DYSTONIA AND PERIPHERAL NEUROPATHY

INHERITED

Autosomal dominant

- SCA3
- SCA1
- SCA11

Autosomal recessive

- MPAN
- Arginase deficiency
- POLG mutations
- AFG3L2 homozygous mutations
- Chorea-acanthocytosis
- Friedreich's ataxia
- Ataxia-telangiectasia
- Ataxia-oculomotor apraxia type 1
- Ataxia-oculomotor apraxia type 2
- Ataxia-telangiectasia like syndrome
- Cerebrotendinous xanthomatosis
- Tay-Sach's disease (late onset form)
- Cockayne syndrome
- Xeroderma pigmentosum
- Sterol carrier protein X (SCPx) mutations
- Niemann-Pick type C
- Metachromatic leukodystrophy
- Globoid cell leukodystrophy (Krabbe disease)
- Chediak-Higashi disease

Mitochondrial

- Mitochondrial disease

Chromosomal

- Chr 18q deletion

ACQUIRED

Infection

- HIV