Autosomal dominant optic atrophy plus syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant optic atrophy plus syndrome. ORPHA:1215

Autosomal dominant optic atrophy plus syndrome (ADOA plus) is a variant of autosomal dominant optic atrophy (ADOA; see this term) associating the typical optic atrophy with other extra-ocular manifestations such as sensorineural deafness, myopathy, chronic progressive external ophthalmoplegia, ataxia and peripheral neuropathy. More rarely, other manifestations have been associated with this condition, such as spastic paraplegia, multiple-sclerosis like illness.