

Call NCBB at (402) 486-9453 and fax this form to (402) 486-9439 as soon as possible.

When completed, fax the transfusion reaction investigation report and supporting documentation to (402) 486-9439.

SERIOUS OUTCOME OF TRANSFUSION REPORT

REASON(S) FOR REPORT: (check appropriate boxes for suspected reaction types)

- Anaphylaxis or severe allergic reaction
- Death potentially related to transfusion [Notify FDA by email: fatalities2@cber.fda.gov]
- Transfusion-associated circulatory overload (TACO)
- Transfusion-associated sepsis
- Transfusion-related acute lung injury (TRALI)
- Transfusion-transmitted disease (TTD) [STOP HERE and complete Suspected Transfusion Transmitted Disease Report (NE-Form-0493)]
- Other (describe) _____

Facility Reporting Serious Outcome _____

Person completing form _____ Date _____

Enter units involved, date(s) of transfusion, and time that transfusion started and ended:

Unit or Pool #	Component	Date	Time (start and end)
			/
			/
			/

COMPLETE WITH AVAILABLE INFORMATION:

Recipient's name _____ DOB _____

Recipient's hospital identification # _____ Gender M F

Recipient's diagnosis _____

Indication for transfusion _____

Describe transfusion facility investigation, treatment and recipient response to date. (Use additional pages if needed.)
(A transfusion reaction investigation form is available on the NCBB website).

Have bags and/or segments been retained for further investigation? Y N
If yes, record date and time blood bag/segments refrigerated: _____

Please attach a copy of the transfusion record.

Transfusion Service Medical Director (or designee) Signature

Date

SOOT _____