CODING REFERENCE CARD
HEMOLYTIC & HEMORRHAGIC DISORDERS

Hemolytic Anemias	
D55.0	Anemia due to glucose-6-phosphate dehydrogenase [G6PD] deficiency
D55.1	Anemia due to other disorders of glutathione metabolism
D55.2	Anemia due to disorders of glycolytic enzymes
D55.3	Anemia due to disorders of nucleotide metabolism
D55.8	Other anemias due to enzyme disorders
D55.9	Anemia due to enzyme disorder, unspecified
D58.8	Other specified hereditary hemolytic anemias
D58.9	Hereditary hemolytic anemia, unspecified
D59.0	Drug-induced autoimmune hemolytic anemia
D59.10	Autoimmune hemolytic anemia, unspecified
D59.11	Warm autoimmune hemolytic anemia
D59.12	Cold autoimmune hemolytic anemia (Rare)
D59.13	Mixed type autoimmune hemolytic anemia
D59.19	Other autoimmune hemolytic anemia
D59.2	Drug-induced nonautoimmune hemolytic anemia
D59.30	Hemolytic-uremic syndrome, unspecified
D59.31	Infection-associated hemolytic-uremic syndrome
D59.32	Hereditary hemolytic-uremic syndrome
D59.39	Other hemolytic-uremic syndrome *Secondary hemolytic-uremic syndrome*
D59.4	Other nonautoimmune hemolytic anemias
D59.6	Hemoglobinuria due to hemolysis from other external causes
D59.8	Other acquired hemolytic anemias
D59.9	Acquired hemolytic anemia, unspecified

Thalassemia	
D56.0	Alpha thalassemia
D56.1	Beta thalassemia (Thalassemia Major)
D56.2	Delta-beta thalassemia
D56.3	Thalassemia minor
D56.5	Hemoglobin E-beta thalassemia
D56.8	Other thalassemias
D56.9	Thalassemia, unspecified
D57.40	Sickle-cell thalassemia without crisis
D57.411	Sickle-cell thalassemia, unspecified, with acute chest syndrome
D57.412	Sickle-cell thalassemia, unspecified, with splenic sequestration
D57.413	Sickle-cell thalassemia, unspecified, with cerebral vascular involvement
D57.418	Sickle-cell thalassemia, unspecified, with crisis with other specified complication
D57.419	Sickle-cell thalassemia, unspecified, with crisis
Other Hemolytic Disorders	
D57.00	Hb-SS disease with crisis, unspecified
D57.01	HB-SS disease with acute chest syndrome
D57.02	Hb-SS disease with splenic sequestration
D57.03	Hb-SS disease with cerebral vascular involvement
D57.09	Hb-SS disease with crisis with other specified complication
D57.1	Sickle-cell disease without crisis
D57.3	Sickle-cell trait
D75.821	Non-immune heparin-induced thrombocytopenia
D75.822	Immune-mediated heparin-induced thrombocytopenia
D75.828	Other heparin-induced thrombocytopenia syndrome *Autoimmune HIT syndrome*
D75.829	Heparin-induced thrombocytopenia, unspecified
D75.84	Other platelet-activating anti-PF4 disorders

Thrombocytopenia	
D69.0	Allergic purpura
D69.1	Qualitative platelet defects
D69.2	Other nonthrombocytopenic purpura
D69.3	Immune thrombocytopenic purpura *Hemorrhagic purpura* *Idiopathic thrombocytopenic purpura*
D69.41	Evans syndrome
D69.42	Congenital and hereditary thrombocytopenia purpura
D69.49	Other primary thrombocytopenia
D69.51	Posttransfusion purpura
D69.59	Other secondary thrombocytopenia
D69.6	Thrombocytopenia, unspecified
D69.8	Other specified hemorrhagic conditions *Vascular pseudohemophilia*
D69.9	Hemorrhagic condition, unspecified

Note: The listed codes are for the most common forms of sickle-cell disease. Additional codes exist. Please consult official coding sources to identify a more specific code if required.

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