

Form Approved OMB No. 0920-0666 Exp. Date: 12/31/2026 www.cdc.gov/nhsn

Hemovigilance Module - Annual Facility Survey Non-Acute Care Facility

*Required for saving		•			
*Facility ID#:		*Survey Year:			
For all questions, use information from	n previous full calendar year.				
Facility Characteristics					
*1. Ownership: (check one)					
☐ Government ☐ Military	☐ Not for pr	rofit, including church			
☐ For profit ☐ Veteran's Affairs ☐ Physician-owned					
*2. Community setting of facility:	☐ Urban ☐ Suburban	☐ Rural			
*3. Total number of operating rooms	s at time of survey completi	ion:			
*4. Total number of procedure room	s at time of survey complet	tion:			
*5. Total number of patient admission	ons in this survey year:				
*6. Check all the specialty(ies) curre	ently performed in your facil	lity:			
☐ Bariatrics	☐ General surgery	☐ Gastroenterology			
☐ Gynecology	□ Neurology	☐ Orthopedic			
☐ Plastic surgery	Spine	☐ Urology			
Other (specify)					
Transfusion Service Characteris	stics				
*7. Does your facility provide all of it	ts own transfusion services	, including all laboratory f	functions?		
☐ Yes ☐ No, we contract w	with a blood center for some	<u>e</u> transfusion service fund	ctions.		
☐ No, we contract with anoth	er healthcare facility for son	<u>me</u> transfusion service fu	nctions.		
☐ No, we contract with anoth	er blood center for <u>all</u> trans	fusion service functions.			
☐ No, we contract with anoth	er healthcare facility for all	transfusion service functi	ions.		
*8. How many dedicated transfusion	n service staff members are	there? (Count full-time equiva	alents; include s	supervisors.)	
	Technologists:	Medical Laborator	ry Technicia	ıns:	
 Does your facility have a dedica function (e.g., TSO) for investigation 			☐ Yes	□No	
*10. Does your facility have a dedic function (e.g., TSO) for investig			☐ Yes	□No	

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)).

Public reporting burden of this collection of information is estimated to average 35 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 H21-8, Atlanta, GA 30333, ATTN: PRA (0920-0666).



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Transfusion Service Characteristics (continued)		
*11. Does your facility have a committee that reviews blood utilization?	☐ Yes	☐ No
*12. Total number of patient samples collected for type and screen or crossmatch:		
*13. Does your facility perform point-of-issue bacterial testing on platelets prior to transfusion?	☐ Yes	☐ No
Transfusion Service Computerization		
*14. Is the transfusion service computerized? Yes No (If No, skip to question 17)		
If Yes, select system(s) used: (check all that apply) ☐ BBCS [®] ☐ BloodTrack Tx [®] (Hand the content of the co	aemonetics	s)
☐ Cerner Classic [®] ☐ Cerner Millennium [®] ☐ HCLL [®] ☐ Horizon BB [®]	Hemoc	are®
☐ Lifeline [®] ☐ Meditech [®] ☐ Misys [®] ☐ Safetrace Tx [®] (Haemonetics) ☐ S	Softbank [®]	
☐ Western Star [®] ☐ Other (specify)		
*15. Is the system ISBT-128 compliant?		
*16. Does the transfusion service system interface with the patient registration system?	es 🗌 No	
*17. Does your facility use positive patient ID technology for transfusion?		
☐ Yes, facility wide ☐ Yes, certain areas ☐ Not used		
If Yes, select purpose(s): (check all that apply) Specimen collection Produc	t administra	ation
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)		
Transfusion Service Specimen Handling and Testing		
*18. Are transfusion service specimens drawn by a dedicated phlebotomy team?		
☐ Always ☐ Sometimes, approximately% of the time ☐ Never		
*19. What specimen labels are used at your facility? (check all that apply)		
☐ Handwritten ☐ Addressograph ☐ Computer generated from laboratory te	st request	
☐ Computer generated by bedside device ☐ Other (specify)		
*20. Are phlebotomy staff members allowed to correct patient identification errors on pre-transfulabels?	sion specir	men
☐ Yes ☐ No		
*21. What items can be used to verify patient identification during specimen collection and prior administration at your facility? (check all that apply)	to product	
☐ Medical record (or other unique patient ID) number ☐ Date of birth		
☐ Gender identity ☐ Sex at birth		
☐ Patient first name ☐ Patient last name ☐ Transfusion specimen ID system (e	.g., Typend	ex®)
☐ Patient verbal confirmation of name or date of birth ☐ Other (specify)		