**National Notifiable Diseases Surveillance System (NNDSS)**

**OMB Control Number 0920-0728**

**Expiration Date: 03/31/2024**

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**Circumstances of Change Request for OMB 0920-0728**

This is a non-substantive change request for OMB No. 0920-0728, expiration date 03/31/2024, for the reporting of Nationally Notifiable Diseases. Information on proposed disease-specific data elements to be added through this non-substantive change request is enumerated in the table below:

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| --- | --- | --- | --- | --- | --- | --- | --- |
| **Disease Name**  **in NNDSS Collection** | Nationally Notifiable (NNC) OR Under Standardized Surveillance (CSS) | Current Case Notification (Y/N) | Proposed Case Notification (Y/N) | Current Disease-specific Data Elements (Y/N) | Proposed Disease-specific Data Elements (Y/N) | Number of Existing Data Elements in NNDSS | Proposed Number of new NNDSS Data Elements |
| Campylobacteriosis | NNC |  |  | Y |  | 14 | 5 |
| Cryptosporidiosis | NNC |  |  | Y |  | 154 | 5 |
| Cyclosporiasis | NNC |  |  | Y |  | 130 | 1 |
| Hansen’s Disease | NNC |  |  | Y |  | 76 | 7 |
| Hepatitis | NNC |  |  | Y |  | 131 | 49 |
| Listeriosis | NNC |  |  | Y |  | 1573 | 11 |
| *S*. Paratyphi Infection | NNC |  |  | Y |  | 62 | 2 |
| *S*. Typhi Infection | NNC |  |  | Y |  | 68 | 2 |
| Salmonellosis | NNC |  |  | Y |  | 154 | 3 |
| Shiga toxin-producing *Escherichia Coli* (STEC) | NNC |  |  | Y |  | 335 | 1 |
| Shigellosis | NNC |  |  | Y |  | 25 | 3 |

The National Notifiable Diseases Surveillance System (NNDSS) is the nation’s public health surveillance system that enables all levels of public health (local, state, territorial, federal and international) to monitor the occurrence and spread of the diseases and conditions that CDC and the Council of State and Territorial Epidemiologists (CSTE) officially designate as “nationally notifiable” or as under “standardized surveillance.” The NNDSS program creates the infrastructure for the surveillance system and facilitates the submission and aggregation of case notification data voluntarily submitted to CDC from 60 jurisdictions: public health departments in every U.S. state, New York City, Washington DC, 5 U.S. territories (American Samoa, the Commonwealth of Northern Mariana Islands, Guam, Puerto Rico, and the U.S. Virgin Islands), and 3 freely associated states (Federated States of Micronesia, the Republic of the Marshall Islands, and the Republic of Palau). The NNDSS also facilitates relevant data management, analysis, interpretation and dissemination of the information. The data are used to monitor the occurrence of notifiable conditions and to plan and conduct prevention and control programs at the state, territorial, local and national levels.

This request is for the addition of 91 new data elements: 1 new laboratory data element for all conditions, 1 new core data element for all conditions, and 89 new disease-specific data elements. The 89 new disease-specific data elements include: 5 new disease-specific data elements for Campylobacteriosis, 5 new disease-specific data elements for Cryptosporidiosis, 1 new disease-specific data element for Cyclosporiasis, 7 new disease-specific data elements for Hansen’s Disease, 49 new disease-specific data elements for Hepatitis, 11 new disease-specific data elements for Listeriosis, 2 new disease-specific data elements for *S*. Paratyphi Infection, 2 new disease-specific data elements for *S*. Typhi Infection, 3 new disease-specific data elements for Salmonellosis, 1 new disease-specific data element for Shiga toxin-producing Escherichia Coli (STEC), and 3 new disease-specific data elements for Shigellosis.

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| --- | --- | --- | --- | --- |
| **Core: 1 Data Element** | |  | | |
| The impetus/urgency for CDC to add data element for all conditions | | * To make surveillance more comprehensive and informative for public health actions * To provide more information about risk factors (related cases and conditions, high acuity care needs, healthcare facility exposure, travel, and specimen testing) that have been associated with colonization or infection * To monitor epidemiology * To update guidance on infection control and prevention | | |
| **Data Element Name** | **Data Element Description** | | **Value Set Code** | **CDC Priority[[1]](#endnote-1)** | |
| NORS ID | CDC National Outbreak Reporting System (NORS) Outbreak ID Number | | N/A | 1 | |

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| **Laboratory: 1 Data Element** | | |  | | |
| The impetus/urgency for CDC to add data element for all conditions | | | * To make surveillance more comprehensive and informative for public health actions * To provide more information about risk factors (related cases and conditions, high acuity care needs, healthcare facility exposure, travel, and specimen testing) that have been associated with colonization or infection * To monitor epidemiology * To update guidance on infection control and prevention | | |
| **Data Element Name** | **Data Element Description** | | **Value Set Code** | **CDC Priorityi** | |
| Isolate sent to State Public Health Lab | Was the isolate sent to a state public health laboratory? (Answer 'Yes' if it was sent to any state lab, even if it was sent to a lab outside of the case's state of residence) | | PHVS\_YesNoUnknown\_CDC | TBD | |

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| **Campylobacteriosis: 5 Data Elements** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention |

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| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Travel State | Domestic destination, state(s) traveled to | PHVS\_State\_FIPS\_5-2 | 3 |
| International Destination(s) of Recent Travel | International destination or countries the patient traveled to | PHVS\_Country\_ISO\_3166-1 | 3 |
| Date of Arrival to Travel Destination | Date of arrival to travel destination | N/A | 3 |
| Date of Departure from Travel Destination | Date of departure from travel destination | N/A | 3 |
| Reason for travel related to current illness | Reason for travel related to current illness | PHVS\_TravelPurpose\_FDD | 3 |

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| **Cryptosporidiosis: 5 Data Elements** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention |

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| --- | --- | --- | --- |
| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Travel State | Domestic destination, state(s) traveled to | PHVS\_State\_FIPS\_5-2 | 3 |
| International Destination(s) of Recent Travel | International destination or countries the patient traveled to | PHVS\_Country\_ISO\_3166-1 | 3 |
| Date of Arrival to Travel Destination | Date of arrival to travel destination | N/A | 3 |
| Date of Departure from Travel Destination | Date of departure from travel destination | N/A | 3 |
| Reason for travel related to current illness | Reason for travel related to current illness | PHVS\_TravelPurpose\_FDD | 3 |

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| **Cyclosporiasis: 1 Data Element** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention |

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| --- | --- | --- | --- |
| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Reason for travel related to current illness | Reason for travel related to current illness | PHVS\_TravelPurpose\_FDD | 3 |

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| --- | --- | --- | --- | --- |
| **Hansen’s Disease: 7 Data Elements** | |  | | |
| The impetus/urgency for CDC to add data elements for this condition | | * Improve CDC’s understanding of Hansen’s disease epidemiology. * Identify challenges to diagnoses. * Possibly prevent further transmission and lifelong neuropathy and disability given the increase in disease incidence, and lack of information related to medication dosages, recipients, duration, and frequency of administration that is received via current notifications. | | |
| **Data Element Name** | | **Data Element Description** | | **Value Set Code** | **CDC Priorityi** | |
| Medication Frequency | | Frequency of medication administered for this condition. | | N/A | 2 | |
| Medication Frequency Unit | | Unit of measure for the frequency of medication administered (e.g. daily, weekly, monthly). | | TBD | 2 | |
| Medication Duration | | Duration of medication treatment or post-exposure prophylaxis. | | N/A | 2 | |
| Medication Duration Units | | Unit of measure for the duration of medication administered (e.g. days, weeks, months). | | TBD | 2 | |
| Medication Recipient | | Specify recipient of medication for Hansen’s disease (e.g. household contact, case subject). | | TBD | 1 | |
| Medication Dose | | Dosage of medication received. | | N/A | 2 | |
| Medication Dosage Unit | | Unit of measure for medication received (e.g. milligram [mg], milligram/kilogram [mg/kg]) | | TBD | 2 | |

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| --- | --- | --- | --- | --- |
| **Hepatitis: 49 Data Elements** |  | | | |
| The impetus/urgency for CDC to add data elements for this condition | * To improve the collection and sending of data elements for those jurisdictions funded through PS21-2103 “Integrated Viral Hepatitis Surveillance and Prevention Funding for Health Departments” * To make surveillance more comprehensive and informative for public health actions * To monitor perinatal hepatitis C using the new Council of State and Territorial Epidemiologists/CDC definition * To provide more information about risk factors (related cases and conditions, incarceration, travel, not prescribed injection/non-injection drug use, contact with a hepatitis confirmed/suspected person, and homelessness) * To describe the epidemiology and risk factors for hepatitis A related to unprecedented multi-state outbreaks * To update guidance on infection control and prevention | | | |
| **Data Element Name** | | | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Gender Identity | | | Patient identified gender identity (i.e., an individual’s personal sense of being a man, woman, or other gender, regardless of the sex that person was assigned at birth) | TBD | 2 |
| CSTE Case Definition | | | Did the patient meet the CSTE case definition(s) for any of the following in a previous reporting year? (*select all that apply*) | TBD | 2 |
| Information Source for Data | | | Source of Laboratory Test: (*select all that apply*) | TBD | 2 |
| Signs and Symptoms | | | Signs and symptoms associated with the illness being reported | TBD | 1 |
| Signs and Symptoms Indicator | | | Response for each of the signs and symptoms. | [Yes No Unknown (YNU) https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888](https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888) | 1 |
| Date of Symptom Onset | | | The date and time, if available, of the symptom onset (clinical manifestation) | N/A | 1 |
| Date of Jaundice Onset | | | What was the date of jaundice onset? | N/A | 1 |
| Case Patient a Healthcare Worker | | | Was the patient employed as a healthcare worker during the TWO WEEKS prior to onset of symptoms to ONE WEEK after onset of JAUNDICE? (If no jaundice, use two weeks after onset of symptoms) | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 2 |
| Patient Epidemiological Risk Factors | | | Exposed risk factors for the patient - Please provide a response for all risk factors in the value set with an associated indicator. In the 15 to 50 days before symptom onset date for hepatitis A. In the 60 to 150 days (2 to 5 months) before symptom onset date for hepatitis B.  In the 14 to 182 days (2 weeks to 6 months) before symptom onset date for hepatitis C. | TBD | 1 |
| Patient Epidemiological Risk Factors Indicator | | | Provide a response for each value in the patient epidemiological risk factors value set. | [Yes No Unknown (YNU) https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888](https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888) | 1 |
| Contact Type | | | If the patient was a contact of a person with confirmed or suspected hepatitis virus infection, was the contact: (select all that apply) | TBD | 2 |
| Men who have Sex with Men | | | Was the patient a man who reported sexual activity with men? | [Yes No Unknown (YNU) https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888](https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888) | 1 |
| Multiple Sex Partners | | | Did the patient report multiple sex partners? | [Yes No Unknown (YNU) https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888](https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888) | 1 |
| Previous STD History | | | Was the patient diagnosed with a sexually transmitted disease? | [Yes No Unknown (YNU) https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888](https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888) | 2 |
| Antiviral Medication | | | Did the gestational parent receive hepatitis B antiviral therapy during the third trimester of pregnancy? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Birth Weight (unit) | | | The patient's birth weight units | TBD | 1 |
| Vaccinated within 12 Hours of Birth | | | Did the patient receive the hepatitis B vaccine within 12 hours of birth? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Treatment within 12 Hours of Birth | | | Did the patient receive the hepatitis B immune globulin within 12 hours of birth? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Seroconversion | | | If hepatitis B case, did the patient meet the acute hepatitis B seroconversion criteria? (*i.e., documented negative HBsAg laboratory test result within 6 months prior to a positive test [HBsAg, HBeAg, or nucleic acid test for HBV DNA (including qualitative, quantitative, and genotype testing)] in someone without a prior diagnosis of HBV infection*) If hepatitis C case, did the patient meet the acute hepatitis C seroconversion criteria? (e.g., documented negative anti-HCV followed within 12 months by a positive anti-HCV test; or documented negative anti-HCV or negative HCV detection test [in someone without a prior diagnosis of HCV infection] followed within 12 months by a positive HCV detection test; or, in the case of presumed reinfection, at least two sequential negative HCV detection tests [in someone with a prior diagnosis of HCV infection] followed by a positive HCV detection test). | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Occupation and Industry Category | | | Was the patient employed as a food handler or a healthcare worker during the TWO WEEKS prior to onset of symptoms to ONE WEEK after the onset of JAUNDICE? (If no jaundice, use two weeks after onset of symptoms) | TBD | 2 |
| Occupation and Industry Category Indicator | | | Please indicate for each occupation: | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 2 |
| Positive Results 6 Months Apart | | | Did the patient have two positive results at least 6 months apart from any of the following tests: (1) HBsAg; (2) nucleic acid test for HBV DNA (including qualitative, quantitative, and genotype testing); (3) HBeAg?  (*Any combination of these positive tests performed at least 6 months apart is acceptable*) | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Mother's Local Record ID | | | Provide the local record ID used for reporting mother's case of hepatitis (DE Identifier "N/A: OBR-3"). This will be used for linking the reported perinatal case to the mother's reported hepatitis case. | N/A | 3 |
| Mother Nucleic Acid Test | | | For hepatitis B, perinatal, did the gestational parent receive nucleic acid testing for HBV DNA during pregnancy?  For hepatitis C, perinatal, did the gestational parent receive nucleic acid testing for HCV RNA (including qualitative or quantitative PCR, or genotype testing) during pregnancy? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 2 |
| Mother Nucleic Acid Test Result | | | For hepatitis B, perinatal, if the gestational parent received nucleic acid testing for HBV DNA during pregnancy, then indicate the result. For hepatitis C, perinatal, if the gestational parent received nucleic acid testing for HCV RNA (including qualitative or quantitative PCR, or genotype testing) during pregnancy, then indicate the result. | TBD | 2 |
| Mother Nucleic Acid Test Viral Load | | | If the gestational parent received nucleic acid testing for HBV DNA during pregnancy, then indicate the viral load: | TBD | 2 |
| Mother HBeAg Test | | | Did the gestational parent receive HBeAg testing during pregnancy? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 2 |
| Mother HBeAg Test Result | | | If the gestational parent received HBeAg testing during pregnancy, indicate the result. | TBD | 2 |
| Infant HBsAg Test | | | Did the patient receive an HBsAg test between age 1–24 months (only if ≥4 weeks after the last dose of hepatitis B vaccine)? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Infant HBsAg Test Result | | | If the patient received an HBsAg test between age 1–24 months (only if ≥4 weeks after the last dose of hepatitis B vaccine), indicate the result. | TBD | 1 |
| Infant HBsAg Positive Date | | | If positive, then indicate the date of the first positive HBsAg test between age 1-24 months. | N/A | 1 |
| Infant HBeAg Test | | | Did the patient receive an HBeAg test between age 9–24 months? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Infant HBeAg Test Result | | | If the patient received an HBeAg test between age 9–24 months, indicate the result. | TBD | 1 |
| Infant HBeAg Positive Date | | | If positive, then indicate the date of the first positive HBeAg test between age 9-24 months. | N/A | 1 |
| Infant HBV DNA Test | | | Did the patient receive an HBV DNA test between age 9–24 months? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Infant HBV DNA Test Result | | | If the patient received an HBV DNA test between age 9–24 months, indicate the result. | TBD | 1 |
| Infant HBV DNA Positive Date | | | If detected/positive, then indicate the date of the first positive HBV DNA test between age 9-24 months. | N/A | 1 |
| Infant anti-HCV Test | | | Did the patient receive an anti-HCV test between age 18-36 months? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Infant anti-HCV Test Result | | | If the patient received an anti-HCV test between age 18-36 months, indicate the result. | TBD | 1 |
| Infant anti-HCV Positive Date | | | If positive, then indicate the date of the first positive anti-HCV test between age 18-36 months. | N/A | 1 |
| Infant Nucleic Acid Test | | | Did the patient receive nucleic acid testing for HCV RNA (including qualitative or quantitative PCR, or genotype testing) between age 2-36 months? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Infant Nucleic Acid Test Result | | | If the patient received nucleic acid testing for HCV RNA (including qualitative or quantitative PCR, or genotype testing) between age 2-36 months, indicate the result. | TBD | 1 |
| Infant Nucleic Acid Positive Date | | | If detected/positive, then indicate the date of the first positive nucleic acid test for HCV RNA between age 2-36 months. | N/A | 1 |
| Infant HCV Antigen Test | | | Did the patient receive HCV antigen test between age 2-36 months? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Infant HCV Antigen Test Result | | | If the patient received HCV antigen test between age 2-36 months, indicate the result. | TBD | 1 |
| Infant HCV Antigen Positive Date | | | If positive, then indicate the date of the first positive HCV antigen test between age 2-36 months. | N/A | 1 |
| Tissue or organ transplant | | | Did the patient receive tissue or organ transplant(s)? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 2 |
| Non-injection Drug Use | | | Did the patient use non-injection drugs not prescribed by a doctor or engage in nonmedical use of prescription drugs?   v1.0 only: During the 2-6 weeks prior to the onset of symptoms, did the subject inject drugs not prescribed by a doctor? | Yes No Unknown (YNU) <https://phinvads.cdc.gov/vads/ViewValueSet.action?oid=2.16.840.1.114222.4.11.888> | 1 |
| Specimen From Mother or Infant | | | Is the specimen from the gestational parent or the infant? | PHVS\_SpecimenFromMotherOrInfant\_CRS | 1 |

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| --- | --- |
| **Listeriosis: 11 Data Elements** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To bring current LI Case Report Form in alignment with 2019 CSTE case definition changes * To monitor trends related to presumptive and suspected cases in accordance to 2019 CSTE case definition change * To track epi-linked maternal and neonatal cases more accurately * To provide more information about risk factors of Listeriosis * To monitor epidemiology |

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| --- | --- | --- | --- |
| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| CaseStatusAPMother | Case classification of Pregnant mother | PHVS\_CaseClassStatus\_NND | TBD |
| CaseStatusAPNeonate | Case classification of Neonate | PHVS\_CaseClassStatus\_NND | TBD |
| CaseStatusNP | Case classification | PHVS\_CaseClassStatus\_NND | TBD |
| LabCriteria | Laboratory Criteria for Diagnosis | N/A | TBD |
| APNeonateAgeAtCollection | Neonatal age at time of laboratory specimen collection | N/A | TBD |
| ResultCulture | Result of culture-based test on specimen | PHVS\_PosNegUnkNotDone\_CDC | TBD |
| ResultCIDT | Result of CIDT-based test on specimen | PHVS\_PosNegUnkNotDone\_CDC | TBD |
| EpiLink | Indicates the case is epi-linked to a confirmed or probable case | PHVS\_YesNoUnknown\_CDC | TBD |
| PrInfantOutcomeDeathDate | Pregnant: If infant died, when was the date of death (Date) | N/A | TBD |
| LocalRecordIDMother | Pregnant: If mother and infant are counted as separate cases provide the State Epi Case ID of the mother | N/A | TBD |
| LocalRecordIDNeonate | Pregnant: If mother and infant are counted as separate cases provide the State Epi Case ID of the neonate | N/A | TBD |

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| ***S.* Paratyphi Infection: 2 Data Elements** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention |

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| --- | --- | --- | --- |
| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Date of Arrival to Travel Destination | Date of arrival to travel destination | N/A | 3 |
| Travel State | Domestic destination, state(s) traveled to | PHVS\_State\_FIPS\_5-2 | 3 |

|  |  |
| --- | --- |
| ***S.* Typhi Infection: 2 Data Elements** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention |

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| --- | --- | --- | --- |
| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Date of Arrival to Travel Destination | Date of arrival to travel destination | N/A | 3 |
| Travel State | Domestic destination, state(s) traveled to | PHVS\_State\_FIPS\_5-2 | 3 |

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| --- | --- |
| **Salmonellosis: 3 Data Elements** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention |

|  |  |  |  |
| --- | --- | --- | --- |
| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Date of Arrival To Travel Destination | Date of arrival to travel destination | N/A | 3 |
| Date of Departure From Travel Destination | Date of departure from travel destination | N/A | 3 |
| Reason for travel related to current illness | Reason for travel related to current illness | PHVS\_TravelPurpose\_FDD | 3 |

|  |  |
| --- | --- |
| **Shiga toxin-producing *Escherichia Coli* (STEC): 1 Data Element** |  |
| The impetus/urgency for CDC to add data elements for this condition | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention |

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| --- | --- | --- | --- |
| **Data Element Name** | **Data Element Description** | **Value Set Code** | **CDC Priorityi** |
| Reason for travel related to current illness | Reason for travel related to current illness | PHVS\_TravelPurpose\_FDD | 3 |

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| --- | --- | --- | --- | --- | --- |
| **Shigellosis: 3 Data Elements** | |  | | | |
| The impetus/urgency for CDC to add data elements for this condition | | * To make surveillance more comprehensive and informative for public health actions * To monitor epidemiology * To update guidance on infection control and prevention | | | |
| **Data Element Name** | **Data Element Description** | | **Value Set Code** | **CDC Priorityi** |
| Date Of Arrival To Travel Destination | Date of arrival to travel destination | | N/A | 2 |
| Date Of Departure From Travel Destination | Date of departure from travel destination | | N/A | 2 |
| Reason for travel related to current illness | Reason for travel related to current illness | | PHVS\_TravelPurpose\_FDD | 3 |

Burden

The burden to add 91 data elements to NNDSS is applicable to all 50 states, 5 territories, 3 freely associated states, and 2 cities. Although not all territories and freely associated states use electronic, automated transmission for their case notifications, it is expected that they will adopt electronic, automated transmission in the next three years. This burden includes the one-time burden incurred by the respondents to add the data elements to their surveillance system and modify their case notification message. A one-time average burden of 9 hours is incurred for respondents to add 91 data elements to their surveillance system and modify their electronic case notification message to accommodate those 91 additional data elements. This one-time burden of 9 hours is noted in the following table:

One-Time Burden to Add 91 Data Elements to NNDSS

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Type of Respondents** | **Number of Respondents** | **Number of Responses per Respondent** | **Average Burden Per Response (in hours): One-time Addition of 91 Data Elements** |  |
| States | 50 | 1 | 9 |  |
| Territories | 5 | 1 | 9 |  |
| Freely Associated States | 3 | 1 | 9 |  |
| Cities | 2 | 1 | 9 |  |
| Total |  |  |  |  |

The total annualized one-time burden is 180 hours (150 hours for states, 15 hours for territories, 9 hours for freely associated states and 6 hours for cities) as noted in the table below.

Annualized One-Time Burden to Add 91 Data Elements to NNDSS

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Type of Respondents** | **Number of Respondents** | **Number of Responses per Respondent** | **Average Burden Per Response (in hours): Annualized One-time Addition of 91 Data Elements** | **Total Annualized One-Time Burden (in hours)** |
| States | 50 | 1 | 3 | 150 |
| Territories | 5 | 1 | 3 | 15 |
| Freely Associated States | 3 | 1 | 3 | 9 |
| Cities | 2 | 1 | 3 | 6 |
| Total |  |  |  | 180 |

180 hours were added to the existing burden hours in Table A.12A and Table A.12B below.

A.12A. Estimates of Annualized Burden Hours

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Type of Respondents** | **Form Name** | **Number of Respondents** | **Number of Responses per Respondent** | **Average Burden Per Response (in hours)** | **Total Burden (in hours)** |
| States | Weekly (Automated) | 50 | 52 | 20/60 | 867 |
| States | Weekly (Non- automated) | 10 | 52 | 2 | 1,040 |
| States | Weekly (NMI Implementation) | 50 | 52 | 4 | 10,400 |
| States | Annual | 50 | 1 | 75 | 3,750 |
| States | One-time Addition of Diseases and Data Elements | 50 | 1 | 15 | 750 |
| Territories | Weekly (Automated) | 5 | 52 | 20/60 | 87 |
| Territories | Weekly, Quarterly (Non-automated) | 5 | 56 | 20/60 | 93 |
| Territories | Weekly (NMI Implementation) | 5 | 52 | 4 | 1,040 |
| Territories | Annual | 5 | 1 | 5 | 25 |
| Territories | One-time Addition of Diseases and Data Elements | 5 | 1 | 15 | 75 |
| Freely Associated States | Weekly (Automated) | 3 | 52 | 20/60 | 52 |
| Freely Associated States | Weekly, Quarterly (Non-automated) | 3 | 56 | 20/60 | 56 |
| Freely Associated States | Annual | 3 | 1 | 5 | 15 |
| Freely Associated States | One-time Addition of Diseases and Data Elements | 3 | 1 | 15 | 45 |
| Cities | Weekly (Automated) | 2 | 52 | 20/60 | 35 |
| Cities | Weekly (Non-automated) | 2 | 52 | 2 | 208 |
| Cities | Weekly (NMI Implementation) | 2 | 52 | 4 | 416 |
| Cities | Annual | 2 | 1 | 75 | 150 |
| Cities | One-time Addition of Diseases and Data Elements | 2 | 1 | **15** | **30** |
| **Total** |  |  |  |  | **19,134** |

A.12B. Estimates of Annualized Cost Burden

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| **Type of Respondents** | **Form Name** | **Number of Respondents** | **Number of Responses per Respondent** | **Average Burden Per Response (in hours)** | **Total Burden Hours** | **Hourly Wage Rate** | **Respondent Cost** |
| States | Weekly (Automated) | 50 | 52 | 20/60 | 867 | $46.23 | $40,081 |
| States | Weekly (Non-automated) | 10 | 52 | 2 | 1,040 | $37.64 | $39,146 |
| States | Weekly (NMI Implementation) | 50 | 52 | 4 | 10,400 | $46.23 | $480,792 |
| States | Annual | 50 | 1 | 75 | 3,750 | $37.64 | $141,150 |
| States | One-time Addition of Diseases and Data Elements | 50 | 1 | 15 | 750 | $46.23 | $34,673 |
| Territories | Weekly (Automated) | 5 | 52 | 20/60 | 87 | $46.23 | $4,022 |
| Territories | Weekly, Quarterly (Non-automated) | 5 | 56 | 20/60 | 93 | $37.64 | $3,501 |
| Territories | Weekly (NMI Implementation) | 5 | 52 | 4 | 1,040 | $46.23 | $48,079 |
| Territories | Annual | 5 | 1 | 5 | 25 | $37.64 | $941 |
| Territories | One-time Addition of Diseases and Data Elements | 5 | 1 | 15 | 75 | $46.23 | $3,467 |
| Freely Associated States | Weekly (Automated) | 3 | 52 | 20/60 | 52 | $46.23 | $2,404 |
| Freely Associated States | Weekly, Quarterly (Non-automated) | 3 | 56 | 20/60 | 56 | $37.64 | $2,108 |
| Freely Associated States | Annual | 3 | 1 | 5 | 15 | $37.64 | $565 |
| Freely Associated States | One-time Addition of Diseases and Data Elements | 3 | 1 | 15 | 45 | $46.23 | $2,080 |
| Cities | Weekly (Automated) | 2 | 52 | 20/60 | 35 | $46.23 | $1,618 |
| Cities | Weekly (Non-automated) | 2 | 52 | 2 | 208 | $37.64 | $7,829 |
| Cities | Weekly (NMI Implementation) | 2 | 52 | 4 | 416 | $46.23 | $19,232 |
| Cities | Annual | 2 | 1 | 75 | 150 | $37.64 | $5,646 |
| Cities | One-time Addition of Diseases and Data Elements | 2 | 1 | 15 | 30 | $46.23 | $1387 |
| **Total** |  |  |  |  |  |  | **$838,721** |

1. R=Required; 1=Priority 1, 2=Priority 2, 3=Priority 3, TBD=To be determined [↑](#endnote-ref-1)