

Supplementary Table S1. Demographic and clinical characteristics of the whole cohort of study.

	Count (%)	Mean (SD)	n
Males/Females	84 (64.6%)/46 (35.4%)		130
Age at disease onset (years)		48.69 (19.25)	130
Disease duration (months)		130.96 (150.65)	130
Follow-up (months)		96.98 (84.93)	130
Disease onset			130
Chronic	89 (68.5%)		
Acute	24 (18.5%)		
Subacute	17 (13.1%)		
Phenotype			130
Typical CIDP	84 (64.6%)		
DADS	16 (12.3%)		
MADSAM	15 (11.5%)		
Pure motor CIDP	11 (8.5%)		
Sensory CIDP	2 (1.5%)		
Focal CIDP	2 (1.5%)		
Disease course			130
Monophasic	26 (20.0%)		
Relapsing	30 (23.1%)		
Chronic progressive (or chronic active)	74 (56.9%)		
EFNS/PNS electrodiagnostic criteria			130
Definite	78 (60.0%)		
Probable	18 (13.8%)		
Possible	19 (14.6%)		
Not fulfilled	15 (11.5%)		
EFNS/PNS diagnostic category			130
Definite CIDP	103 (79.2%)		
Probable CIDP	11 (8.5%)		
Possible CIDP	0 (0.0%)		
Not fulfilled	16 (12.3%)		

Legend to the table: SD, standard deviation; CIDP, Chronic Inflammatory Demyelinating Polyneuropathy; DADS, Distal Acquired Demyelinating Symmetric Neuropathy; MADSAM, Multifocal Acquired Demyelinating Sensory And Motor Neuropathy; EFNS/PNS, European Federation of Neurological Societies/Peripheral Nerve Society.