

Intravenous Immune Globulin (IVIG) Order Form

Fax order to: 517-364-8448

Complete and return to UMH- Sparrow Infusion Center. See next page for accepted indications and dosing. For alternate dosing or indications, literature supporting IVIG use is required and subject to approval by pharmacist.

Patient Name (Infusion Center Use Only) UMHSparrow MRN			kg Height	cm
1. Diagnosis IVIG Treatment Indication: Your patient will be scheduled upon confirmation of insurance		ICD-10 diagnosis co	de:	
2. Pre-Medications □ Acetaminophen (Tylenol [®]) mg orally □ Loratadine (Claritin [®]) 10 mg orally x 1 dose □ Other:	🗆 Dij	phenhydramine (Benadryl [®]) _ phenhydramine (Benadryl [®]) _	mg orall mg IV x	y x 1 dose 1 dose
3. IVIG Product Selection				
 Gammagard[®] 10% - Formulary product Gammagard[®] S/D Powder 10% - Patient 	must be previously	stabilized on this product or hav	e contraindication to	o formulary product.
√ IV/IG arams/ka □	One time only Daily for Every	days weeks for □ on	e year □	doses
For patients ≥18 years, IVIG dose will be determined based determined based on (1) Actual BW for patients < 60 kg; (2)				
5. ANSWER REQUIRED: Patient may receive IVIG at *Accelerated Rate after patient's first IVIG infusion. YES NO *Accelerated Rate Policy a. Use ideal Body weight to determine rate b. Start at rate of 0.5 ml/kg/hr. c. After 30 min increase to 1 ml/kg/hr. d. After 30 min increase to 2 ml/kg/hr. e. After 30 min increase to 4 ml/kg/hr. f. After 30 min increase to 5 ml/kg/hr. f. After 30 min increase to 5 ml/kg/hr. for remaining volume 6. Labs Other:				
Physician Signature X			Date	Time
Physician Printed Name				
Office Contact			Office Phone	
Pharmacy Use Only: 1. Circle weight to use: Ideal BWkg 2. Dosage calculation:grams/kg x 3. Round to nearest 10 grams (if >20 grams) = □ IVIG use NOT approved □ Patient approved for IVIG use:	kg = grams		UMH-Sparrow IVI Infusion Center	IG Orders –
Order expires Rph Signature		e, and frequency)	PH-2000.10 (rev.	AS 07/24)

Copied from: Sparrow Health System IVIG Medication Use Policy (updated 5/2023) Table 1. Accepted Indications and Dosing (FDA Approved)			
Replacement therapy in primary immune deficiency syndrome	0.4 g/kg every 3 to 4 weeks with dose adjustments to maintain trough IgG level of 500 mg/dL and/or reduction in the incidence of infection		
Immune Thrombocytopenic	Acute ITP: 1 g/kg once. Dose may be repeated in 48 hours if no		
Purpura (ITP)-Adult	response. In the situation of life-threatening bleeding, the second		
	dose may be repeated at 24 hours.		
Criteria for use of IVIG			
Platelet count less than 30,000 AND severe bleeding	Chronic ITP: 0.4 g/kg every 3 to 4 weeks		
Platelet count less than 10,000 with no minimal bleeding			
Platelet count less than 50,000 and pending surgery			
Immune Thrombocytopenic	Acute ITP: 0.8-1 g/kg once		
Purpura (ITP)-Pediatrics	Chronic ITP: 0.4 g/kg every 3 to 4 weeks		
Criteria for use of IVIG			
Platelet count less than 20,000 AND significant bleeding			
Platelet count less than 10,000 with no or minimal bleeding			
Prophylaxis of infection and	0.4 g/kg every 3 to 4 weeks		
treatment of hypo-gammaglobulinemia in B-Cell CLL			
Kawasaki Disease	2 g/kg once		
CIDP	Loading dose: 2 g/kg over 2 to 5 days		
	Maintenance: 1 g/kg over 1 to 2 days every 3 weeks		
Multifocal motor neuropathy	2 g/kg over 5 days		

Table 2. Other Accepted Indications and Dosing (Non-FDA Approved)			
Indication	Dosing		
Anemia, hemolytic neonatal	0.5-1 g/kg once. Dose may be repeated in 12 hours.		
Dermatomyositis	2 g/kg over 2 days every 4 weeks for 3		
	doses in severe active illness for whom other interventions have		
	been unsuccessful or intolerable		
Guillain-Barre or Acute inflammatory demyelinating polyneuropathy (AIDP)	2 g/kg over 5 days		
Paraneoplastic opsoclonus-myoclonus-ataxia associated with neuroblastoma	2 g/kg over 1 to 2 days		
Severe pseudomembranous colitis	0.4 g/kg for 1 or 2 doses		
Stiff-person (Moersch-Woltmann) syndrome	2 g/kg over 2-5 days every month		
Systemic lupus erythematosus	2 g/kg over 2 to 5 days every month in severe active illness for whom other interventions have been unsuccessful or intolerable		
Thrombocytopenia, fetal-neonatal	1 g/kg every week (use actual body weight of		
alloimmune/autoimmune	mother) from week 20 of pregnancy until delivery		
Streptococcal bacteremia, toxic shock syndrome	1 g/kg on day 1, 0.5 g/kg on days 2 and 3		
Transplantation, acute humoral rejection, renal	Used in combination with plasmapheresis and rituximab. Daily plasmapheresis followed by 0.1 g/kg for 4 doses then IVIG 2 g/kg every 3 weeks for 4 doses		
Transplantation, solid organ, CMV infection/prevention	0.4 g/kg every 3 weeks		
Secondary hypogammaglobulinemia (Multiple myeloma, other lympho-proliferative disorders, HIV- infected children, etc. to prevent serious infections) Documentation of low IgG levels required	0.4 g/kg every 3 to 4 weeks with dose adjustments to maintain trough IgG level of 500 mg/dL and/or reduction in the incidence of infection		
Myasthenia gravis	Loading dose: 2 g/kg over 2 to 5 days, Maintenance: 0.4 g/kg every 3-6 weeks		
	Note: When IVIG supply is sufficient to meet the health-system's demand, maintenance doses can be increased to 1 g/kg every 3-4 weeks. When IVIG supply is insufficient to meet the health-system's demand maintenance doses will be limited to 0.4 g/kg every 3-4 weeks.		
Multisystem Inflammatory Syndrome in children (MIS-C) associated with SARS-CoV-2	1-2 gm/kg once. Dose may be repeated in 24-36 hours.		