



Hemovigilance Module Adverse Reaction Hypotensive Transfusion Reaction

| *Required for saving | | |
|---|---|---|
| *Facility ID#: NHS | N Adverse Reaction #: | |
| Patient Information | | |
| *Patient ID: | *Gender: M F | Other *Date of Birth:// |
| Social Security #: | | Medicare #: |
| Last Name: | First Name: | Middle Name: |
| Ethnicity Hispanic or Latin | o Not Hispanic or Not Latino | 0 |
| Race American Indian | Alaska Native 🔲 Asian | Black or African American |
| Native Hawaiian | Other Pacific Islander | White |
| *Blood Group: A- A+ | B- B+ AB- AB+ | O- O+ Blood type not done |
| | | |
| Patient Medical History (Us | e worksheet on page 4 for additiona | al codes and descriptions.) |
| (part 1) List the patient's adr | nitting diagnosis. (Use ICD-10 Diagn | ostic codes/descriptions) |
| Code: | Description: | |
| Code: | | |
| Code: | Description: | |
| | erlying indication for transfusion. (Us | e ICD-10 Diagnostic codes/descriptions) |
| Code: | Description: | |
| Code: | | |
| Code: | Description: | |
| | norbid conditions at the time of the tra O Diagnostic codes/descriptions) | ansfusion related to the UNKNOWN NONE |
| Code: | Description: | |
| Code: | | |
| Code: | Description: | |
| | | Continued >> |
| of any individual or institution is collections and will not otherwise be discleded by the sections 304, 306 and 308(d) of the Fublic reporting burden of this collection reviewing instructions, searching exist collection of information. An agency runless it displays a currently valid OM | ted with a guarantee that it will be held in so sed or released without the consent of the tublic Health Service Act (42 USC 242b, 24 on of information is estimated to average 2 ing data sources, gathering and maintaining not conduct or sponsor, and a person in B control number. Send comments regard to gestions for reducing this burden to CDC, | is surveillance system that would permit identification strict confidence, will be used only for the purposes individual, or the institution in accordance with 42k, and 242m(d)). To minutes per response, including the time for any the data needed, and completing and reviewing the is not required to respond to a collection of information ling this burden estimate or any other aspect of this Reports Clearance Officer, 1600 Clifton Rd., MS D-74, |



| Patient Medical History (Use worksheet on page 4 for additional codes and descriptions.) |
|---|
| (part 4) List the patient's relevant medical procedure including past procedures and procedures to be performed during the current hospital or outpatient stay. (Use ICD-10 Procedure codes/descriptions) |
| Code: Description: |
| Code: Description: |
| Code: Description: |
| (part 5) Additional Information |
| |
| |
| Transfusion History (Use worksheet on page 4 for additional transfusion history.) |
| Has the patient received a previous transfusion?YESNOUNKNOWN |
| **If yes, provide information about the transfusion event. If not, skip to Reaction Details section. |
| Blood Product: |
| Date of Transfusion:// UNKNOWN |
| Did the patient experience a transfusion adverse reaction? |
| If yes, provide information about the transfusion adverse reaction. |
| Type of transfusion adverse reaction: Allergic AHTR DHTR DSTR FNHTR |
| HTR TTI PTP TACO TAD TA-GVHD TRALI UNKNOWN |
| OTHER Specify |
| Reaction Details |
| *Date reaction occurred:// *Time reaction occurred:: Time unknown |
| *Facility location where patient was transfused: |
| *Is this reaction associated with an incident? Yes No If Yes, Incident #: |
| After recognition of the transfusion reaction, was the current transfusion: |
| Continued Stopped and restarted Stopped indefinitely |
| Investigation Results |
| * Hypotensive transfusion reaction |
| *Case Definition |
| Check all that occurred during or within 1 hour of cessation of transfusion: |
| All other adverse reactions presenting with hypotension are excluded. |
| Hypotension |
| Check all that apply: |
| Hypotension occurs, does not meet the criteria above. Other, more specific reaction definitions do not apply. |
| None of the above |
| Continued >> |



| Investigation Results (d | ontinued) | | | | | | | |
|--|---|--|--|--|--|--|--|--|
| Other signs and symptoms: | (check all that apply) | | | | | | | |
| Generalized: | Chills/rigors | Fever | Nausea/vomiting | | | | | |
| Cardiovascular: | Shock | | | | | | | |
| Cutaneous: | Edema | Flushing | Jaundice | | | | | |
| Odta 100doi | Other rash | Pruritus (itching) | Urticaria (hives) | | | | | |
| Hemolysis/Hemorrhage: | Disseminated intra | vascular coagulation | Hemoglobinemia | | | | | |
| | | Positive antibody screen | | | | | | |
| Pain: | Abdominal pain | Abdominal pain Back pain Flank pain Infusion site pain | | | | | | |
| Renal: | Hematuria | Hemoglobinuria | Oliguria Oliguria | | | | | |
| Respiratory: | Bilateral infiltrates | | nchospasm Cough | | | | | |
| | Hypoxemia | Hypoxemia Shortness of breath | | | | | | |
| Other: (specify) | | | | | | | | |
| | | | | | | | | |
| <u>*Severity</u> | | | | | | | | |
| | | • , | definitions listed in protocol) | | | | | |
| Symptomatic trea | | | prolonged hospitalization | | | | | |
| Life-threatening r | | | and/or incapacitation | | | | | |
| Congenital anom | aly or birth defect(s) of | the fetus | Death | | | | | |
| Other medically i | mportant conditions | | Unknown or not stated | | | | | |
| <u>*Imputability</u> | | | | | | | | |
| Which best describes the | e relationship between | the transfusion and the | reaction? | | | | | |
| The patient has no | other conditions that c | ould explain hypotensio | n. | | | | | |
| | tential causes present | that could explain hypot | ension, but transfusion is the most likely | | | | | |
| cause. | | | | | | | | |
| | | n hypotension are prese | | | | | | |
| | | | but transfusion cannot be excluded. | | | | | |
| | • | | e other than the transfusion. | | | | | |
| The relationship be | tween the adverse rea | action and the transfusio | n is unknown or not stated. | | | | | |
| How did the patient resp | ond the cessation of tra | ansfusion and supportive | e treatment? | | | | | |
| Responds rapidly (i.e., within 10 minutes) to cessation of transfusion and supportive treatment. | | | | | | | | |
| The patient does not respond rapidly to cessation of transfusion and supportive treatment. | | | | | | | | |
| Did the transfusion occu | at your facility? | YES NO | | | | | | |
| When did the reaction od | cur in relation to the tra | ansfusion? | | | | | | |
| Occurs less than 2 | L5 minutes after the sta | art of the transfusion. | | | | | | |
| Onset is between | Occurs less than 15 minutes after the start of the transfusion. Onset is between 15 minutes after start and 1 hour after cessation of transfusion. | | | | | | | |
| | | | Continued >> | | | | | |



| Investigation Results (continued) |
|--|
| Designations for case definition, severity, and imputability will be automatically assigned in the NHSN application based on responses in the corresponding investigation results section above. |
| Do you agree with the case definition designation? |
| Please indicate your designation |
| Do you agree with the severity designation? |
| Please indicate your designation |
| Do you agree with the imputability designation? |
| Please indicate your designation |
| Additional Information |
| |
| |
| |
| Patient Treatment |
| *Did the patient receive treatment for the transfusion reaction? YES NO UNKNOWN |
| If yes, select treatment(s): Medication (Select the type of medication) |
| Medication (Select the type of medication) |
| Antipyretics Antihistamines Inotropes/Vasopressors Bronchodilator Diuretics |
| Intravenous |
| Immunoglobulin |
| Antithymocyte globulin Cyclosporin H1 receptor blockers Other |
| Volume resuscitation (Intravenous colloids or crystalloids) |
| Respiratory support (Select the type of support) |
| Mechanical ventilation Noninvasive ventilation Oxygen |
| Renal replacement therapy (Select the type of therapy) |
| |
| Hemodialysis Peritoneal Continuous Veno-Venous Hemofiltration |
| Phlebotomy |
| |
| Other Specify: |
| |
| Outcome |
| *Outcome: Death Major or long-term sequelae sequelae Not determined |
| Date of Death: / / |
| ^*If recipient died, relationship of transfusion to death: |
| Definite Probable Possible Doubtful Ruled Out Not determined |



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|---------------------------|-----|----|------------------|--------|
| Cause of death: | | | | |
| Was an autopsy performed? | Yes | No | | |
| | | | Continu | ıed >> |



| Component Details (Use worksheet on page 4 for additional units.) | | | | | | | | | |
|---|--|-------------------------------------|--------------------|-------------------------------|---------|-----------------|------------|-------------------------|--|
| | ular unit implicated in | | ble for) the adver | se reaction? | Yes | | No [| N/A | |
| Transfusion Start and End Date/Time | *Component code (check system used) | Amount transfused at reaction onset | Unit number | *Unit expiration Date/Time | *Blood | group | of unit | Implic ated Unit? | |
| ^IMPLICATED UNIT | | | | | | | | | |
| : | ISBT-128 Codabar | Entire unit Partial unitmL | | | A | A+ AB- O+ | B- AB+ | Y | |
| | ISBT-128 Codabar | Entire unit Partial unitmL | | | A[| A+ AB- O+ | B- AB+ N/A | N | |
| | ISBT-128 Codabar | Entire unit Partial unitmL | | | A- B+ A | A+ AB- O+ | B- AB+ N/A | N | |
| Custom Fields | | | | | | | | | |
| Label | | | Label | | | | | | |
| | | <u> </u> | | | | | | | |
| Comments | | | | | | | | | |
| | | | | | | | | | |
| | | | | | | | | | |



Hemovigilance Module Additional Worksheet

| | story |
|--|--|
| (part 1) List the pa | ient's admitting diagnosis. (Use ICD-10 Diagnostic codes/descriptions) |
| Code: | Description: |
| Code: | |
| (part 2) List the pat | ent's underlying indication for transfusion. (Use ICD-10 Diagnostic codes/descriptions) |
| Code: | Description: |
| Code: | Description: |
| Code: | |
| Code: | |
| Code: | |
| Code: | |
| | ient's comorbid conditions at the time of the transfusion related to the |
| adverse reaction. (| Jse ICD-10 Diagnostic codes/descriptions) |
| adverse reaction. (| Jse ICD-10 Diagnostic codes/descriptions) |
| | Jse ICD-10 Diagnostic codes/descriptions) NONE Description: |
| Code: | Jse ICD-10 Diagnostic codes/descriptions) NONE Description: Description: |
| Code: Code: | Jse ICD-10 Diagnostic codes/descriptions) Description: Description: Description: |
| Code: Code: Code: | Description: Description: Description: Description: Description: |
| Code: Code: | Description: Description: Description: Description: Description: Description: Description: |
| Code: Code: Code: Code: Code: Code: (part 4) List the pa | Description: |
| Code: Code: Code: Code: Code: Code: (part 4) List the partocedures to be possible. | Description: Description: |
| Code: Code: Code: Code: Code: Code: Code: Procedures to be portional procedure codes/displayed. | Description: |
| Code: Code: Code: Code: Code: (part 4) List the partocedures to be perfocedure codes/de Code: | Description: |
| Code: Code: Code: Code: Code: Code: (part 4) List the partocedures to be performed by the performance by the performa | Description: |
| Code: Code: Code: Code: Code: Code: (part 4) List the partocedures to be porcedure codes/dictional code: Code: Code: Code: Code: Code: Code: | Description: |



Hemovigilance Module Additional Worksheet

| Transfusion History |
|--|
| Has the patient received a previous transfusion? YES NO |
| **If yes, provide information about the transfusion event. If not, skip to Reaction Details section. |
| Blood Product: WB RBC Platelet Plasma Cryoprecipitate Granulocyte |
| Date of Transfusion:// UNKNOWN |
| Did the patient experience a transfusion adverse reaction? YES NO |
| If yes, provide information about the transfusion adverse reaction. |
| Type of transfusion adverse reaction: Allergic AHTR DHTR DSTR FNHTR |
| ☐ HTR ☐ TTI ☐ PTP ☐ TACO ☐ TAD ☐ TA-GVHD ☐ TRALI ☐ UNKNOWN |
| OTHER Specify |
| |
| Has the patient received a previous transfusion? YES NO |
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| |
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| Date of Transfusion://UNKNOWN |
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| ☐ HTR ☐ TTI ☐ PTP ☐ TACO ☐ TAD ☐ TA-GVHD ☐ TRALI ☐ UNKNOWN |
| OTHER Specify |
| |



Hemovigilance Module Additional Worksheet

| Component Details | | | | | | | |
|-------------------------------------|--|--|----------------------------|----------------------------------|------------------------------------|-------------------------|--|
| *Was a particul | ar unit implicated in (| (i.e., responsi | ible for) the adverse read | ction? Y | es No N | I/A | |
| Transfusion Start and End Date/Time | *Component code (check system used) | Amount transfused at reaction onset | Unit number | *Unit expiration Date/Time | *Blood group of unit | Implic ated Unit? | |
| | ISBT-128 Codabar | Entire unit Partial unit mL | | <i>l 1</i> | A- A+ B- B AB- AB+ O- O+ N/A | N | |
| | ISBT-128 Codabar | Entire unit Partial unit mL | | | A- A+ B- B- AB- AB+ O- O+ N/A | N | |
| | ISBT-128 Codabar | Entire unit Partial unit mL | | | A- A+ B- B- AB- AB+ O- O+ N/A | N | |
| | ISBT-128 Codabar | Entire unit Partial unitmL | | | A- A+ B- B- AB- AB+ O- O+ N/A | N | |
| | ISBT-128 Codabar | Entire unit Partial unit mL | | <i>I. I.</i> | A- A+ B- B- AB- AB+ O- O+ N/A | N | |
| | ISBT-128 Codabar | Entire unit Partial unitmL | | | A- A+ B- B AB- AB+ O- O+ N/A | N | |
| | SBT-128 | Entire | | | A+ B- | IN | |

