gamunex®-C
immune globulin injection (human), 10%
caprylate/chromatography purified

Copay Assistance Program now available for your patients

START their IG therapy here

- PATIENTS WITH CIDP OR PIDD are eligible with no financial qualification or paperwork necessary
- SAVINGS UP TO $2500 over 12 months to cover deductibles, copayment, and coinsurance medication costs*
- ELIGIBILITY—patients must be enrolled in Gamunex Connexions and have commercial insurance that:
  - Covers medication costs for GAMUNEX-C
  - Allows for copay or coupon assistance

STAY and SUCCEED with additional support

- PATIENT ASSISTANCE PROGRAM (PAP) is available for patients with CIDP or PIDD who are underinsured or without insurance
- SUPPORT offered through Gamunex Connexions provides various resources throughout the patient journey, including a primary point of contact

*For terms and conditions, please see inside.
IG=immune globulin, CIDP=chronic inflammatory demyelinating polyneuropathy, PIDD=primary immunodeficiency disease, ITP=idiopathic thrombocytopenic purpura.

NOW COVERS SC FOR PIDD TOO!

gamunex® connexions

Helping patients **start**, **stay**, and **succeed** on IG therapy

Please see inside for Important Safety Information and refer to accompanying full Prescribing Information for GAMUNEX-C.

GRIFOLS

Terms and conditions for Copay Assistance Program

This manufacturer coupon program is not valid for prescriptions reimbursed, in whole or in part, by Medicaid, Medicare, Medigap, VA, DoD, TRICARE, or any other federal or state healthcare programs, including state pharmaceutical assistance programs, and where prohibited by the health insurance provider or by law. This coupon program provides a maximum benefit of $2500 for eligible out-of-pocket costs and expires 12 months from the date of first assisted fill. Eligible costs include deductible, copayment, and coinsurance costs for GAMUNEX®-C (immune globulin injection [human], 10% caprylate/chromatography purified) medication. Nonmedication expenses, such as ancillary supplies or administration-related costs, are not eligible. To be eligible, patients must 1) be starting or receiving treatment with (and have a current prescription for) GAMUNEX-C with an ICD9 or ICD10, as applicable of a chronic inflammatory demyelinating polyneuropathy (CIDP) or primary immunodeficiency disease (PIDD); and 2) have commercial insurance that covers medication costs for GAMUNEX-C treatment and allows for copay/coupon assistance. Acceptance of this offer must be consistent with the terms of benefits provided by patient's health insurance provider. Offer limited to outpatient IG treatment and may not be combined with any other coupon, discount, prescription savings card, rebate, free trial, or other offer. This program is only valid for residents of the United States, excluding Puerto Rico and other US territories. Grifols reserves the right to change or discontinue this program at any time without notice. This is not health insurance.

Important Safety Information

GAMUNEX®-C (immune globulin injection [human], 10% caprylate/chromatography purified) is indicated for the treatment of primary humoral immunodeficiency disease (PIDD) in patients 2 years of age and older, idiopathic thrombocytopenic purpura (ITP) in adults and children, and chronic inflammatory demyelinating polyneuropathy (CIDP) in adults.

Thrombosis may occur with immune globulin products, including GAMUNEX-C. Risk factors may include: advanced age, prolonged immobilization, hypercoagulable conditions, history of venous or arterial thrombosis, use of estrogens, indwelling central vascular catheters, hyperviscosity, and cardiovascular risk factors. Thrombosis may occur in the absence of known risk factors. For patients at risk of thrombosis, administer GAMUNEX-C at the minimum dose and infusion rate practicable. Ensure adequate hydration in patients before administration. Monitor for signs and symptoms of thrombosis and assess blood viscosity in patients at risk for hyperviscosity.

Renal dysfunction, acute renal failure, osmotic nephrosis, and death may occur with immune globulin intravenous (IVIG) products in predisposed patients. Patients predisposed to renal dysfunction include those with any degree of preexisting renal insufficiency, diabetes mellitus, age greater than 65, volume depletion, sepsis, paraproteinemia, or patients receiving known nephrotoxic drugs. Renal dysfunction and acute renal failure occur more commonly in patients receiving IVIG products containing sucrose. GAMUNEX-C does not contain sucrose. For patients at risk of renal dysfunction or failure, administer GAMUNEX-C at the minimum concentration available and the minimum infusion rate practicable.

GAMUNEX-C is contraindicated in patients who have had an anaphylactic or severe systemic reaction to the administration of human immune globulin. It is contraindicated in IgA-deficient patients with antibodies against IgA and history of hypersensitivity.

Severe hypersensitivity reactions may occur with IVIG products, including GAMUNEX-C. In case of hypersensitivity, discontinue GAMUNEX-C infusion immediately and institute appropriate treatment.

Important Safety Information (cont.)

Monitor renal function, including blood urea nitrogen (BUN), serum creatinine, and urine output in patients at risk of developing acute renal failure.

Hyperproteinemia, increased serum viscosity, and hyponatremia may occur in patients receiving IVIG treatment, including GAMUNEX-C.

There have been reports of aseptic meningitis, hemolytic anemia, and noncardiogenic pulmonary edema (transfusion-related acute lung injury [TRALI]) in patients administered with IVIG, including GAMUNEX-C.

The high-dose regimen (1g/kg x 1-2 days) is not recommended for individuals with expanded fluid volumes or where fluid volume may be a concern.

Because GAMUNEX-C is made from human blood, it may carry a risk of transmitting infectious agents, eg, viruses, the variant Creutzfeldt-Jakob disease (vCJD) agent, and, theoretically, the Creutzfeldt-Jakob disease (CJD) agent.

Do not administer GAMUNEX-C subcutaneously in patients with ITP because of the risk of hematoma formation.

Periodic monitoring of renal function and urine output is particularly important in patients judged to be at increased risk of developing acute renal failure. Assess renal function, including measurement of BUN and serum creatinine, before the initial infusion of GAMUNEX-C and at appropriate intervals thereafter.

Consider baseline assessment of blood viscosity in patients at risk for hyperviscosity, including those with cryoglobulins, fasting chylomicronemia/markedly high triacylglycerols (triglycerides), or monoclonal gammopathies, because of the potentially increased risk of thrombosis.

If signs and/or symptoms of hemolysis are present after an infusion of GAMUNEX-C, perform appropriate laboratory testing for confirmation.

If TRALI is suspected, perform appropriate tests for the presence of antineutrophil antibodies and anti-HLA antibodies in both the product and patient's serum.

After infusion of IgG, the transitory rise of the various passively transferred antibodies in the patient's blood may yield positive serological testing results, with the potential for misleading interpretation.

In clinical studies, the most common adverse reactions with GAMUNEX-C were headache, pyrexia, hypertension, chills, rash, nausea, arthralgia, and asthenia (in CIDP); cough, rhinitis, pharyngitis, headache, asthma, nausea, fever, diarrhea, and sinusitis with intravenous use (in PIDD) and local infusion-site reactions, fatigue, headache, upper respiratory tract infection, arthralgia, diarrhea, nausea, sinusitis, bronchitis, depression, allergic dermatitis, migraine, myalgia, viral infection, and pyrexia with subcutaneous use (in PIDD); and headache, ecchymosis, vomiting, fever, nausea, rash, abdominal pain, back pain, and dyspepsia (in ITP).

The most serious adverse reactions in clinical studies were pulmonary embolism (PE) in 1 subject with a history of PE (in CIDP), an exacerbation of autoimmune pure red cell aplasia in 1 subject (in PIDD), and myocarditis in 1 subject that occurred 50 days post-study drug infusion and was not considered drug related (in ITP).

Please see accompanying full Prescribing Information for GAMUNEX-C.

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Billing and reimbursement at a glance

Assistance is available for patients receiving GAMUNEX-C

FOR MEDICAL BILLING:

1. REGISTER your patients at the medical billing portal: gamunex-c.medmonk.com

2. OPT IN your patients to qualify for financial assistance

3. PROCESS a claim with instant adjudication through the medical billing portal

FOR PHARMACY BILLING:

1. SUBMIT a claim to the primary payer first. If patient requires copay assistance, submit the balance to Medmonk as a secondary payer coordination of benefits (COB) using the following:

BIN: 016664 | PCN: MEDMONK | CARDHOLDER: MEDMONK

gamunex® connexions

For technical support regarding the Copay Assistance Program, call
1-866-ADHERE2 • (234-3732)
(available 9 AM to 5 PM ET)
or email support@medmonk.com

For all other questions regarding Gamunex Connexions, call
1-888-MYGAMUNEX • (694-2686)
(available 8 AM to 8 PM ET)
or visit GAMUNEX-C.com

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