

DYSTONIA AND PARKINSONISM IN OLDER ADULTS

Presenting as Parkinson's plus syndrome

With focal dystonia

Sporadic

- Early onset Parkinson's disease
- Multiple system atrophy
- Progressive supranuclear palsy

Presenting as corticobasal syndrome

Sporadic

- Corticobasal degeneration
- Progressive supranuclear palsy
- Alzheimer's disease

Autosomal dominant

- Tau gene mutations
- Progranulin gene mutations
- c9orf72 gene mutations

Infection

- Prion disease

With ataxia

Sporadic

- Multiple system atrophy

Autosomal dominant

- SCA2
- SCA3
- SCA17
- SCA6

Presenting with variable mix of dystonia and parkinsonism

INHERITED

Autosomal dominant

- Fahr's syndrome
- Neuroferritinopathy

Autosomal recessive

- Wilson's disease
- Neuroacanthocytosis
- Niemann-Pick type C
- Manganese transporter deficiency

X-linked

- Lubag (DYT3)

ACQUIRED

Infection

- HIV
- Prion disease

Drug-induced

- Typical and atypical neuroleptics
- Dopamine blocking anti-emetics

Toxic

Wasp sting

Carbon monoxide (delayed onset)

Methanol (delayed onset)

Disulfiram (delayed onset)

Cyanide (delayed onset)

Manganese

- Manganese miners

- Welders

- Chronic liver disease (hepatolenticular degeneration)

- TPN

- Ephedrone recreational use

Metabolic

Hypoxia (often delayed onset)

- Asphyxia

- Perinatal hypoxia-ischemia

Extrapontine myelinosis

Hepatocerebral syndrome

Neoplastic

Glioma

Lymphoma