
Absence of sensation
Jarcho−Levin syndrome
Tuberculosis, Multidrug−Resistant
Chondrodysplasia
Attention Deficit and Disruptive Behavior Disorders
Urea Cycle Disorders, Inborn
Leber Congenital Amaurosis
Nonsyndromic Deafness
Photopsia
Epilepsy, Rolandic
Alkaline Phosphatase Adverse Event
Drug withdrawal syndrome
Autoimmune thyroid disease
Fanconi Syndrome
Common Migraine
Active tuberculosis
Infectious Otitis Media
Constitutional delay of growth and puberty
Manic
Psychosexual Disorders
Brugada Syndrome (disorder)
Hypothyroidism, Autoimmune
Acidosis
Chronic pain
Corneal dystrophy
Acth−Independent Macronodular Adrenal Hyperplasia
Aganglionosis, Colonic
Abnormal involuntary movement
Erythroblastosis, Fetal
Laryngitis
Albinism
Hirschsprung disease 1
Hematocrit level
Angelman Syndrome
Double Outlet Right Ventricle
Hyperbilirubinemia
Congenital adrenal hyperplasia due to 21 hydroxylase deficiency
Acute type B viral hepatitis
CONOTRUNCAL HEART MALFORMATIONS (disorder)
Ambiguous Genitalia
Congenital adrenal hyperplasia
Hyperbilirubinemia, Neonatal
Placental Steroid Sulfatase Deficiency
21−hydroxylase deficiency
Blood pressure finding
Absence Epilepsy
Cutaneous hypersensitivity
Rheumatoid Vasculitis
Idiopathic generalized epilepsy