CODING REFERENCE CARD- IMMUNODEFICIENCY STATUS

Immunodeficiency with predominantly antibody defects	
D80.0	Hereditary hypogammaglobulinemia
D80.1	Nonfamilial hypogammaglobulinemia
D80.2	Selective deficiency of immunoglobulin A (IgA)
D80.3	Selective deficiency of immunoglobulin G (IgG) subclasses
D80.4	Selective deficiency of immunoglobulin M (IgM)
D80.5	Immunodeficiency with increased immunoglobulin M (IgM)
D80.6	Antibody deficiency with near-normal immunoglobulins or with hyperimmunoglobulinemia
D80.7	Transient hypogammaglobulinemia of infancy
D80.8	Other immunodeficiencies with predominantly antibody defects (Kappa light chain deficiency)
D80.9	Immunodeficiency with predominantly antibody defects, unspecified

Combined immunodeficiencies	
D81.0	Severe combined immunodeficiency [SCID] with reticular dysgenesis
D81.1	Severe combined immunodeficiency [SCID] with low T- and B-cell numbers
D81.2	Severe combined immunodeficiency [SCID] with low or normal B-cell numbers
D81.30	Adenosine deaminase deficiency, unspecified (ADA deficiency NOS)
D81.31	Severe combined immunodeficiency due to adenosine deaminase deficiency
D81.32	Adenosine deaminase 2 deficiency **Code also any associated manifestations such as polyarteritis nodosa or stroke**
D81.39	Other adenosine deaminase deficiency
D81.4	Nezelof's syndrome
D81.5	Purine nucleoside phosphorylase [PNP] deficiency
D81.6	Major histocompatibility complex class I deficiency
D81.7	Major histocompatibility complex class II deficiency

Other combined immunodeficiencies	
D81.810	Biotinidase deficiency
D81.818	Other biotin-dependent carboxylase deficiency
D81.819	Biotin-dependent carboxylase deficiency, unspecified
D81.89	Other combined immunodeficiency
D81.9	Combined immunodeficiency, unspecified (Severe combined immunodeficiency disorder [SCID] NOS)

Immunodeficiency Associated with other major defects	
D82.0	Wiskott-Aldrich syndrome
D82.1	Di George's syndrome
D82.2	Immunodeficiency with short-limbed stature
D82.3	Immunodeficiency following hereditary defective response to Epstein-Barr virus
D82.4	Hyperimmunoglobulin E (IgE) Syndrome
D82.8	Immunodeficiency associated with other specified major defects
D82.9	Immunodeficiency associated with major defect, unspecified

NOTE: This tool is intended to assist with documentation only and not intended to take the place of clinical analysis. Information regarding any law or regulation does not constitute legal or tax advice and is subject to change based upon the issuance of new guidance and/or change in laws or regulations. Reference Official ICD-10-CM coding guidelines and manuals or electronic medical coding software for accurate ICD-10-CM codes and specificity.

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