

## Hemovigilance Module



### Annual Facility Survey

\*Required fields

\*Tracking # / Facility ID: \_\_\_\_\_

\*Survey Year: \_\_\_\_\_

**Facility Characteristics:** (For all questions use past full calendar year annual statistics)

\*1. Ownership: (Check one)

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

\*2. Is your hospital affiliated with a medical school?    Yes    No

If yes, type of affiliation:    Major    Graduate    Limited

\*3. Community setting of facility:    Urban    Suburban    Rural

\*4. Total beds set up and staffed: \_\_\_\_\_

\*5. Number of surgeries performed per year:   Inpatient \_\_\_\_\_   Outpatient \_\_\_\_\_

\*6. At what trauma level is your facility certified? \_\_\_\_\_    N/A

**Transfusion Services Characteristics:**

\*7. Primary classification of facility areas served by Transfusion Services: (Check all that apply)

- General medical and surgical    Obstetrics and gynecology    Orthopedic  
 Cancer center    Chronic disease    Children's general medical and surgical  
 Children's orthopedic    Children's cancer center  
 Children's chronic disease    Other (specify)\_\_\_\_\_

\*8. Is your Transfusion Services part of the facility's core laboratory?    Yes    No

\*9. How many dedicated Transfusion Services staff are there?

Number of technical FTEs (including supervisors): \_\_\_\_\_

Number of dedicated physician FTEs: \_\_\_\_\_

Number of MLT: \_\_\_\_\_   Number of MT: \_\_\_\_\_

**Assurance of Confidentiality:** The 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)).

Public reporting burden of this collection of information is estimated to average 2 hours 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).

- \*10. Is your Transfusion Services Laboratory Accredited?  Yes  No  
If Yes, select all that apply:  College of American Pathologists (CAP)  AABB
- \*11. How is your hospital accredited? (Check one)  
 The Joint Commission  Centers for Medicare and Medicaid Services
- \*12. Do you have a committee that reviews blood utilization?  Yes  No
- \*13. What is the total number of samples collected in the past year: \_\_\_\_\_
- \*14. Products and Units Transfused: (Check all that apply)
- Red Blood Cells (RBCs)  
Total number of units transfused in the past year: \_\_\_\_\_  
Number of units from which aliquots were made: \_\_\_\_\_  
Number of aliquots transfused: \_\_\_\_\_
- Platelets  
Number of units of whole blood derived platelet concentrates transfused: \_\_\_\_\_  
What is your average pool size? \_\_\_\_\_  
Number of units of apheresis platelets transfused: \_\_\_\_\_
- Plasma            Number of units transfused: \_\_\_\_\_ (Incl. FFP, thawed, etc.)  
 Cryoprecipitate    Number of units transfused: \_\_\_\_\_  
 Granulocytes      Number of units transfused: \_\_\_\_\_  
 Lymphocytes        Number of units transfused: \_\_\_\_\_
- \*15. Are any of the following administered through Transfusion Services? (Check all that apply)
- Albumin     Factors (VIIa, VIII, IX, ATIII, etc)     Immunoglobulin (IV)  
 Immunoglobulin (IM or subcutaneous)     RHlg     None
- \*16. Does your facility attempt to transfuse only leukocyte-reduced cellular components?  
 Yes     No

17. Units Transfused by Department/Service

Department/Service	Samples Collected	Units Transfused						
		Platelets		RBCs	Plasma	Cryo	Grans	Lymphs
		Apheresis	Whole Blood Derived					
Emergency Room/Trauma								
Hematology/Oncology (Incl. Bone marrow transplant & apheresis)								
ICU								
Nephrology/Dialysis								
Obstetrics/Gynecology								
Pediatrics/Neonatology								
Surgery – cardiac								
Surgery – general								
Surgery – orthopedic								
Surgery – other								
Transplant – solid organ								

\*18. Are all units stored in the Transfusion Services area?  Yes  No

If No, indicate the location of satellite storage: (Check all that apply)

- Operating room  Emergency room  Ambulatory care  
 Other: \_\_\_\_\_

\*19. To what extent does Transfusion Services modify products? (Check all that apply)

- Aliquot  Deglycerolizing  Irradiation  Leukoreduction  
 Plasma reduction  Pooling  Washing  None of these

\*20. Do you collect blood for transfusion at your facility?  Yes  No

If Yes, check all that apply:  Allogeneic  Autologous  Directed

\*21. Does your facility perform viral testing on blood for transfusion?  Yes  No

**Transfusion Services Computerization:**

\*22. Is Transfusion Services computerized?

- Yes  No SKIP to next section

If Yes, system used: (Check all that apply)  Cerner Classic®  Cerner Millenium®  HCLL®

Horizon BB®  Hemocare®  Lifeline®  Meditech®  Mysis®

Wingate® (Safetrace TX)  Softbank®  Western Star®

Other (specify) \_\_\_\_\_

- \*23. Is your system ISBT-128 compliant?  Yes  No
- \*24. Does the Transfusion Services system interface with the patient registration system?  
 Yes  No
- \*25. Are Transfusion Services adverse events entered into a hospital-wide reporting computer system?  
 Yes Specify system used: \_\_\_\_\_  No
- \*26. Do you use positive patient ID technology for transfusion services?  
 Yes, hospital wide  Yes, certain areas  Not used [SKIP to Q. 27]  
If Yes, used for: (Check all that apply)  Specimen collection  Product administration  
Indicate system 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) \_\_\_\_\_
- \*27. Do you have physician on-line order entry for test requesting?  Yes  No
- \*28. Do you have physician on-line order entry for product requesting?  Yes  No

**Transfusion Service Specimens: Handling/Testing**

- \*29. Are the Transfusion Service specimens drawn by a dedicated phlebotomy team?  
 Always  Sometimes, approximately \_\_\_\_% of the time  Never
- \*30. 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) \_\_\_\_\_
- \*31. Are phlebotomy staff allowed to correct errors in patient identification on pre-transfusion specimen labels?  Yes  No
- \*32. 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  Other (specify) \_\_\_\_\_
- \*33. How are routine type and screen done? (Check all that apply)  
 Manual technique % done \_\_\_\_\_  Automatic technique % done \_\_\_\_\_

Automatic and manual % done \_\_\_\_\_

\*34. Is the ABO group of a pre-transfusion specimen routinely confirmed?

Yes (Check one)

No

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 RBC unit of Group A, B or AB is issued for transfusion?  Yes  No

\*35. How many RBC type and crossmatch procedures were performed at your facility by any method?

RBC type: \_\_\_\_\_

Estimated % done by each method: (Check all that apply)

electronically \_\_\_\_\_  serologically \_\_\_\_\_  Don't know

RBC crossmatch: \_\_\_\_\_

Estimated % done by each method: (Check all that apply)

electronically \_\_\_\_\_  serologically \_\_\_\_\_  Don't know