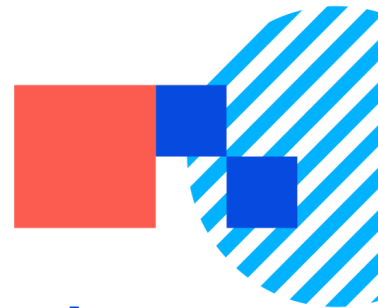




Immune Globulins Prior Authorization Drug List A

Drug(s) Applied:	Gamunex-C Flebogamma Dif, Gammagard, Gammaplex
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Criteria:

Drug(s) Applied will be approved when the requested medication is being used for a compendia-supported indication and all of the following criteria are met:

I. Initial Therapy Criteria

A. Primary Immunodeficiency as indicated by chart notes within past 120 days

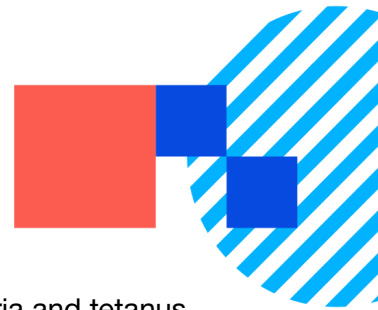
1. ONE of the following:

- a) Total IgG less than 200mg/dL at baseline prior to immune globulin therapy **or**
- b) Patient has abnormal Bruton tyrosine kinase (BTK) gene/absence of BTK protein (X-linked agammaglobulinemia) **or**
- c) Patient has an absence of B lymphocytes **or**
- d) ALL of the following:

(1) Patient has ONE of the following:

- (a) Selective IgG subclass deficiency [Defined as deficiency of 1 or more IgG subclasses (e.g., IgG1, IgG2, IgG3, or IgG4) by more than 2 standard deviations (SD) below age-specific mean, assessed on 2 separate occasions during infection free period] **and**
- (b) Specific antibody deficiency (SAD) with normal levels of both immunoglobulin and total IgG subclasses **and**
- (c) Hypogammaglobulinemia defined as total IgG less than 700 mg/dL OR more than 2 standard deviations below mean for the patient's age at baseline prior to immune globulin therapy **and**
- (d) Another primary immunodeficiency [e.g., Common variable immunodeficiency (CVID), X-linked immunodeficiency, severe combined immunodeficiency (SCID), combined immunodeficiency syndromes (e.g., Ataxia Telangiectasia (A-T), DiGeorge syndrome, Wiskott-Aldrich Syndrome)] **and**

- (2) Patient has a lack of response or inability to mount an adequate response to protein and/or polysaccharide antigens (e.g., inability



- to make IgG antibody against either diphtheria and tetanus toxoids, or pneumococcal polysaccharide vaccine, or both) **and**
- (3) Patient has evidence of recurrent, persistent, severe, difficult-to-treat infections (e.g., recurring otitis media, bronchiectasis, recurrent infections requiring IV antibiotics) despite aggressive prophylactic management and treatment with antibiotics **and**
 - 2. If requested agent is Flebogamma Dif, Gammagard, Gammaplex:
 - a) Patient has tried and had an inadequate response to Gamunex-C for at least 12 consecutive weeks **or**
 - b) Patient has an intolerance or FDA labeled contraindication to Gamunex-C **and**
 - 3. Prescriber has submitted support of use of SCIG over preferred IVIG with ONE of the following after at least 6 months of consistent IVIG therapy:
 - a) History of multiple upper respiratory symptoms or malaise during the week prior to the next infusion with the patient being adherent to an infusion schedule of every 3 to 4 weeks **or**
 - b) IgG trough level is below the patient's individualized goal IgG trough level (e.g. IgG trough level > 500mg/dL) with the patient being adherent to an infusion schedule of every 3 to 4 weeks and no infections during the time the trough level was drawn **and**
 - 4. Patient does NOT have any FDA labeled contraindications to the requested agent, including but not limited to IgA deficiency with antibodies against IgA and a history of hypersensitivity **and**
 - 5. Prescriber is a specialist or has consulted with a specialist in the area of the patient's diagnosis (e.g., allergy, immunology)

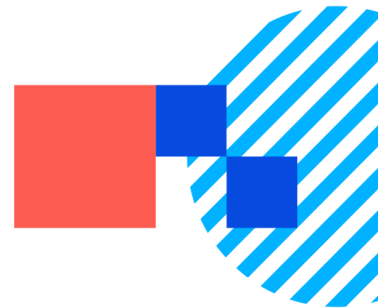
Approval Duration:

Gamunex-C: 12 months

Flebogamma, Gammagard, Gammaplex: 2 months

B. Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) as indicated by chart notes within past 120 days

- 1. Patient has progressive symptoms present for at least 2 months **and**
- 2. Patient has progressive or relapsing motor sensory impairment of more than one limb **and**
- 3. Patient has electrodiagnostic findings indicating at least ONE of the following are present:
 - a) Motor distal latency prolongation in 2 nerves



- b) Reduction of motor conduction velocity in 2 nerves
 - c) Prolongation of F-wave latency in 2 nerves
 - d) Absence of F-waves in at least 1 nerve
 - e) Partial motor conduction block of at least 1 motor nerve
 - f) Abnormal temporal dispersion in at least 2 nerves
 - g) Distal CMAP duration increase in at least 1 nerve **and**
- 4. Prescriber has submitted support of use of SCIG over preferred IVIG with ONE of the following after at least 6 months of consistent IVIG therapy:
 - a) History of multiple upper respiratory symptoms or malaise during the week prior to the next infusion with the patient being adherent to an infusion schedule of every 3 to 4 weeks **or**
 - b) IgG trough level is below the patient's individualized goal IgG trough level (e.g. IgG trough level > 500mg/dL) with the patient being adherent to an infusion schedule of every 3 to 4 weeks and no infections during the time the trough level was drawn **and**
- 5. If requested agent is Flebogamma Dif, Gammagard, Gammaplex:
 - a) Patient has tried and had an inadequate response to Gamunex-C for at least 12 consecutive weeks **or**
 - b) Patient has an intolerance or FDA labeled contraindication to Gamunex-C **and**
- 6. ONE of the following:
 - a) Patient has tried and had an inadequate response to oral or IV corticosteroid therapy **or**
 - b) Patient has an intolerance or FDA labeled contraindication to corticosteroid therapy **and**
- 7. Patient does NOT have any FDA labeled contraindications to the requested agent, including but not limited to IgA deficiency with antibodies against IgA and a history of hypersensitivity
- 8. Prescriber is a specialist or has consulted with a specialist in the area of the patient's diagnosis (e.g., neurology)

Approval Duration: Gamunex-C: 6 months

Flebogamma, Gammagard, Gammaplex: 2 months

C. Dermatomyositis/Polymyositis as indicated by chart notes within past 120 days

- 1. ONE of the following:
 - a) Patient has tried and had an inadequate response to a conventional therapy (e.g. prednisone, azathioprine, mycophenolate) **or**
 - b) Patient has an intolerance or FDA labeled contraindication to ALL conventional therapies (e.g. prednisone, azathioprine, mycophenolate) **and**



2. Prescriber has submitted support of use of SCIG over preferred IVIG with ONE of the following after at least 6 months of consistent IVIG therapy:
 - a) History of multiple upper respiratory symptoms or malaise during the week prior to the next infusion with the patient being adherent to an infusion schedule of every 3 to 4 weeks **or**
 - b) IgG trough level is below the patient's individualized goal IgG trough level (e.g. IgG trough level > 500mg/dL) with the patient being adherent to an infusion schedule of every 3 or 4 weeks and no infections during the time the trough level was drawn **and**
3. If requested agent is Flebogamma Dif, Gammagard, Gammaplex:
 - a) Patient has tried and had an inadequate response to Gamunex-C for at least 12 consecutive weeks **or**
 - b) Patient has an intolerance or FDA labeled contraindication to Gamunex-C **and**
4. Patient does NOT have any FDA labeled contraindications to the requested agent, including but not limited to IgA deficiency with antibodies against IgA and a history of hypersensitivity **and**
5. Prescriber is a specialist or has consulted with a specialist in the area of the patient's diagnosis (e.g., dermatology, immunology)

Approval Duration: Gamunex-C: 4 months

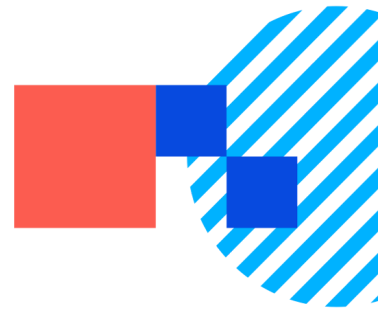
Flebogamma, Gammagard, Gammaplex: 2 months

II. Continued Therapy Criteria

A. All indications as indicated by chart notes within past 12 months

For Maryland Fully Insured groups only, chart notes not required.

1. Patient has been previously approved for the requested agent through the plan's Prior Authorization process or meets the initial therapy criteria above **and**
2. Chart notes indicate patient has been treated with multiple doses of the requested agent within the past 120 days and is a continuation of therapy (starting on samples is not approvable) **and**
3. If the patient has received more than 6 months of therapy, there is documented clinical benefit or disease stabilization since starting the requested agent (i.e., IgG level has improved from pre-treatment levels with the requested agent, reduction in the number and/or severity of difficult to treat infections) **and**
4. Patient does NOT have any FDA labeled contraindications to the requested agent, including but not limited to IgA deficiency with antibodies against IgA and a history of hypersensitivity



Approval Duration: Gamunex-C: 12 months
Flebogamma, Gammagard, Gammaplex: 2 months

References:

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3. Hizentra prescribing information. CSL Behring LLC. April 2023.
4. Perez EE, Orange JS, Bonilla F, et al. Update on the use of immunoglobulin in human disease: A review of evidence. J Allergy Clin Immunol. 2017;139(3S):S1-S46.
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8. Cassard L, Seraly N, Riegert M, Patel A, Fernandez AP. Dermatomyositis: practical guidance and unmet needs. Immunotargets Ther. 2024;13:151-172. doi:10.2147/ITT.S381472

Policy Owned by: Curative PBM team