



Hemovigilance Module - Annual Facility Survey Acute Care Facility

*Required for saving

*Facility ID#: _____

*Survey Year: _____

For all questions, use information from previous full calendar year.

Facility Characteristics

NOTE: Questions 1 – 7 are completed automatically (i.e., auto-populated) in the NHSN application with responses from the previous year's survey.

*1. Ownership: (check one)

- Government Military Not for profit, including church
 For profit Veteran's Affairs Physician-owned

*2. Is your hospital a teaching hospital for physicians and/or physicians-in-training? Yes No

If Yes, check type:

- Major Graduate Undergraduate

*3. Community setting of facility: Urban Suburban Rural

*4. How is your hospital accredited? (check one)

- The Joint Commission American Osteopathic Association (AOA)
 National Integrated Accreditation for Healthcare Organizations (DNV) Other Accrediting Organization

*5. Total beds served by the transfusion service. _____

*6. Number of surgeries performed per year:

Inpatient: _____ Outpatient: _____

*7. At what trauma level is your facility certified? I II III IV N/A

Transfusion Service Characteristics

*8. Primary classification of facility areas served by the transfusion service: (check all that apply)

- Cancer center Orthopedic General medical and surgical
 Children's cancer center Children's orthopedic Children's general medical and surgical
 Chronic disease Burn center Obstetrics/Gynecology
 Children's chronic disease Trauma/Emergency Other (specify) _____

*9. Does your healthcare facility provide all of its own transfusion services, including all laboratory functions?

- Yes No, we contract with a blood center for some transfusion service functions.
 No, we contract with another healthcare facility for some transfusion service functions.

*10. Is the transfusion service part of the facility's core laboratory? Yes No

*11. How many dedicated transfusion service staff members are there? (Count full-time equivalents; include supervisors.)

Physicians: _____ Medical Technologists: _____ Medical Laboratory Technicians: _____

Assurance of Confidentiality: The voluntarily provided information obtained in this surveillance system that would permit identification of any individual or institution is collected with a guarantee that it will be held in strict confidence, will be used only for the purposes stated, and will not otherwise be disclosed or released without the consent of the individual, or the institution in accordance with Sections 304, 306 and 308(d) of the Public Health Service Act (42 USC 242b, 242k, and 242m(d)).



The burden of this collection of information is estimated to average 1 hour and 25 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC, Reports Clearance Officer, 1600 Clifton Rd., MS D-74, Atlanta, GA 30333 ATTN: PRA (0920-0666).

- *12. Does your hospital have a dedicated position or FTE in a quality or patient safety function (e.g., TSO) for investigation of transfusion-related adverse reactions? Yes No
- *13. Does your hospital have a dedicated position or FTE in a quality or patient safety function (e.g., TSO) for investigation of transfusion errors (i.e., incidents)? Yes No
- *14. Is the transfusion service laboratory accredited? Yes No
 If Yes, select all that apply: College of American Pathologists (CAP) AABB TJC
- *15. Does your facility have a committee that reviews blood utilization? Yes No
- *16. Total number of patient samples collected for type and screen or crossmatch: _____
- *17. Are any of the following issued through the transfusion service? (check all that apply)
 Albumin Factors (VIIa, VIII, IX, ATIII, etc.) Immunoglobulin (IV)
 Immunoglobulin (IM or subcutaneous) Rhlg None
- *18. Does your facility attempt to transfuse only leukocyte-reduced or leuko-poor cellular components? Yes No
- *19. Are all units stored in the transfusion service? Yes No
 If No, indicate the location(s) of satellite storage: (check all that apply)
 Ambulatory Care Cancer Center Cardiac ICU
 Emergency Department Labor and Delivery Medical Flight Facility
 Operating Room Other: (specify) _____
- *20. To what extent does the transfusion service modify products? (check all that apply)
 Aliquot Deglycerolizing Irradiation Leukoreduction
 Plasma reduction Pooling Washing None of these
- *21. Do you collect blood for transfusion at your facility? Yes No
 If Yes, check all that apply: Allogeneic Autologous Directed
- *22. Does your facility perform viral testing on blood for transfusion? Yes No
- *23. Does your facility perform point-of-issue bacterial testing on platelets prior to transfusion? Yes No

Transfusion Service Computerization

- *24. Is the transfusion service computerized? Yes No (If No, skip to next section)
 If Yes, select system(s) used: (check all that apply) BBCS® BloodTrack Tx® (Haemonetics)
 Cerner Classic® Cerner Millennium® HCLL® Horizon BB® Hemocare®



- Lifeline® Meditech® Misys® Safetrace Tx® (Haemonetics) Softbank®
- Western Star® Other (specify) _____

- *25. Is the system ISBT-128 compliant? Yes No
- *26. Does the transfusion service system interface with the patient registration system? Yes No
- *27. Are the transfusion service adverse events entered into a **hospital-wide** electronic reporting system?
 Yes No If Yes, specify system used: _____
- *28. Does your facility use positive patient ID technology for the transfusion service?

- Yes, hospital wide Yes, certain areas Not used
- If Yes, select purpose(s): (check all that apply) Specimen collection Product administration
- If Yes, select system(s) used: (check all that apply)
- Mechanical barrier system (e.g., Bloodloc®)
- Separate transfusion ID wristband system (e.g., Typenex®)
- Radio frequency identification (RFID) Bedside ID band barcode scanning
- Other (specify) _____

- *29. Does your facility have physician online order entry for test requesting? Yes No
- *30. Does your facility have physician online order entry for product requesting? Yes No

Transfusion Service Specimen Handling and Testing

- *31. Are transfusion service specimens drawn by a dedicated phlebotomy team?
 Always Sometimes, approximately _____% of the time Never
- *32. What specimen labels are used at your facility? (check all that apply)
 Handwritten Addressograph Computer generated from laboratory test request
 Computer generated by bedside device Other (specify) _____
- *33. Are phlebotomy staff members allowed to correct patient identification errors on pre-transfusion specimen labels?
 Yes No
- *34. What items can be used to verify patient identification during specimen collection and prior to product administration at your facility? (check all that apply)
 Medical record (or other unique patient ID) number Date of birth Gender
 Patient first name Patient last name Transfusion specimen ID system (e.g., Typenex®)
 Patient verbal confirmation of name or date of birth
 Sex at Birth Gender Identity Other (specify) _____
- *35. How is routine type and screen done? (check all that apply and estimate frequency of each)
 Manual technique _____% Automated technique _____%
 Both automated and manual technique _____% *Total should equal 100%*
- *36. Is the ABO group of a pre-transfusion specimen routinely confirmed? Yes No



If Yes, check one:

- All samples
- If there is no laboratory record of previous determination of patient's ABO group
- If there is no laboratory record of previous determination of patient's ABO group AND the patient is a candidate for electronic crossmatching

If Yes, is the confirmation required on a separately-collected specimen before a unit of Group A, B or AB red blood cells is issued for transfusion? Yes No

*37. How many RBC type and screen and crossmatch procedures were performed at your facility by any method?

RBC type and screen: _____ RBC crossmatch _____

Estimate the % of crossmatch procedures done by each method: (check all that apply)

Electronically _____% Serologically _____% Don't know *Total may be >100%*