

Table S15. Information of rare variants identified in in-house controls.

Table with columns: Case No., ALS, Gene, Chromosomal position, HGVS, dNA change, Amino Acid change, Het/Hom, Variant Type, Minor allele frequencies (ExAC East Asia, gnomAD East Asia, in-house controls), HGMDB, Clinical significance, Functional predictions (SIFT, PPH2, MA, FATHMM, CADD), prediction, Evidence classification, and ACMG.

Table with columns: Case No., ALS, Gene, Chromosomal position, HGVS, dNA change, Amino Acid change, Het/Hom, Variant Type, dBSNP, Minor allele frequencies (ExAC East Asia, gnomAD East Asia, in-house controls), HGMDB, Clinical significance, Functional predictions (SIFT, PPH2, MA, FATHMM, CADD), prediction, Damage R, and ACMG.

Table with columns for gene names, coordinates, and various data points. The table lists numerous genes such as NT09755, NP18846, NP11126, etc., with their corresponding genomic coordinates and associated values.

