# DISEASE:
# Autosomal spastic paraplegia type 72

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| **NAME:** | Autosomal spastic paraplegia type 72 |
| **DESCRIPTION:** | Autosomal spastic paraplegia type 72 is a rare, genetic, pure hereditary spastic paraplegia disorder characterized by early childhood onset of slowly progressive crural spastic paraparesis presenting with spastic gait, mild stiffness at rest, hyperreflexia (in lower limbs), extensor plantar responses and, in some, mild postural tremor, pes cavus, sphincter disturbances and sensory loss at ankles. |
| **ORPHACODE:** | 401849 |
| **SYNONYMS:** | SPG72 |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | REEP2 |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Hereditary Spastic Paraplegia (gene panel)
- Spastic Paraplegia (gene panel)

## Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen

## Related Analytes

- receptor accessory protein 2

## Related Gene Panels

- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Spastic Paraplegia (89 genes) - IPG