

BC Provincial Blood Coordinating Office

A program of the Provincial Health Services Authority

Intravenous Immune Globulin (IVIG) Reaction Chart

All patients should receive information on potential transfusion reactions and how to report a suspected transfusion reaction.

Mild Transient Reactions/Side Effects

- Common mild signs and symptoms that resolve if the flow rate is reduced and/or the patient is medicated.
- Most likely to occur in first 30 to 60 minutes of infusion.
- **Do Not Report** to Transfusion Medicine Services/Laboratory (TMS/Lab). No patient samples required.
- If patient's condition does not improve despite decreasing the rate or medication: **Stop the infusion** and refer to the section on **Acute IVIG Reactions** on this chart.

Signs and Symptoms	Actions	Comments	
 Headache (mild to moderate) Flushing Muscle aches Shivering Nausea Localized Urticaria Pruritus Anxiety Light-headed Dizziness or irritability 	 Decrease the flow rate until a rate at which the symptoms subside is reached Consult physician Medicate appropriately Apply relevant patient comfort measures Frequent vital signs Document as per facility policy Do not report to TMS/Lab 	For subsequent treatments consider: premedication advancing infusion at a slower rate	
 Pain at intravenous site 	Use of a large vein for the infusion may avoid pain at the intravenous site.		

Acute IVIG Reactions - Within 24 hours of Transfusion

- Stop the infusion; Consult Physician; Report to Transfusion Medical Services / Laboratory (TMS/Lab)
- Document as per facility policy.
- Return any unopened product to TMS/Lab.

*Note: **Fever** is defined as an oral temperature ≥ 38°C **AND** ≥ 1°C rise in oral temperature above pre-transfusion baseline

Signs and Symptoms	Reaction	Actions	Comments
Anxiety; fever*;chills; rigors; non localized urticaria/rash; itchiness; flushing; nausea; vomiting; chest, back or abdominal pain; tachycardia; hypotension or hypertension OR any mild reactions/side effects listed above that do not respond to rate decrease or medication.	Moderate to Severe	 Contact the physician for assessment and symptomatic treatment. Comfort measures as applicable. Do not restart without a physician's order. Reassess patient frequently. 	For subsequent treatments consider: premedication increasing infusion at a slower rate changing brand of IVIG the use of SCIG
Facial and/or tongue swelling; difficulty in swallowing; chest tightness; airway edema; dyspnea; hypotension; shock; tachycardia; nausea; vomiting; widespread urticaria/rash (>2/3 body), anxiety; fever*.	Anaphylaxis	 Do not restart. Contact physician for assessment and symptomatic treatment. May require epinephrine. Comfort measures as applicable 	 May be reaction to IgA in an IgA deficient patient. For subsequent treatments consider: changing brand of IVIG reassessing the need for IVIG. consulting an immunologist. measuring IgA levels
Fever*, back pain, dyspnea, red / brown urine.	Acute Hemolysis	 Do not restart. Contact physician for assessment and symptomatic treatment. Send to TMS/Lab: 2 EDTA vials. First voided post-reaction urine sample for routine urinalysis. 	 Due to antibodies in IVIG directed against a patient's red blood cells. Blood group A, B or AB adult patients receiving more than 50g of IVIG or pediatric patients receiving 1g/kg or more are at an increased risk of hemolysis.



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Delayed IVIG Reactions - Greater than 24 hours post Transfusion

- Consult Physician; Report to TMS/Lab; Send Required Patient Samples.
- Symptomatic treatment as ordered by physician. Comfort measures as applicable.
- Document as per facility policy
- Patient may be readmitted to hospital at a later date due to delayed reaction.
- For subsequent administrations consider:
 - premedicating appropriately
 - advancing the infusion at a slower rate
 - reducing maximum infusion rate
 - changing brand of IVIG
 - the use of SCIG

*Note: Fever is defined as an oral temperature ≥ 38°C AND ≥ 1°C rise in oral temperature above pre-transfusion baseline

Signs and Symptoms	Reaction Type	Comments
Prolonged and severe headache that is unresolved by medication	Delayed Headache	 Medicate as ordered as soon as first signs of headache occur. For subsequent IVIG administration, physician may consider prehydration with saline.
Severe and incapacitating headache with nuchal rigidity, drowsiness, fever*, lethargy, photophobia, painful eye movements, nausea, vomiting, diarrhea, pharyngitis, deterioration of mental status	Aseptic Meningitis	 Presents up to 72 hours post transfusion. Usually resolves spontaneously in 1-2 days. Previous history of migraine headaches may be a risk factor. Pre/post medication with corticosteroids/anti-migraine medication may help to prevent/reduce incidence.
Fever*, back pain, dyspnea, red / brown urine, falling haemoglobin, jaundice, unexpected/unexplained fatigue	Delayed Hemolysis	 Occurring within 10 days post transfusion. Often due to antibodies in IVIG directed against a patient's red blood cells. Blood group A, B or AB adult patients receiving more than 50g of IVIG or pediatric patients receiving 1g/kg or more are at an increased risk of hemolysis.
Peripheral edema, periorbital edema, urination changes, increased serum creatinine, hypertension, back pain, flank pain, blood in urine.	Acute Renal Failure	 Predisposing factors: age>65; diabetes mellitus; pre-existing renal insufficiency. Usually seen with products containing sucrose (none currently licensed in Canada).
Symptoms related to: myocardial infarction; transient ischemic attack; stroke; deep vein thrombosis.	Thrombo- embolic events	 Causative relationship not clearly understood. Possibly related to increases in blood viscosity. Risk factors include: arthrosclerosis; advanced age; previous thrombotic event; clotting disorder; hypertension; diabetes; obesity; immobility.
Variable as per specific infectious disease.	Transfusion Transmitted Infections	 Diagnosed through transmissible disease tests. No reported cases of HIV or HBV. No reported HCV since 1995. Effective viral reduction measures. Prion (vCJD) transmission theoretical risk.