

Supplementary Table 1. Number of mutation carriers (1A) and pathological diagnosis (1B) for each clinical diagnosis group.

1A

		Genetic Diagnosis					
		<i>C9orf72</i>	GRN	MAPT	GRN+ <i>C9orf72</i>	<i>C9orf72+SQSTM1</i>	Total
Clinical Diagnosis	FTD-MND	2	0	0	1	1	4
	bvFTD	20	9	23	0	0	52
	PNFA	2	4	1	0	0	7
	PPA-NOS	0	2	0	0	0	2
	SD	0	0	0	0	0	0
	Total	24	15	24	1	1	65

1B

		Pathological Diagnosis			
		TDP-43	tau	FUS	Total
Clinical Diagnosis	FTD-MND	4	0	0	4
	bvFTD	31	31	3	65
	PNFA	8	7	0	15
	PPA-NOS	2	0	0	2
	SD	16	2	0	18
	Total	61	40	3	104

Supplementary Table 2. Volumetric comparisons between the different clinical subgroups in the sporadic cohort for the right, left and total thalamic volume. Volumetric comparisons are adjusted for age, gender and scanner type, and also for disease duration when comparing between FTD groups. Bold represents a significant difference between groups after correcting for multiple comparisons.