



Sutter Roseville Medical Center

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Owner: *Nadera Poirier: Supervisor,  
Laboratory Analytic*Policy Area: *Lab - Transfusion Service*

References:

Applicability: *Sutter Roseville Medical Center*

## Routine Neonatal Transfusion Protocol

### PURPOSE

To provide instruction on required pre-transfusion testing and appropriate selection of blood components for neonatal patients (<4 months of age).

### POLICY

- Neonate pre-transfusion specimen acceptability follows standard collection and labeling requirements. See *Acceptability Criteria for and Labeling of Transfusion Services Specimens SOP*.
  - Specimen must be **peripheral 1.5 mL minimum EDTA** sample
  - Cord blood specimens may NOT be used as a primary or confirmatory ABORh
  - Heel stick specimens are unacceptable for pre-transfusion testing
- Two adult-sized RBC units meeting neonate selection requirements (see *Product Selection*) will be kept in stock through standing order (every 3-4 days) at all times. Units will be marked with the last date they may be used for neonatal transfusion and rotated into adult inventory past that date.
  - Adult units will be issued unless pediatric units are specifically requested by the clinician
  - Pediatric aliquots must be ordered as such from the blood supplier
- Neonate blood filter is to be attached and issued with each blood product. Contact NICU for additional filters, as needed.
- The EDTA tubes for drawing the pre-transfusion and confirmation testing along with job aid will be sent to NICU when order for transfusion testing has been placed or requested by the NICU nurse.
- **Exchange Transfusions:**
  - Product with final hematocrit of 45-60% will be manufactured at the blood supplier upon request using selection criteria noted below
  - Total volume of final product will range from 370-420 mL, containing approximately 220 mL RBCs and 150-175 mL of plasma
  - Units may not be equilibrated to 1-6°C prior to transport due to STAT need and abbreviated expiration of 24 hours from manufacture
  - Exchange transfusions require a blood warmer: add test **PI** and result with ETC **WARM** (Blood Warmer Required)
  - If **intrauterine** exchange transfusion is requested, perform all routine pre-transfusion testing on the *mother's XM accession* and complete AHG crossmatch.
    - Add **BBC**: "Intrauterine exchange transfusion"
    - Add **PI**: "Intrauterine exchange transfusion. Blood warmer required."

# TESTING FREQUENCY/ORDERS

| IF neonate is:                             | THEN pre-transfusion testing is performed:  |
|--|---|
| < 4 months, during current admission       | On the first sample, NOT repeated during admission (unless plasma is needed for testing and no is longer available) |
| < 4 months, discharged and readmitted      | On new sample from readmission, NOT repeated during new admission   |
| ≥ 4 months, regardless of admission status | Every 3 days  |

| NEONATAL ORDER CODE | ADULT EQUIVALENT | COMMENTS   |
|---------------------|------------------|--|
| INWU                | XM/TS            | <ul style="list-style-type: none"> <li>• Required for first order of <b>ANY</b> blood product</li> <li>• Includes ABORh, antibody screen, and IgG DAT</li> <li>• To be performed according to testing frequency noted above</li> <li>• Can be ordered with or without volume of product requested</li> </ul> |
| TINF                | ADDXM            | <ul style="list-style-type: none"> <li>• Routine request for additional infant RBCs</li> <li>• Result UO and HX, then allocate on original INWU accession</li> </ul>   |

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# PRODUCT SELECTION

| COMPONENT                        | SELECTION   | COMMENTS   |            |   |             |   |       |   |       |                    |    |  |
|----------------------------------|---|--|------------|---|-------------|---|-------|---|-------|--------------------|----|--|
| <b>Red Blood Cells (routine)</b> | <ul style="list-style-type: none"> <li>O negative</li> <li>Leukocyte reduced</li> <li>AS-1 or AS-3</li> <li>&lt;7 days post-irradiation</li> <li>Irradiated</li> <li>CMV negative</li> <li>Antigen negative, if indicated</li> </ul>  | <ul style="list-style-type: none"> <li>O negative <b>only</b></li> <li>Consult pathologist if appropriate product unavailable from blood supplier</li> </ul>   |            |   |             |   |       |   |       |                    |    |  |
| <b>Platelets</b>                 | <ul style="list-style-type: none"> <li>Plasma compatible (select in order listed)</li> </ul> <table border="1"> <thead> <tr> <th>NEONATE ABO:</th> <th>SELECTION:</th> </tr> </thead> <tbody> <tr> <td>O</td> <td>O, A, B, AB</td> </tr> <tr> <td>A</td> <td>A, AB</td> </tr> <tr> <td>B</td> <td>B, AB</td> </tr> <tr> <td>AB, INVAL, Unknown</td> <td>AB</td> </tr> </tbody> </table> <ul style="list-style-type: none"> <li>Leukocyte reduced</li> <li>Irradiated</li> <li>CMV negative</li> </ul> | NEONATE ABO:   | SELECTION: | O | O, A, B, AB | A | A, AB | B | B, AB | AB, INVAL, Unknown | AB | <ul style="list-style-type: none"> <li>Rh type not a consideration <b>except</b> in cases of Rh negative female, in which case Rh negative will be selected</li> <li>Consult pathologist if Rh positive must be given to Rh negative female to determine is Rhogam should be administered</li> </ul> |
| NEONATE ABO:                     | SELECTION:  |  |            |   |             |   |       |   |       |                    |    |  |
| O                                | O, A, B, AB   |  |            |   |             |   |       |   |       |                    |    |  |
| A                                | A, AB   |  |            |   |             |   |       |   |       |                    |    |  |
| B                                | B, AB   |  |            |   |             |   |       |   |       |                    |    |  |
| AB, INVAL, Unknown               | AB  |  |            |   |             |   |       |   |       |                    |    |  |
| <b>Plasma</b>                    | AB only   | <ul style="list-style-type: none"> <li>Rh type not a consideration</li> </ul>  |            |   |             |   |       |   |       |                    |    |  |
| <b>Cryoprecipitate</b>           | ABO compatible  | <ul style="list-style-type: none"> <li>Rh type not a consideration</li> </ul>  |            |   |             |   |       |   |       |                    |    |  |
| <b>Whole Blood (exchange)</b>    | <p>RBC requirements:</p> <ul style="list-style-type: none"> <li>O negative</li> <li>Leukocyte reduced</li> <li>CPD or CPDA-1</li> <li>&lt; 3 days old (per nursing SOP)</li> <li>Irradiated</li> <li>CMV negative</li> <li>HgbS negative</li> <li>Antigen negative, if indicated</li> </ul> <p>Plasma requirement:</p> <ul style="list-style-type: none"> <li>Type AB collected in CPD or CPDA-1</li> </ul>   | <ul style="list-style-type: none"> <li>O negative <b>only</b></li> <li>Ordered reconstituted from blood supplier</li> <li>Good for only 24 hours</li> <li><b>DO NOT</b> order until decision to perform procedure at SRMC has been made</li> </ul> |            |   |             |   |       |   |       |                    |    |  |

|                    |  |  |
|--------------------|--|--|
| <b>Granulocyte</b> | <ul style="list-style-type: none"> <li>• O negative</li> <li>• Irradiated</li> <li>• CMV negative</li> <li>• Antigen negative, if indicated</li> </ul> |  |
|--------------------|--|--|

## CROSSMATCH COMPATIBILITY

| NEONATE RESULTS  | CROSSMATCH METHOD   | RESULTING XM   |
|--|---|--|
| <b>AS:</b> Negative<br><b>DIGG:</b> Negative   | Electronic Crossmatch   | Auto-populates grids upon saving                       |
| <b>AS:</b> Positive<br><b>ABID:</b> Non-clinically significant antibody, ANAPS, or low incidence antibody<br>(regardless of <b>DIGG</b> or <b>ELU</b> results) | AHG Crossmatch  | Result according to AHG crossmatch procedure performed |
| <b>AS:</b> Negative<br><b>DIGG:</b> Positive<br><b>ELU:</b> Non-clinically significant antibody, ANAPS, or low incidence antibody                              |   |  |
| <b>AS:</b> Positive<br><b>ABID:</b> Clinically significant antibody<br>(regardless of <b>DIGG</b> or <b>ELU</b> results)                                       | Neonatal No XM Protocol<br>AHG crossmatch not required,<br><b>MUST</b> provide antigen negative units | <b>XIS:</b> NRQ (~)<br><b>Interp:</b> NCMP (shift ~)   |
| <b>AS:</b> Negative<br><b>DIGG:</b> Positive<br><b>ELU:</b> Clinically significant antibody  |   |  |

## PROCEDURE

| Step: | Action:   |
|-------|---|
| 1.    | Perform history check on neonate and mother (if available), then access the INWU accession in <i>Blood Order Processing</i> and result the following fields: <ul style="list-style-type: none"> <li>• <b>EXX</b> (Crossmatch Expiration): Date neonate will be 4 months old               <ul style="list-style-type: none"> <li>◦ Specimen is to be expired upon patient discharge OR exhaustion of neonate plasma, if needed for AHG crossmatches</li> </ul> </li> <li>• <b>UO</b> (Units Ordered): Number requested or <i>HIDE</i> (if none)</li> <li>• <b>HX</b> (History Check): <i>YH</i> or <i>NH</i></li> <li>• Add/order appropriate confirmation testing based on patient history</li> <li>• Add <b>PB</b> and result with free text: "Infant 4 months old on MM/DD/YYYY"</li> <li>• Add <b>PB</b> and result with ETCs for required neonatal attributes: <b>BBIRR, BBCMVN</b></li> </ul> |
| 2.    | Perform automated testing on neonate's specimen.<br>Note: Transcribe ABORh results from Echo. Reverse type is not indicated in neonates and will be result as <i>ND</i> .   |

|    |   |
|----|---|
| 3. | Perform antibody ID if antibody screen demonstrates reactivity. <ul style="list-style-type: none"> <li>• If mother has known specificity and limited neonate plasma is available, PEG selected cells may be used to identify the antibody in the neonate's plasma</li> <li>• All clinically significant antibodies must be ruled out using neonate's current specimen</li> <li>• Result <b>ABI</b> with <b>dated free text</b> including mother's name and medical record number (if available): "<i>MM/DD/YYYY Maternal anti X. Mother's name: LAST, FIRST (MRN: ###)</i>"</li> <li>• Add <b>PI</b> indicating need for antigen negative units, if applicable based on ABID</li> </ul> |
| 4. | Add test <b>ELU</b> and perform an elution if: <ul style="list-style-type: none"> <li>• DAT is positive upon initial workup <i>OR</i></li> <li>• DAT strength has increased since previous peripheral specimen testing</li> </ul>   |
| 5. | Crossmatch and allocate units as needed once all testing had been completed using <i>Crossmatch Compatibility</i> section for guidance.   |
| 6. | Select <i>Save</i> .  |

## RELATED DOCUMENTS

Acceptability Criteria for and Labeling of Transfusion Services Specimens

All revision dates:

### Attachments

No Attachments

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