

Monoallelic and bi-allelic variants in *NCDN* cause neurodevelopmental delay, intellectual disability, and epilepsy

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In the originally published version of this article, family 2 was assigned an incorrect aa substitution as a result of the G>A transition in codon 478 of the *NCDN* gene. The aa substitution in individual F2: II.1 should read p.Arg478Gln, not p.Arg478Glu. The error appeared consistently throughout the text, in Table 1, and in Figures 1, 2, and 3; the error has been corrected online and Figures 1, 2, and 3 appear correctly here as well. The authors regret this error.

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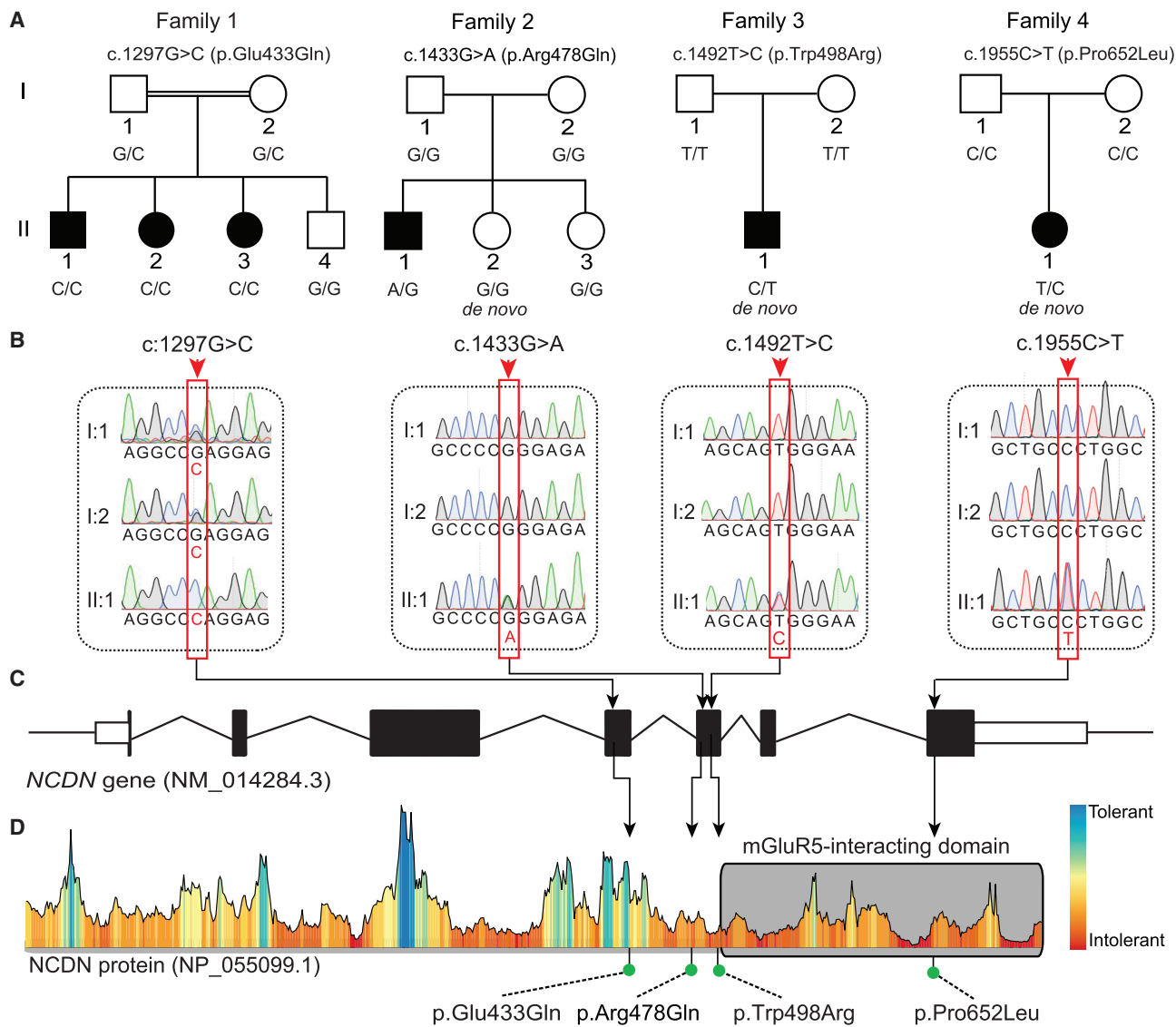


Figure 1. Segregation of rare *NCDN* missense variants in families with neurodevelopmental phenotypes (corrected)

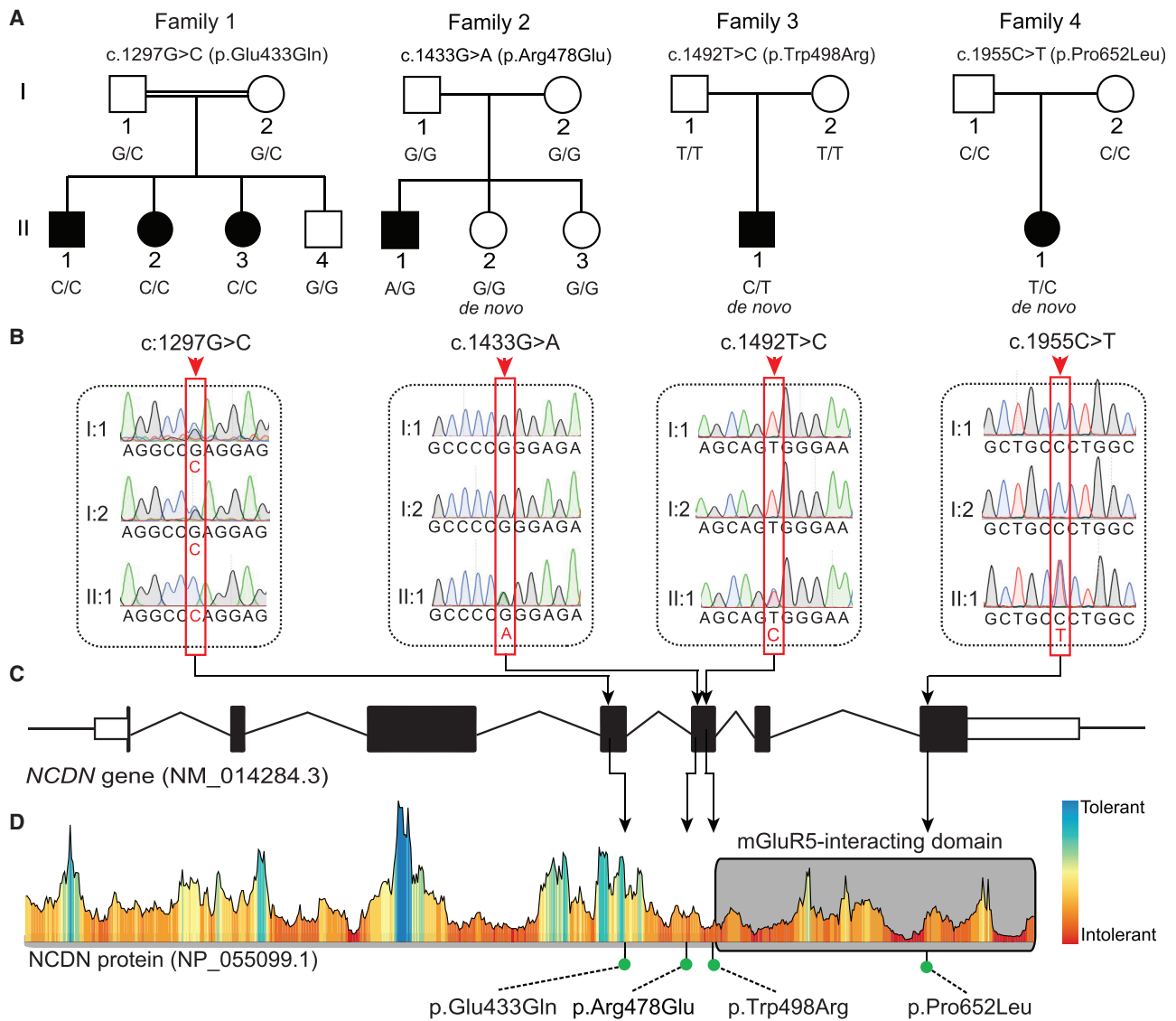


Figure 1. Segregation of rare *NCDN* missense variants in families with neurodevelopmental phenotypes (original)

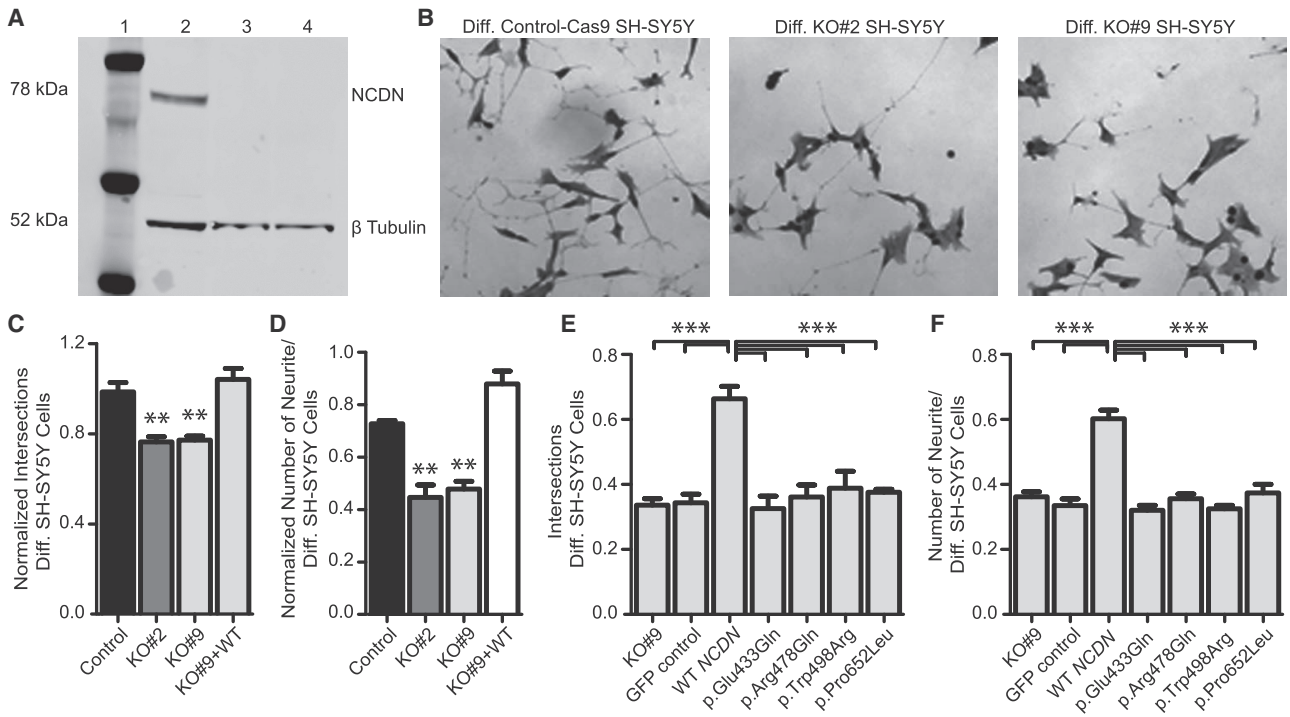


Figure 2. Missense variants in *NCDN* alter length and number of neurites in SH-SY5Y cells (corrected)

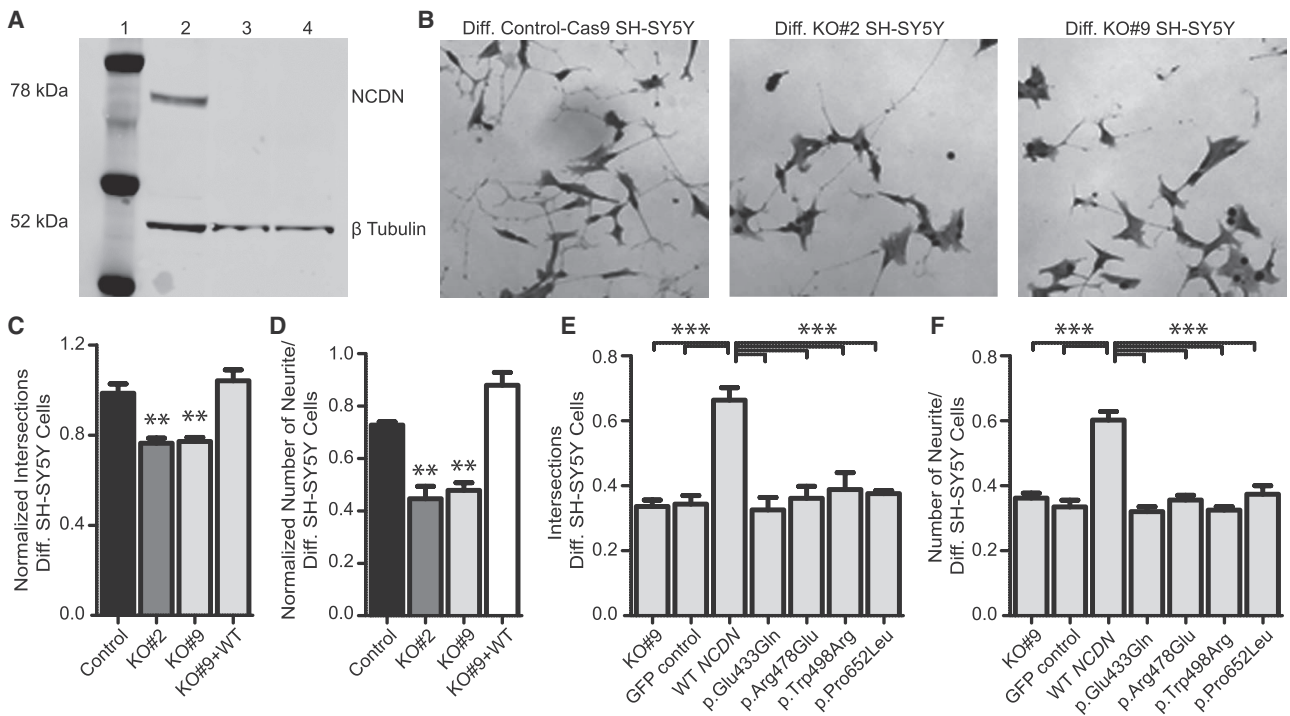


Figure 2. Missense variants in *NCDN* alter length and number of neurites in SH-SY5Y cells (original)

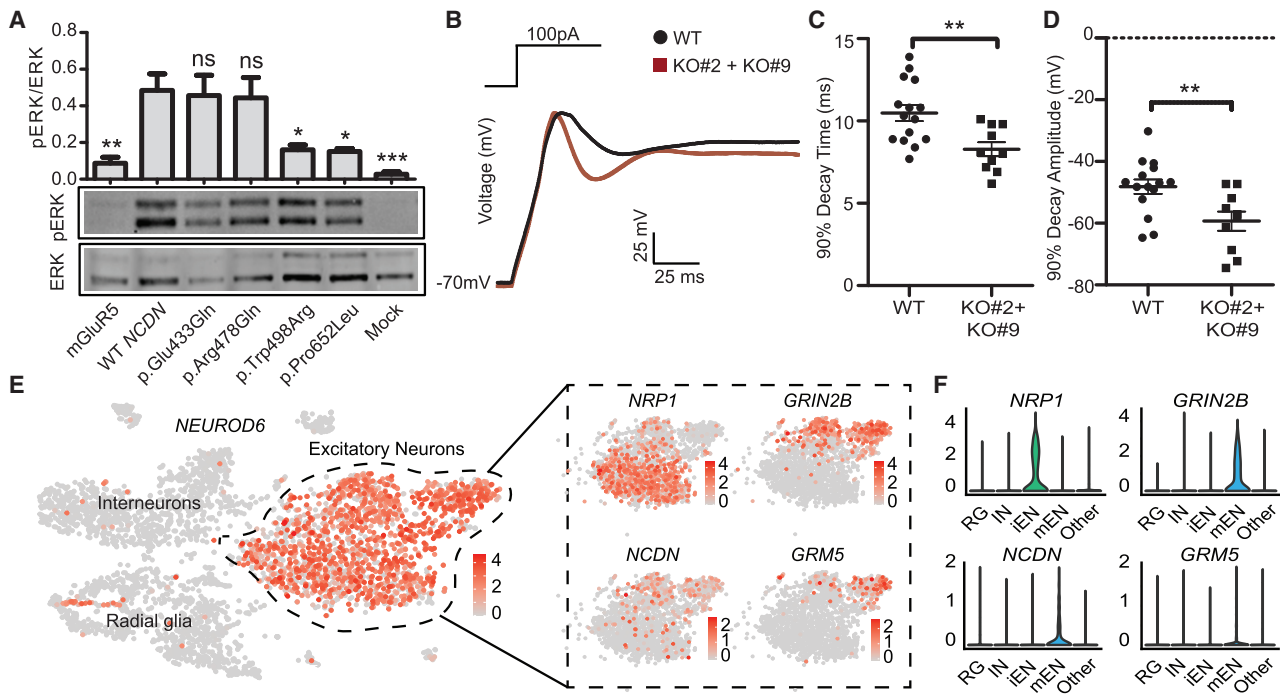


Figure 3. *NCDN* variants interfere with mGluR5 signaling and alter electrophysiological properties of SH-SY5Y cells (corrected)

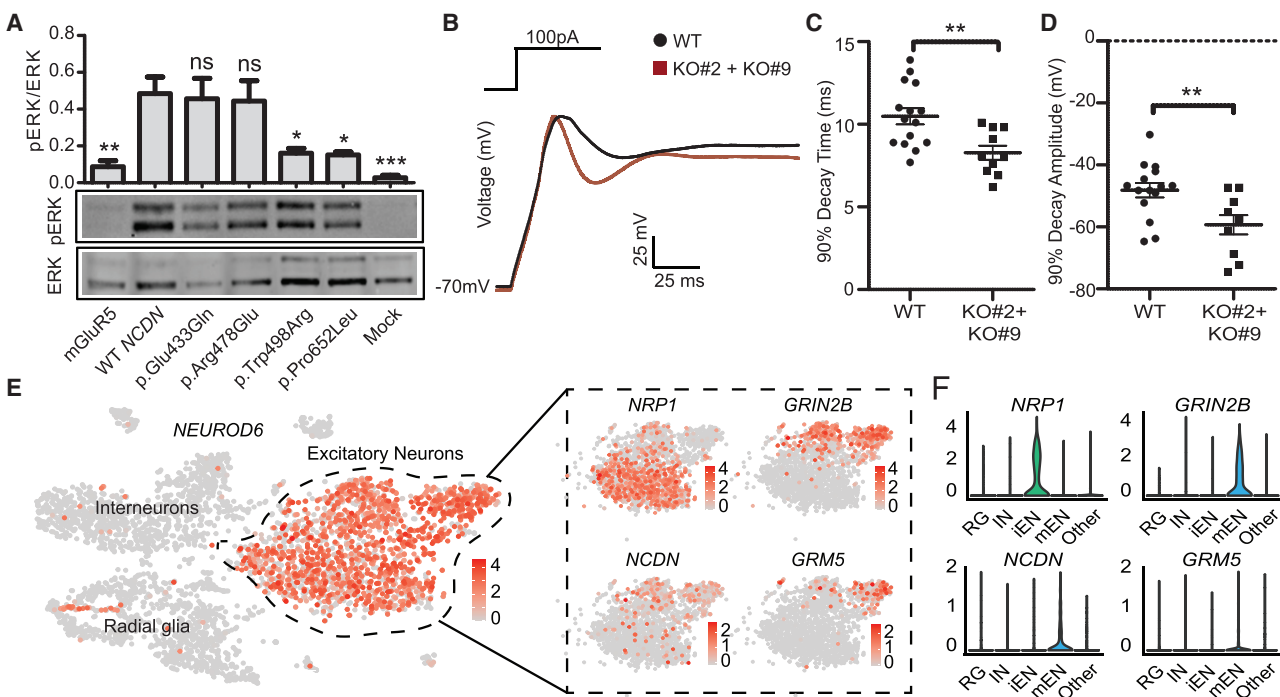


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