Sept. 17, 2009 ICD-9-CM Coordination and Maintenance Committee Meeting

- Primary Hereditary Hemochromatosis
- Acquired (Secondary) Iron Overload
- Other Iron Overload (rarely as severe)

- Primary Hereditary Hemochromatosis
 - HFE-associated (type 1)
 - C282Y homozygosity
 - C282Y/H63D compound heterozygosity
 - Juvenile; hepcidin and hemojuvelin mutations (type 2)
 - transferrin receptor 2 mutations (type 3)
 - ferroportin mutations (type 4)
- Acquired (Secondary) Iron Overload
- Other Iron Overload (rarely as severe)

References:

OMIM articles 235200, 602390, 604250, and 606069. Conn's Current Therapy 2008, Ch. 14, Hemochromatosis.

- Primary Hereditary Hemochromatosis
- Acquired (Secondary) Iron Overload
 - Iron loading anemias
 - Thalassemia major
 - Sideroblastic anemia
 - Chronic hemolytic anemia
 - Dietary iron overload
 - Parenteral iron overload (including multiple blood transfusions)
- Other Iron Overload (rarely as severe)

References:

Conn's Current Therapy 2008, Ch. 14, Hemochromatosis.

- Primary Hereditary Hemochromatosis
- Acquired (Secondary) Iron Overload
- Other Iron Overload (rarely as severe)
 - Long term hemodialysis
 - Chronic liver disease (alcoholic; hepatitis C)
 - Porphyria cutanea tarda
 - Dysmetabolic iron overload syndrome
 - Post portacaval shunting
 - Iron overload in Africa
 - Neonatal iron overload
 - Aceruloplasminemia
 - Congenital atransferrinemia

References:

Conn's Current Therapy 2008, Ch. 14, Hemochromatosis.

Concepts for Coding in Iron Overload

- Hemochromatosis, unspecified
- Primary hereditary hemochromatosis
- Hemochromatosis due to multiple blood transfusions
- Iron overload due to conditions classified elsewhere
 - Code first underlying conditions
- Other iron overload
 - Other hemochromatosis