DYSTONIA WITH CEREBELLAR ATAXIA

INHERITED

Autosomal dominant

SCA1 (ATXN1 mutations)
SCA2 (ATXN3 mutations)
SCA3 (ATXN3 mutations)
SCA6 (CACNA1A mutations)
SCA7 (ATX7 mutations)
SCA11 (TTBK2 mutations)
SCA12 (PPP2R2B mutations)
SCA 14 (PRKCG mutations)
SCA17 (TATA-box binding protein mutations)
Dentatorubropallidoluysian atrophy (DRPLA)
Glucose transporter 1 deficiency (SLC2A1 mutations)

Autosomal recessive

Friedreich's ataxia
Ataxia-telangiectasia
Ataxia-oculomotor apraxia type 1
Ataxia-oculomotor apraxia type 2
Ataxia-telangiectasia like syndrome
Cerebrotendinous xanthomatosis
CoQ10 deficiency (e.g. CABC1/ADCK3 mutations)
SPG7 (paraplegin)
AFG3L2 mutations (SCA28)
Cockayne syndrome
Xeroderma pigmentosum
Polymerase gamma (POLG) mutations
Fatty acid hydroxylase-associated neurological disease (FAHN)
Niemann-Pick Type C
Aceruloplasminemia
Neuronal ceroid lipofuscinosis

X-linked

Fragile-X tremor ataxia syndromes (FXTAS)

Mitochondrial

Mitochondrial disease

IDIOPATHIC

Sporadic

Multiple system atrophy (MSA)
Dystonia with cerebellar atrophy (DYTCA)
Cerebral folate deficiency