

## Davide Pareyson, MD

Davide Pareyson is a Clinical Neurologist working at the Fondazione IRCCS Istituto Neurologico C.Besta (FINCB) of Milan, Italy, where he is currently Head of the Rare Neurological Diseases Unit and of the Functional Department of Neurodegenerative and Rare Neurological Diseases. He obtained his MD degree in Medicine in 1984, completed his Specialisation in Neurology in 1988 and in Clinical Neurophysiology in 1991, and achieved the Italian national certification of eligibility as Full Professor.



His main interest is clinical research on inherited neurological disorders, with special reference to hereditary peripheral neuropathies and motor neuronopathies. He performed studies on phenotype-genotype correlation, clinical findings, electrophysiology, neuropathology of hereditary neuropathies (particularly Charcot-Marie-Tooth disease – CMT - and related neuropathies, but also amyloid neuropathy) and other neurogenetic disorders including spinal and bulbar muscle atrophy (SBMA). During the last years, he has been working on the development of outcome measures for hereditary neuropathies and other rare diseases and has coordinated and participated in clinical trials and natural history studies in inherited and acquired neuropathies.

He was member of board of PNS, ENS, Chair of the CMTR, coordinator of the neurogenetic scientific panel and member of committees of the EAN. He is Co-Chair of the Neuropathy Working Group of the EURO-NMD ERN and Co-Chair of the EAN/PNS Guidelines on Diagnosis and Treatment of ATTRv-related neurological manifestations. He has co-authored more than 340 papers on peer-reviewed Journals mainly on hereditary disorders and neuromuscular diseases.