Genetically Handicapped Persons Program Eligible Medical Conditions Include:

1) Diseases of the Blood, 2) Cystic Fibrosis, 3) Diseases of the Brain and Nerves, 4) Diseases of the Protein Metabolism, 5) Diseases of Carbohydrates Metabolism, 6) Disease of Copper Metabolism, and 7) Von Hippel-Lindau Disease (VHL)

I. Diseases of the Blood

- Hemophilia or Factor Deficiency (FACTORS I, II, V, VII, VIII,IX,X, XI, XIII)
- Von Willebrand's Disease
- Congenital Hereditary Platelet Deficiency Diseases or dysfunction such as Congenital Thrombasthenia (Glanzmann's Thrombasthenia) and Thrombocytopathia
- Hemoglobinopathies with Anemia:
- Sickle Cell Disease (Not sickle cell trait)
- Thalassemia

II. Cystic Fibrosis (CF)

- Cystic Fibrosis (CF) (Non DHCS)

III. Diseases of the Brain and Nerves

- Huntington's Disease (HD)
- Joseph's Disease (Spinocerebellar Ataxia III or SCA III)
- Friedrich's Ataxia, ataxias due to spinocerebellar degeneration
- Hereditary Spastic Paraplegia
- Roussy-Levy Syndrome
- Olivopontocerebellar Degeneration/Atrophy (OPCA)
- Refsum's Disease
- Charcot-Marie Tooth Syndrome (CMT Syndrome)

IV. Diseases of Protein Metabolism

- Phenylketonuria (PKU)
- Tyrosinemia
- Disturbances of Metabolism of Leucine, Isoleucine, Valine
- Hypervalinemia
- Intermittent Branched-Chain Ketonuria
- Leucine Induced/Leucinosis
- Maple Syrup Urine Disease (MSUD)
- Propionic Acidemia and Methylmalonic Acidemia
- Lactic and Pyruvate Metabolism Disorders
- Hereditary Orotic (Pyrimidine Acidemia)
- Homocystinuria
- Hypermethioninemia
- Argininosuccinic Aciduria
- Citrullinemia
- Disorders of Metabolism of Ornithine, Citrulline, Argininosuccinic Acid, Arginine, and Ammonia
- Hyperammonemia
- Hyperornithinemia

V. Diseases of Carbohydrates Metabolism

- Galactosemia
- Galactose-1-Phosphate Uridyltransferase Deficiency
- Galactosuria

VI. Disease of Copper Metabolism

- Wilson's Disease

VII. Von Hippel-Lindau Disease (VHL)