

Covered ICD codes for GAMUNEX®-C (immune globulin injection [human], 10% caprylate/chromatography purified) Copay Assistance

ICD-9

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| <p>279.00 Hypogammaglobulinemia, unspecified
Agammaglobulinemia NOS</p> <p>279.01 Selective IgA immunodeficiency</p> <p>279.02 Selective IgM immunodeficiency</p> <p>279.03 Other selective immunoglobulin deficiencies
Selective deficiency of IgG</p> <p>279.04 Congenital hypogammaglobulinemia
Agammaglobulinemia:
 <ul style="list-style-type: none"> • Bruton's type • X-linked </p> <p>279.05 Immunodeficiency with increased IgM
Immunodeficiency with hyper-IgM:
 <ul style="list-style-type: none"> • Autosomal recessive • X-linked </p> <p>279.06 Common variable immunodeficiency
Dysgammaglobulinemia (acquired)
(congenital) (primary)
 <ul style="list-style-type: none"> • Hypogammaglobulinemia: <ul style="list-style-type: none"> • Acquired primary • Congenital non–sex-linked • Sporadic </p> <p>279.09 Other deficiency of humoral immunity</p> <p>279.12 Wiskott-Aldrich syndrome</p> <p>279.2 Combined immunity deficiency</p> <p>357.81 Chronic inflammatory demyelinating polyneuritis (CIDP)</p> | <p>D80.0 Hereditary hypogammaglobulinemia</p> <p>D80.5 Immunodeficiency with increased immunoglobulin M (IgM)</p> <p>D80.6 Antibody deficiency with near-normal immunoglobulins or with hyperimmunoglobulinemia</p> <p>D80.8 Other immunodeficiencies with predominantly antibody defects</p> <p>D80.9 Immunodeficiency with predominantly antibody defects, unspecified</p> <p>D83.0 Common variable immunodeficiency with predominant abnormalities of B-cell numbers and functions</p> <p>D83.2 Common variable immunodeficiency with autoantibodies to B or T cells</p> <p>D83.8 Other common variable immunodeficiencies</p> <p>D83.9 Common variable immunodeficiency, unspecified</p> <p>D80.7 Transient hypogammaglobulinemia of infancy</p> <p>D82.0 Wiskott-Aldrich syndrome</p> <p>D81.0 Severe combined immunodeficiency (SCID) with reticular dysgenesis</p> <p>D81.1 Severe combined immunodeficiency (SCID) with low T-and B-cell numbers</p> <p>D81.2 Severe combined immunodeficiency (SCID) with low or normal B-cell numbers</p> <p>D81.6 Major histocompatibility complex class I deficiency</p> <p>D81.7 Major histocompatibility complex class II deficiency</p> <p>D81.89 Other combined immunodeficiency</p> <p>D81.9 Combined immunodeficiency, unspecified</p> <p>G61.81 Chronic Inflammatory demyelinating polyneuritis</p> |
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ICD-10

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| <p>D80.1 Nonfamilial hypogammaglobulinemia</p> <p>D80.2 Selective deficiency of immunoglobulin A (IgA)</p> <p>D80.4 Selective deficiency of immunoglobulin M (IgM)</p> <p>D80.3 Selective deficiency of immunoglobulin G (IgG) subclasses</p> | <p>D81.7 Major histocompatibility complex class II deficiency</p> <p>D81.89 Other combined immunodeficiency</p> <p>D81.9 Combined immunodeficiency, unspecified</p> <p>G61.81 Chronic Inflammatory demyelinating polyneuritis</p> |
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