

Supplementary Table 5. CNS features seen in other Charcot-Marie-Tooth disease subtypes.	
CNS feature	Charcot-Marie-Tooth disease-related gene
Hearing loss	PMP22, MPZ, SH3TC2, NDRG1, INF2
Pyramidal features	MFN2, KIF5A, GDAP1, SPG11, GJB1
Ataxia (sensory or cerebellar)	PMP22, MPZ, PRX, GJB1, MME