Mitochondrial Disorders, Diseases and Syndromes

Represented by CMDN / EMDN

Alpers Disease (Progressive Infantile Poliodystrophy)
Barth Syndrome / LIC (Lethal Infantile Cardiomyopathy)
Caritine-Acyl-Carnitine Deficiency
Carnitine Deficiency
Co-Enzyme Q10 Deficiency
Mitochondrial Respiratory Chain Disorders
Complex I Deficiency
Complex II Deficiency
Complex III Deficiency
Complex IV / COX Deficiency
Complex V Deficiency
CPEO (Chronic Progressive External Ophthalmoplegia Syndrome)
CPT I Deficiency
CPT II Deficiency
KSS (Kearns-Sayre Syndrome)
Lactic Acidosis
LCAD (Long-Chain AycI-CoA Dehydrogenase Deficiency)
LCHAD
Leigh Disease or Syndrome (Subacute Necrotizing Encephalmyelopathy)
LHON (Leber Hereditary Optic Neuropathy)
Luft Disease
MAD / Glutaric Aciduria Type II (Multiple Acyl-CoA Dehydrogenase Deficiency)
MACD (Medium Chain Acyl-CoA Dehydrogenase Deficiency)
MELAS (Mitochondrial Encephalomyopathy, Lactic Acidosis and Stroke Like Episodes)
MERRF (Myoclonic Epilepsy and Ragged Red Fibre Disease)
Mitochondrial Cytopathy
Mitochondrial DNA Depletion
Mitochondrial Encephalopathy
Mitochondrial Myopathy
MINGIE (Myoneurogastointestinal Disorder and Encephalopathy)
NARP (Neuropathy, Ataxia and Retinitis Pigmentosa)
Pearson Syndrome
Pyruvate Carboxylase Deficiency
Pyruvate Dehydrogenase Deficiency
SCAD (Short-Chain Acyl-CoA Dehydrogenase Deficiency)
SCHAD
VLCAD (Very Long-Chain Aycl-CoA Dehydrogenase Deficiency

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