

## Intravenous Immunoglobulin/Subcutaneous Immunoglobulin Infusion

### Prior Authorization Request

Date \_\_\_\_\_

Member information		
Member name (print)	SMID	Date of birth (month/day/year)
Provider information		
Provider name (print)	Telephone number	Fax number

#### Medication requested

Intravenous immunoglobulin (IVIG):

- Bivigam™     
  Carimune®     
  Flebogamma®     
  Gammagard®     
  Gammaked®  
 Gammaplex®     
  Gamunex®     
  Octagam®     
  Privigen®

Subcutaneous immunoglobulin (SQIG):

- Gammagard®     
  Gammaked®     
  Gamunex®     
  Hizentra®

#### Indicate the diagnosis

- Primary immunodeficiency syndrome
- Include medical records with the following:
    - History and physical examination, documenting the severity of the condition, including frequency and severity of infections
    - Clinical deficiency of immunity as evidenced by one of the following:
      - Laboratory results supporting the diagnosis for which immune globulin is requested
      - Documented failure to produce antibodies to specific antigens
      - History of significant recurrent infections
- Idiopathic thrombocytopenic purpura (ITP)
- Include medical records with the following:
    - Documentation of diagnosis of idiopathic thrombocytopenic purpura (ITP)
    - Documentation of platelet count less than  $50 \times 10^9/L$
- Chronic inflammatory demyelinating polyneuropathy
- Initial request: The immune globulin dose does not exceed 2,000 mg/kg given over 2 to 5 consecutive days administered in up to 3 monthly infusions
- Include medical records confirming any checked boxes below:
    - Member has had symptoms present for at least 2 months
    - Symptomatic polyradiculoneuropathy as indicated by progressive or relapsing motor or sensory impairment of more than one limb
- Electrophysiologic findings:
- Partial conduction block of  $\geq 1$  motor nerve
  - Reduced conduction velocity of  $\geq 2$  motor nerves
  - Prolonged distal latency of  $\geq 2$  motor nerves
  - Prolonged F-wave latencies of  $\geq 2$  motor nerves or the absence of F waves
- Lumbar puncture:
- White blood cell count  $<10/mm^3$
  - Elevated CSF protein

Continuation request:

– Include medical records confirming any checked boxes below:

- Documentation of positive clinical response to therapy as measured by an objective scale (e.g. Rankin, Modified Rankin, Medical Research Council [MRC] scale)
- For long-term treatment, documentation that the dose has been periodically reduced or the treatment withdrawn, and the effects measured
- Concomitant immunomodulator therapy (e.g. azathioprine, cyclosporine, methotrexate, prednisone), unless contraindicated
- Immune globulin dose does not exceed 2,000 mg/kg given over 2 to 5 consecutive days. IVIG administration may be repeated monthly as needed to prevent exacerbation. Dosing interval may need to be adjusted in patients with severe comorbidities

Other \_\_\_\_\_

Initial request:

– Include medical records confirming any checked boxes below:

- Medical records documenting the severity of the diagnosis, including frequency and severity of infections
- Laboratory results supporting the diagnosis for which immune globulin is requested
- Documented failure to produce antibodies to specific antigens

Continuation request:

– Include medical records confirming any checked boxes below:

- Medical records indicating objective response to therapy, including decreasing frequency, and severity of infections
- Documentation of titration to the minimum dose and frequency to maintain a sustained clinical effect
- Serum immunoglobulin levels prior to immune globulin therapy and most recent serum IgG trough levels

HCPCS/J-code	
ICD code	
Strength requested	
Dosing schedule/frequency	
Duration of therapy	
Weight of member	
Administration site	<input type="checkbox"/> Provider office <input type="checkbox"/> Home infusion _____ <input type="checkbox"/> Outpatient infusion center _____ <input type="checkbox"/> Other _____

Comments \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

Provider signature

Date

**Pre-service decisions:** Initial review is received and a coverage determination is made within fourteen (14) calendar days of receipt of request. The member and/or provider are notified in writing of a denial decision within fourteen (14) calendar days of receipt of the request.

**Urgent pre-service decisions:** Initial review is received and a coverage determination is made within seventy-two (72) hours of receipt of request.

**Mail or fax form to:** Security Health Plan  
Health Services Department  
PO Box 8000  
Marshfield, WI 54449-8000  
Fax 715.221.9918

**Marshfield Clinic providers route to:**  
Health Services Department  
Routing location, SHP

**If you have any questions, please contact Provider Assistance at 1.800.472.2363.**