Opis choroby *

Definicja

Frontotemporal dementia with motor neuron disease (FTD-MND) is a type of frontotemporal lobar degeneration characterized by the insidious onset (between the ages of 38-78 years) of dementia-associated psychiatric symptoms (e.g. personality changes, uninhibited behavior, irritability, aggressiveness), memory difficulties, global intellectual impairment, emotional disorders and transcortical motor aphasia that eventually leads to mutism, in addition to the manifestations of motor neuron disease such as neurogenic muscular wasting (similar to what is seen in amyotrophic lateral sclerosis; see this term). The disease is progressive, with death occurring 2-5 years after onset.

Dane

Klasyfikacja
Choroba

Synonimy
FTD-ALS
FTD-ALS
FTD-MND
Otępienie czołowo-skroniowe ze stwardnieniem zanikowym bocznym
FTD-MND
Frontotemporal dementia with amyotrophic lateral sclerosis

Kod ORPHA
275872

Kod OMIM
616439

Kod ICD10
G31.0

Kod ICD11
6D83

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*Źródło

orphanet