

National Healthcare Safety Network (NHSN) Overview

CDC's NHSN is a secure, web-based surveillance application that is the nation's most widely used healthcare-associated infection (HAI) tracking system. The application is managed and maintained by the Division of Healthcare Quality Promotion (DHQP) at the Centers for Disease Control and Prevention (CDC).

While ensuring data security, integrity, and confidentiality, NHSN gives healthcare facilities the ability to see their data in real-time and share that information with clinicians and facility leadership, as well as with other facilities (for example, a multihospital system) and partners such as health departments or quality improvement organizations. CDC provides the standard national measures for HAIs as well as analytic tools that enable each facility to assess its progress and identify where additional efforts are needed. In addition, NHSN is the conduit for facilities to comply with Centers for Medicare and Medicaid Services (CMS) infection reporting requirements.

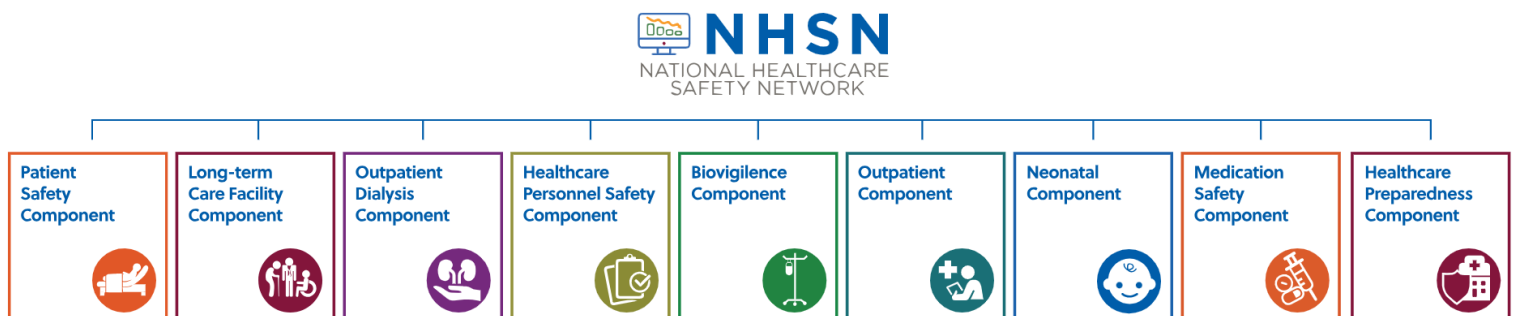
NHSN provides medical facilities, states, regions, and the nation with data collection and reporting capabilities needed to:

- identify infection prevention problems by facility, state, or specific quality improvement project
- benchmark progress of infection prevention efforts
- comply with state and federal public reporting mandates, and ultimately,
- drive national progress toward elimination of HAIs.

[NHSN Fact Sheet](#)

NHSN includes the following components: Patient Safety, Long-term Care Facility, Outpatient Dialysis, Healthcare Personnel Safety, Biovigilance, Outpatient Procedure, Neonatal, Medication Safety, and Healthcare Preparedness (Figure 1).

Figure 1: NHSN Components



The **Patient Safety Component (PSC)** includes modules that focus on events associated with medical devices, surgical procedures, antimicrobial agents used during healthcare, and multidrug resistant organisms.

- Device-associated Modules:
 - Bloodstream Infection (CLABSI – Central line-associated bloodstream infection)
 - Urinary Tract Infection (CAUTI – Catheter-associated urinary tract infection)
 - Pediatric Ventilator-associated Events (PedVAE) (NICU and pediatric locations only)
 - Ventilator-associated Events (VAE) (adult locations only)
 - Pneumonia (VAP – Ventilator-associated Pneumonia) - in pediatric locations (in-plan* or off-plan*), or NICU and adult locations (off-plan* only)
- Procedure-associated Module:
 - Surgical Site Infection (SSI)
- Antimicrobial Use and Resistance Module (AUR)
- Multidrug-Resistant Organism and *Clostridioides difficile* Infection (MDRO/CDI) Module
- Hospital Respiratory Data (HRD) Module

***Note:** “In-plan” surveillance means that the facility has committed to following the NHSN surveillance protocol, in its entirety, for that particular event, as shown in the facility’s NHSN monthly reporting plan. “Off-plan” surveillance is surveillance that is done because a facility has decided to track a particular event for internal use. Data that are entered into NHSN “off-plan” are not included in NHSN annual reports or other NHSN publications. A facility makes no commitment to follow the NHSN protocol for “off-plan” events. Further, “off-plan” data cannot be uploaded into NHSN via Clinical Document Architecture (CDA) and must be manually entered. Instructions and standardized surveillance methods and definitions for each module of the Patient Safety Component are provided in this manual and on the NHSN website (www.cdc.gov/nhsn).

The NHSN **Long-term Care Facility Component** provides long-term care facilities (LTCFs) with standardized surveillance methods and definitions for four modules: 1) Respiratory Pathogens and Vaccination (RPV); 2) Multidrug resistant organism (MDRO) and *Clostridioides difficile* Infection (CDI) Laboratory-identified (LabID) Events; 3) Urinary Tract Infections (UTI); and 4) Prevention Process Measures. The component is accessible to nursing homes, skilled nursing facilities, chronic care facilities, assisted living and residential care facilities, intermediate care facilities for individuals with intellectual disabilities, psychiatric residential treatment facilities, and State Veteran’s Homes. LTCF surveillance protocols, training materials, data collection forms, instructions, and other supporting materials are provided on the Long-term Care Facility Component website: <https://www.cdc.gov/nhsn/ltc/index.html>.

Outpatient hemodialysis centers have surveillance options that are tailored for the specific facility patient population and clinical setting within the **Dialysis Component**. The Dialysis component consists of the following: 1) Dialysis Event (Outpatient Hemodialysis and Acute Kidney Injury (AKI)); 2) Prevention Process Measures; and 3) Summary Data. These modules focus on monitoring and reporting adverse events for the purpose of evaluating prevention efforts among hemodialysis patients. Facilities that treat hemodialysis outpatients should refer to the Dialysis Component protocol, instructions and standardized surveillance methods and definitions at www.cdc.gov/nhsn/dialysis/index.html.

There are two modules in the **Healthcare Personnel Safety (HPS) Component** of NHSN: The Healthcare Personnel Exposure Module and the Healthcare Personnel Vaccination Module. These modules may be used separately or simultaneously. Data collected in this surveillance component can assist healthcare facilities, health systems, and public health agencies to review data on blood/body fluid exposures and antiviral medication use for exposures to influenza, and to monitor influenza and COVID-19 vaccination coverage among healthcare personnel.

The Healthcare Personnel Exposure Module includes Blood/Body Fluid Exposure Only; Blood/Body Fluid Exposure with Exposure Management; and Influenza Exposure Management. This module was retired in June 2025, but facilities can continue to access their historical exposure data for analysis.

The Healthcare Personnel Vaccination Module includes the Influenza Vaccination Summary and COVID-19 Vaccination Summary. Information on reporting influenza vaccination data for healthcare personnel is found here: <https://www.cdc.gov/nhsn/hps/vaccination/index.html>. Information on reporting COVID-19 vaccination data for healthcare personnel is found here: <https://www.cdc.gov/nhsn/hps/weekly-covid-vac/index.html>.

The **NHSN Biovigilance Component**, Hemovigilance Module facilitates national surveillance of transfusion-related recipient adverse events. The Hemovigilance Module is designed for transfusion service staff to collect data on annual facility and transfusion service characteristics, individual reports on adverse transfusion reactions, and yearly counts of transfused or discarded components. The Hemovigilance Module surveillance protocol, training materials, data collection forms, instructions, and other supporting materials are provided on the Hemovigilance Module website: <https://www.cdc.gov/nhsn/biovigilance/blood-safety/index.html>.

The **Outpatient Procedure Component (OPC)** includes two modules that focus on adverse events associated with surgical procedures performed in Ambulatory Surgery Centers (ASCs). The two modules include Same Day Outcome Measures and Surgical Site Infections.

- Same Day Outcome Measures (OPC-SDOM) are a grouping of outpatient care quality indicators that represent a broad range of risks encountered by patients accessing care in various outpatient settings. The four individual outcome measures are:
 - Patient Burn
 - Patient Fall
 - Wrong Site, Wrong Side, Wrong Patient, Wrong Procedure, Wrong Implant
 - All-Cause Hospital Transfer/Admission
- Surgical Site Infection (OPC-SSI) - SSI surveillance for outpatient operative procedures using the Outpatient Procedure Component (OPC).

The OPC surveillance protocols, training materials, data collection forms, instructions, and other supporting materials are provided on the Outpatient Procedure Component website: <https://www.cdc.gov/nhsn/ambulatory-surgery/index.html>.

The **Neonatal Component** includes one module, Late-Onset Sepsis/ Meningitis (LOS/MEN). This module will track late-onset sepsis and meningitis events in very low birthweight neonates housed in Level II/III, Level III, and Level IV nursery locations. The following events will be tracked in the LOS/MEN module:

- Late-Onset Sepsis Event: In an eligible infant, a recognized pathogen or common commensal identified from one or more blood specimens by a culture or non-culture based microbiologic testing method which is performed for purposes of clinical diagnosis or treatment. Under this major type of infection, there are two specific types of infection (see below).
 - NLCBI 1
 - NLCBI 2
- Meningitis Event: In an eligible infant, a recognized pathogen or common commensal identified from a CSF specimen by a culture or non-culture based microbiologic testing method which is performed for purposes of clinical diagnosis or treatment. Under this major type of infection, there are two specific types of infection (see below).
 - NLCM 1
 - NLCM 2

The LOS/MEN surveillance protocols, training materials, data collection forms, instructions, and other supporting materials are provided on the Neonatal Component website:

<https://www.cdc.gov/nhsn/neonatal/index.html>.

The **Medication Safety Component** (MSC) comprises a single module, the Glycemic Control Module. This module focuses on surveillance, measurement, and benchmarking of inpatient medication-related hypoglycemia and hyperglycemia events within healthcare facilities using the Fast Healthcare Interoperability Resources (FHIR) data- exchange standard. Manual and CDA reporting are not available for the Medication Safety Component.

As NHSN gains more experience with the MSC, new areas of focus that can be supported by digital quality measurement will be added. All inpatient facilities enrolled in NHSN are eligible to enroll and participate in the MSC. Long-term care facilities (LTCFs) and outpatient dialysis facilities are not yet eligible to participate in the Medication Safety Component.

The Medication Safety Component protocol is available at the NHSN website:

<https://www.cdc.gov/nhsn/medication-safety-component/index.html>.

The **Healthcare Preparedness Component** is designed to collect data on hospital readiness, operational and critical infrastructure status. The Healthcare Preparedness component will provide NHSN users with resources that support the collection of standardized data on hospital readiness and resource availability. These resources are designed to assess the operational status of U.S. healthcare facilities under normal conditions and during emergencies, including natural disasters and public health crises.

The first module in this component is the Daily Facility Operating Status. The Daily Facility Operating Status module allows facilities to enter daily status updates on evacuation status and critical elements such as power, water, and structural integrity. The report from the module enhances situational awareness and informs coordination and response efforts at the federal, state, and local levels.

The Daily Facility Operating Status module is available for optional reporting of critical infrastructure information for acute care or other short-term stay hospitals. Facilities report every 12 hours during normal operations and during emergency events. The data elements in this module include the overall facility operating status (normal, contingency, emergency) and the following:

- Structural
- Evacuation
- Power
- Water
- Sewer
- Immediate needs

The Daily Facility Operating Status protocols, training materials, data collection forms, instructions, and other supporting materials are provided on the Healthcare Preparedness Component website.

Surveillance Techniques

Surveillance for healthcare associated conditions/infections require a combination of active, concurrent, prospective, or retrospective approaches and surveillance techniques and resources. Trained Infection Preventionists (IPs) and designees shall seek out infections/conditions by screening a variety of data sources, such as laboratory, pharmacy, admission/discharge/transfer, radiology/imaging, and pathology databases, as well as patient charts, including history and physical exam notes, nurses'/ physicians' notes, temperature charts, etc. Others may be trained to screen data sources for these events, but the IP must make the final determination. Laboratory-based surveillance should not be used alone, unless all possible criteria for identifying an infection are solely determined by laboratory evidence (for example, LabID event detection in the MDRO/CDI Module). Retrospective chart reviews should be used only when patients are discharged before all information can be gathered. NHSN forms should be used to collect all required data, using the NHSN definitions of each data field. To minimize the IP's data collection burden, others may be trained to collect the denominator data and process of care data; additionally, electronic capture of data is an option for reporting as an aide to optimizing available resources.

Procedure-Associated Module

Surgical site infection (SSI) monitoring is offered through this module. SSI surveillance requires active, patient-based, prospective surveillance techniques (see Surveillance Techniques above).

Concurrent and post-discharge surveillance methods should be used to detect SSIs following inpatient operative procedures and post-discharge surveillance for outpatient operative procedures. These methods may include 1) direct examination of patients' wounds during hospitalization, or follow-up visits to either surgery clinics or physicians' offices, 2) review of medical records or surgery clinic patient records, 3) visits to the ICU and wards; interview primary care staff, 4) surgeon surveys by mail or telephone, and 5) patient surveys and/or reports (though patients may have a difficult time assessing their infections). Any combination of these methods (or other methods identified by the facility) with the capacity to identify all SSIs is acceptable for use; however, NHSN criteria for SSI must be used. See Surgical Site Infection Event (SSI) protocol for additional examples of concurrent and post-discharge surveillance methods (www.cdc.gov/nhsn/pdfs/pscmanual/9pscscsscurrent.pdf). To minimize IPs' workload of collecting denominator data, operating room data may be downloaded (see file specifications at <https://www.cdc.gov/nhsn/pdfs/ps-analysis-resources/ImportingProcedureData.pdf>).

Device-Associated Module

Medical instrumentation increases the risk of developing an HAI and most patients admitted to a healthcare facility will have a medical device used in the course of their admission. Such devices include, but are not limited to, vascular access devices, indwelling urinary catheters, and ventilators. NHSN enables facilities to monitor for infections associated with the use of these medical devices and to monitor processes related to their use which might increase infection risk. Specifically, NHSN allows facilities to perform surveillance on central line-associated bloodstream infection (CLABSI), catheter-associated urinary tract infection (CAUTI), ventilator-associated events (VAE and PedVAE), and/or ventilator-associated pneumonia (VAP). See the Dialysis Component for detailed instructions for Dialysis Event (DE) surveillance of hemodialysis outpatients (www.cdc.gov/nhsn/dialysis/index.html).

Device-associated denominator data should be collected for CLABSI, CAUTI, VAE, PedVAE, and VAP surveillance (see the CLABSI, CAUTI, VAE, PedVAE, and PNEU protocols for guidance) at the same time each day, or by weekly sampling methods in certain locations. When denominator data are available from electronic databases (for example, ventilator days from respiratory therapy), these sources may be used as long as the counts are not substantially different (+/- 5%) from manually collected counts that have been validated for a minimum of three months. See the respective device-associated event protocols for detailed surveillance instructions.

Antimicrobial Use and Resistance (AUR) Module

The use of antimicrobial agents has a direct effect on antimicrobial resistance patterns of pathogens. The observed increase in multidrug resistance is in part due to inappropriate prescription of, as well as only partial completion of courses of antibiotics.

The AUR Module allows facilities to collect information on the amount of antimicrobials that are used for patient care within their systems, as well as to collect data on the prevalence of drug-resistant organisms in their inpatient and outpatient areas. Electronic capture and reporting of microbiology and pharmacy data are the only available options for reporting data into this module.

See the [Antimicrobial Use and Resistance](#) protocol for detailed surveillance instructions.

Multidrug-resistant Organism and *Clostridioides difficile* Infection (MDRO/CDI) Module

The NHSN MDRO/CDI Module offers a means for facilities to meet criteria and metrics that are outlined in several organizational guidelines to control and measure the spread of MDROs and CDI within their healthcare system. The module has two separate and independent reporting options, Laboratory-identified (LabID) Event and Infection Surveillance, that may be tailored to meet the needs of participating NHSN facilities.

In addition, the following process measures are available: (1) adherence to hand hygiene; (2) adherence to contact precautions when caring for patients infected or colonized with an MDRO or *C. difficile*; and (3) adherence to active surveillance testing (AST) of MRSA and/or VRE. Active surveillance testing outcome measures is also available in locations where AST adherence is being performed and enables facilities to use the results of AST to monitor the incidence and prevalence of positive MRSA and/or VRE cultures. See the [MDRO/CDI](#) protocol for detailed surveillance instructions.