Supplementary Table S2. Family history (FHx) characteristics of subjects from the cohort who completed germline genetic testing

Clinical Characteristics	***FH* wild type (n=37)**	***FH* pathogenic/likely pathogenic variant (n=12)**	**p-values**
FHx of uterine leiomyoma	13 (35%)	7 (58%)	0.19
FHx of cutaneous leiomyoma	0 (0%)	3 (25%)	0.01
FHx of renal cell carcinoma	5 (14%)	1 (8%)	1
FHx of pheochromocytoma/ paraganglioma	0 (0%)	0 (0%)	n/a