

## Hemovigilance Module Monthly Reporting Denominators

### \*Required for saving

#### Table 1

Products			*Units Transfused	*Aliquots Transfused	*Total Discards
Whole Blood		TOTAL			
Red blood	ed blood ells Whole blood derived Irradiate derived Irradiate Irradiate Apheresis Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate Irradiate	TOTAL			
		Not irradiated or leukocyte reduced			
		Irradiated			
		Leukocyte reduced			
		Irradiated and leukocyte reduced			
cells		TOTAL			
		Not irradiated or leukocyte reduced			
		Irradiated			
		Leukocyte reduced			
		Irradiated and leukocyte reduced			
	Whole blood derived	TOTAL			
		Not irradiated or leukocyte reduced			
		Irradiated			
		Leukocyte reduced			
Platelets		Irradiated and leukocyte reduced			
T latelets		sis Irradiated Leukocyte reduced Irradiated and leukocyte reduced TOTAL Not irradiated or leukocyte reduced Irradiated Leukocyte reduced Irradiated and leukocyte reduced TOTAL Not irradiated or leukocyte reduced Sis Irradiated			
	Apheresis	Not irradiated or leukocyte reduced			
		Irradiated			
		Leukocyte reduced			
		Irradiated and leukocyte reduced			
Plasma	Total whole blood derived				
(all types)	Total apheresis				
Cryoprecipitate					

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 70 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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\*Does your facility transfuse blood products treated with pathogen reduction technology? \_\_\_\_\_ Yes \_\_\_\_\_ No ^If yes, then complete Table 2. **Table 2** 

			Units	Aliquots	Total
Products			Transfused	Transfused	Discards
Red blood cells	Whole	TOTAL			
	blood	S-303-treated			
	derived	Riboflavin-treated			
		TOTAL			
	Apheresis	S-303 -treated			
		Riboflavin-treated			
	Whole	TOTAL			
	blood	Psoralen-treated			
Distalata	derived	Riboflavin-treated			
Platelets		TOTAL			
	Apheresis	Psoralen-treated			
		Riboflavin-treated			
	Whole	TOTAL			
	blood	Psoralen-treated			
Plasma	derived	Riboflavin-treated			
(all types)		TOTAL			
	Apheresis	Psoralen-treated			
		Riboflavin-treated			
Cryoprecipitate		TOTAL			
		Psoralen-treated			
		Riboflavin-treated			
		Pathogen Reduction			
		Cryoprecipitated Fibrinogen Complex			

^If your facility transfused pathogen reduced apheresis platelets (e.g., the apheresis platelet total in table 2 is greater than 0), then complete Table 3.

### Table 3

Products			Units Transfused	Aliquots Transfused	Total Discards
	Apheresis	Psoralen-treated			
		Psoralen-treated and in Plasma			
Platelets		Psoralen-treated and in Platelet additive solution			
		Riboflavin-treated			
		Riboflavin-treated and in Plasma			
		Riboflavin-treated and in Platelet additive solution			

\*Patient samples collected for type and screen or crossmatch: \_\_\_\_\_

\*Total crossmatch procedures: \_\_\_\_\_

Total patients transfused: \_\_\_\_\_



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Custom Fields					
Label		Label			