



Advancing Transfusion and Cellular Therapies Worldwide

AABB Supplementary Questionnaire Transfusion-Transmitted Bacterial Infection

Reasons for Investigation of Bacterial Sepsis

[Why are you investigating this as Blood Component Associated Bacterial Sepsis?]

In the left column, please check the event(s) that triggered your report of sepsis associated with transfusion. In the middle column, please provide all available sign/symptom information, whether or not it was the primary trigger for the investigation.

Event	Signs and Symptoms	Time - sign/symptom noted after the start of the transfusion
<input type="checkbox"/> Temperature Change	Pre-transfusion: _____ °F /°C Post-reaction max: _____ °F /°C	
<input type="checkbox"/> Blood Pressure Change	Pre-transfusion: _____ mmHg (systolic/diastolic) Post-reaction: BP Max: _____ mmHg BP Min: _____ mmHg (systolic/diastolic)	
<input type="checkbox"/> Respiration rate	Pre-transfusion: _____ Post-reaction: _____ (breaths per minute)	
<input type="checkbox"/> Shock ¹		
<input type="checkbox"/> Chills/ Rigors		
<input type="checkbox"/> CNS signs and symptoms	Describe using NHSN Instructions:	
<input type="checkbox"/> Multiple organ dysfunction syndrome (progressive dysfunction of two or more major organ systems in a critically ill patient that makes it impossible to maintain homeostasis without medical intervention)	Describe:	
<input type="checkbox"/> Other:	Describe:	

¹ NHSN definition: A drop in blood pressure accompanied by a drop in cardiac output including rapid heart rate (increase to 100 beats per minute or more), rapid breathing, cutaneous vasoconstriction, pallor, sweating, decreased or scanty urine production, agitation and/or loss of consciousness that required fluid resuscitation, with or without inotropic support (9_2013).

Organization ID: _____
 NHSN Adverse Reaction: # _____ Date of reaction: _____

Did clinicians at your hospital diagnose sepsis in this patient using SIRS criteria²?

Yes No Don't Know

Suspected component (check all that apply):

RED BLOOD CELLS PLATELETS PLASMA OTHER: _____

Laboratory Testing: Bacteria				
	Culture Performed (check one)	Culture Results (check one)	Identity of Organism	Date/Time of Sampling
Patient Pre-transfusion: (check all that apply)				
<input type="checkbox"/> Blood	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Don't Know	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Don't Know		
<input type="checkbox"/> Urine	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Don't Know	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Don't Know		
<input type="checkbox"/> Other: _____	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Don't Know	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Don't Know		
Patient Post-transfusion:				
<input type="checkbox"/> Blood <input type="checkbox"/> Other: _____	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Don't Know	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Don't Know		
Suspect Unit(s) ³ :	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Don't Know	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Don't Know		
Suspect Unit(s): Final Result from platelet QC culture done at collection facility (if known and applicable).	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Don't Know	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Don't Know		
Point-of-issue bacterial screening result if performed (e.g. PGD or BACTx)?	<input type="checkbox"/> BACTx <input type="checkbox"/> PGD <input type="checkbox"/> Other:	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Don't Know		
Did co-components from the involved donation cause septic reaction(s) in other recipients?				<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Don't Know
If bacterial strain(s) were isolated from the patient and the implicated component and/or co-components, were they evaluated for identity? Strain _____ If yes, please check the method was used:				<input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Antimicrobial Sensitivity <input type="checkbox"/> RFLP/PFGE <input type="checkbox"/> Sequencing <input type="checkbox"/> Other (please describe): _____				
If agents were evaluated for identity, were the donor and patient agents identical?				<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Were there other documented, concurrent infections in the patient? If yes, please describe (type of infection/organism and time):				<input type="checkbox"/> Yes <input type="checkbox"/> No

² ACP/SCCM Consensus Panel Guidelines.
 Sepsis is culture-documented infection plus two or more of the following

- T of >38C or <36C
- Heart rate >90
- RR >20
- WBC count >12000 or <4000 or ≥10% band forms

³ Sample from implicated unit. Cultures performed only with an integral segment, unless positive, should be considered NOT performed since, due to low concentrations of bacteria, they may be sterile at the time they are sealed in the collection facility.