DISEASE:
Parkinsonian-pyramidal syndrome

NAME:	Parkinsonian-pyramidal syndrome
DESCRIPTION:	Parkinsonian-pyramidal syndrome is a rare, genetic, neurological disorder characterized by the association of both parkinsonian (i.e. bradykinesia, rigidity and/or rest tremor) and pyramidal (i.e. increased reflexes, extensor plantar reflexes, pyramidal weakness or spasticity) manifestations, which vary according to the underlying associated disease (e.g. neurodegenerative disease, inborn errors of metabolism).
ORPHACODE:	171695
SYNONYMS:	Pallidopyramidal syndrome
XREF(S):	Orphanet ICD-10 OMIM OMIM OMIM
ANALYTE(S):	SNCA FBXO7
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RELATED CONTENT

Related Genetic Tests

- Neurodegeneration (gene panel)
- Parkinson (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- F-box protein 7
- synuclein alpha

Related Gene Panels

- Neurodegeneration (99 genes) - IPG
- Parkinson (18 genes) - KUL