

Health PEI

BLOOD TRANSFUSION SERVICE LABORATORY

Queen Elizabeth Hospital
Charlottetown, PEI
Phone (902) 894-2300
Fax (902) 894-2415

Prince County Hospital
Summerside, PEI
Phone (902) 438-4280
Fax (902) 438-4281

Address for Non-PEI Residents Required

Name: _____

Street: _____ **Place Label Here**

City: _____ Prov./State _____

Postal Code/Zip: _____

Allergies: _____

ORDER SET

Intravenous Immunoglobulin (IVIG) Hematology – Adult

Patient Name: _____ Patient MRN: _____ DOB: YYYY/MON/DD

Items preceded by a **bullet** (•) are active orders. Items preceded by a **checkbox** (☐) are only to be carried out if checked.

• **Any change to indication, dose, duration or frequency requires a new order.**

Note: IVIG dose is calculated using the patient's DOSING BODY WEIGHT (DBW) for all indications. **If** patient height **under 152.4 cm**, Dosing Body Weight equals Actual Weight or pre-pregnancy weight. **If** patient height **over 152.4 cm**, use the DBW Calculator to obtain a clinically appropriate Dosing Body Weight. To obtain the DBW calculator, search "NS Health IVIG Dose Calculator" in an internet search engine.

| Actual Weight (kg): | | Pre-pregnancy Weight (kg – if applicable): | Height (cm): |
|--|--|---|--------------|
| Dosing Body Weight (kg – see note above): | | IVIG Rounded Dose (g): | Gender: |
| IgA Deficient Product Required: <input type="checkbox"/> Yes <input type="checkbox"/> No | Is this a repeat dose due to lack of expected response? <input type="checkbox"/> Yes <input type="checkbox"/> No | Intended Treatment Start Date (YYYY/MON/DD): | |
| Dosage and Duration of Therapy | | | |
| • Infuse _____ g/kg = _____ g daily for _____ days OR Infuse _____ g/kg = _____ g divided over _____ days | | | |
| • If indicated, repeat this regimen every _____ days for a total of _____ treatments | | | |
| Indicated Conditions | Prerequisites – checkboxes must be checked / completed as appropriate. Missing information will result in delays or denial of product. PATIENT MUST MEET THE FOLLOWING: | Dose | |
| <input type="checkbox"/> Immune Thrombocytopenia (ITP)* | <input type="checkbox"/> Major bleeding and platelets less than 50x10 ⁹ /L OR <input type="checkbox"/> Failed to respond to steroids after 3 or more days OR <input type="checkbox"/> To produce an increase in platelet count to a level considered safe | Acute: 1 g/kg/day for 1 or 2 consecutive days depending on response Chronic: 1 to 2 g/kg no more frequently than every 2 weeks | |
| <input type="checkbox"/> Pregnancy – Associated ITP* | <input type="checkbox"/> There is major bleeding OR <input type="checkbox"/> Platelet counts fall below 10x10 ⁹ /L anytime during pregnancy OR 10 to 30x10 ⁹ /L during second or third trimester OR <input type="checkbox"/> Rapid elevation of platelets required before delivery or any invasive procedure (e.g. amniocentesis) | 1 g/kg/day x 2 consecutive days (dosing body weight is based on the pre-pregnancy weight for determining IVIG dose) (no maximum dose) | |
| <input type="checkbox"/> Post-Transfusion Purpura (PTP)* | No prerequisites are required | 1 g/kg repeated if necessary | |
| <input type="checkbox"/> Fetal Alloimmune Thrombocytopenia (FAIT)* | <input type="checkbox"/> Mother has been found to have anti-platelet alloantibodies through a prior affected pregnancy or close family member (e.g. sister) with an affected pregnancy AND <input type="checkbox"/> Treatment is under the direction of a maternal fetal medicine centre | 1 to 2 g/kg/week throughout the pregnancy (dosing body weight is based on the pre-pregnancy weight for determining IVIG dose; disease severity also considered) (no maximum dose) | |

* May be considered URGENT if notified by ordering prescriber

Authorized Prescriber's Signature: _____ Reg. No.: _____

Prescriber's Name: _____ Date (YYYY/MON/DD): _____ Time: _____
Print

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| Possibly Indicated Conditions are approved for a 3 month period <u>only</u> at which time a clinical outcome questionnaire must be provided for the patient to continue treatment | | |
|---|--|---------------------------------|
| Possibly Indicated Conditions | Prerequisites – checkboxes must be completed PATIENT MUST MEET THE FOLLOWING: | Dose |
| <input type="checkbox"/> Acquired Hemophilia with Factor VIII Inhibitor* | <input type="checkbox"/> Order must be in consultation with a Hematologist Name: _____ | 2 g/kg divided over 2 to 5 days |
| <input type="checkbox"/> Factor XIII Inhibitor* | <input type="checkbox"/> Order must be in consultation with a Hematologist Name: _____ | 2 g/kg divided over 2 to 5 days |
| <input type="checkbox"/> Secondary Immune Deficiency (SID) | <input type="checkbox"/> Order must be in consultation with a Hematologist Name: _____ | 0.4 g/kg every 3 to 4 weeks |
| <input type="checkbox"/> Warm Autoimmune Hemolytic Anemia | Resistant to steroids and symptomatic anemia | Up to 2 g/kg |
| <input type="checkbox"/> Hemophagocytic Lymphohistiocytosis (HLH)* | <input type="checkbox"/> Order must be in consultation with a Rheumatologist, Hematologist or General Internist Name: _____ | 2 g/kg divided over 2 to 5 days |

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Prescriber's Name: _____ Date (YYYY/MON/DD): _____ Time: _____
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