

### Mutations in $\beta$ -Tubulin Cause Abnormal Neuronal Migration in Mice and Lissencephaly in Humans

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Figure S1

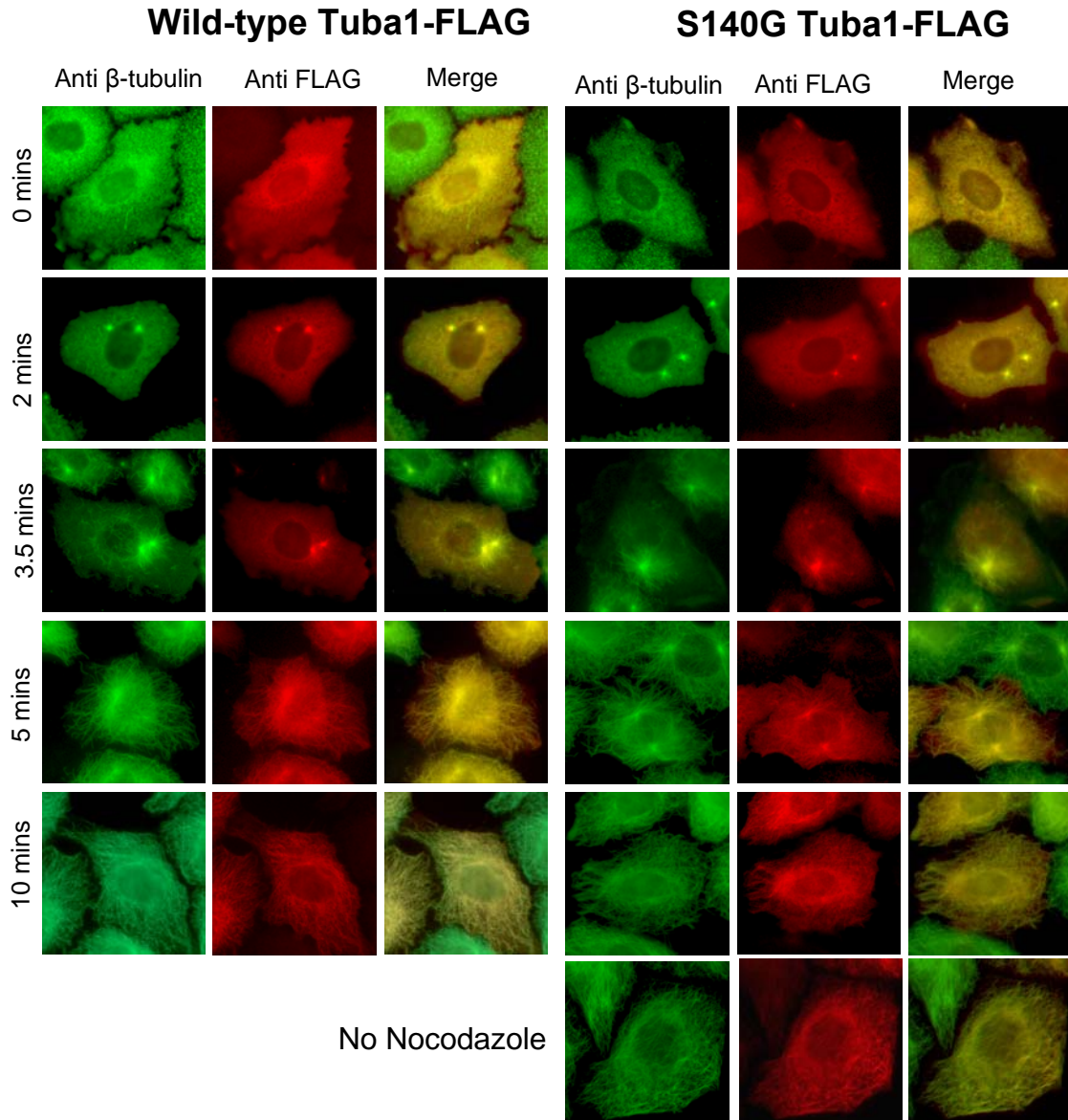


Figure S1: HeLa cells grown on glass coverslips were transfected with constructs encoding FLAG-tagged wild type or S140G mutant Tuba1 as described (see Methods). 36 hours post transfection, the cultures were treated for 2 hours with nocodazole (10  $\mu$ M) to completely depolymerize microtubules. Cells were restored to warm (37°C) drug-free medium and processed for immunofluorescence using anti-FLAG and anti- $\beta$ -tubulin antisera at the times shown on the left of the Figure.

Figure S2

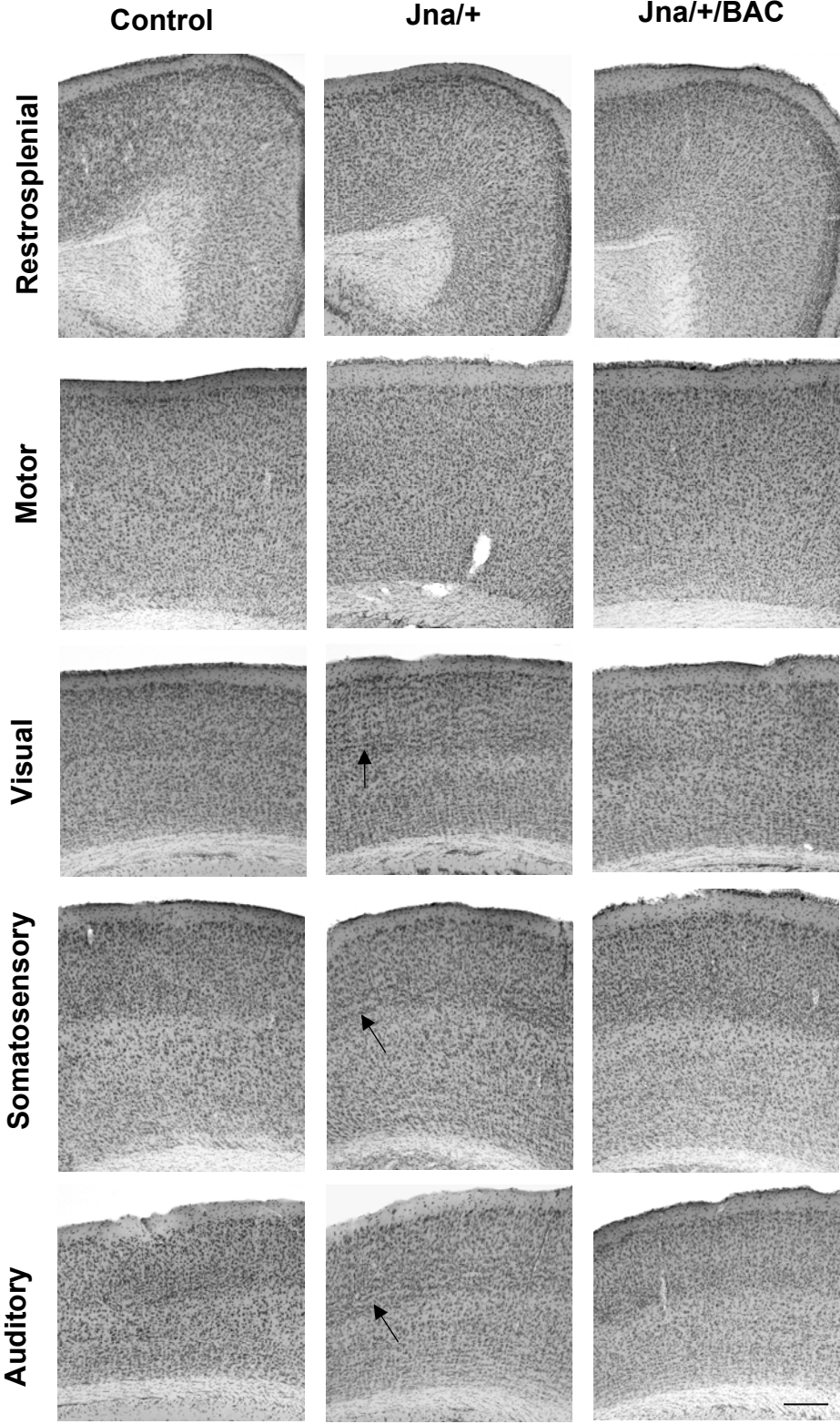


Figure S2: Nissl stains of the retrosplenial, motor, visual, somatosensory and auditory cortices. Arrows indicate wave-like perturbations in layer IV of the visual, somatosensory and auditory cortices. Scale bar shows 200  $\mu$ m.

Figure S3

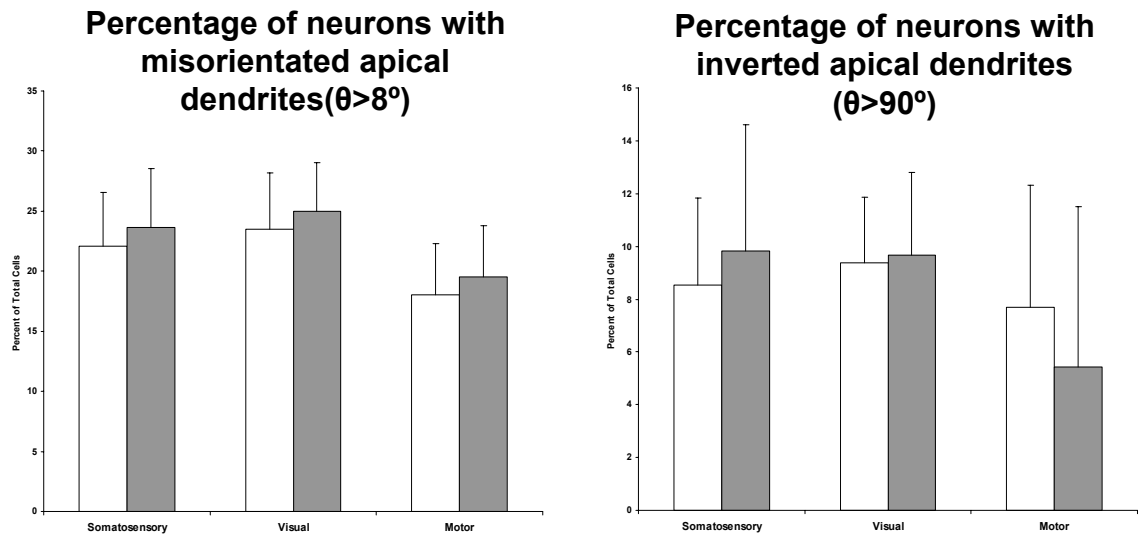
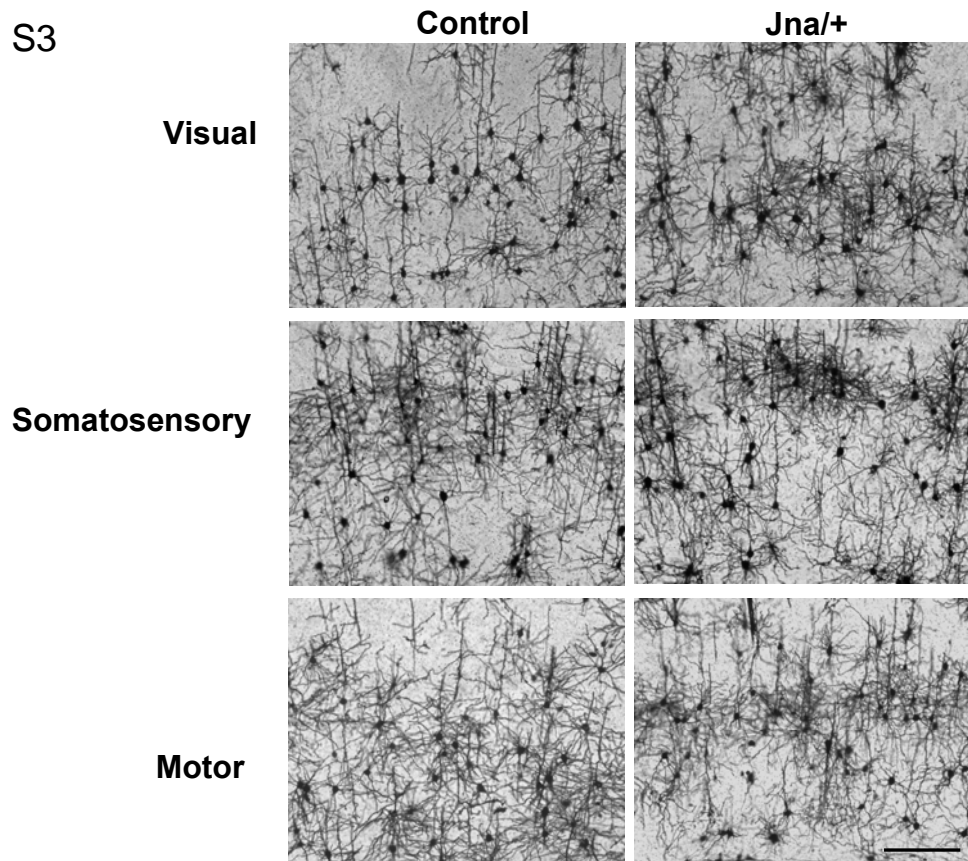


Figure S3: Dendritic orientation in the visual, somatosensory and motor cortices in littermate control (n=3) and *Jna/+* mice (n=3). No differences were observed in the percentage of neurons with misorientated ( $\theta > 8^\circ$ ) apical dendrites or with inverted apical dendrites ( $\theta > 90^\circ$ ). Scale bar shows 200  $\mu\text{m}$ .

Figure S4

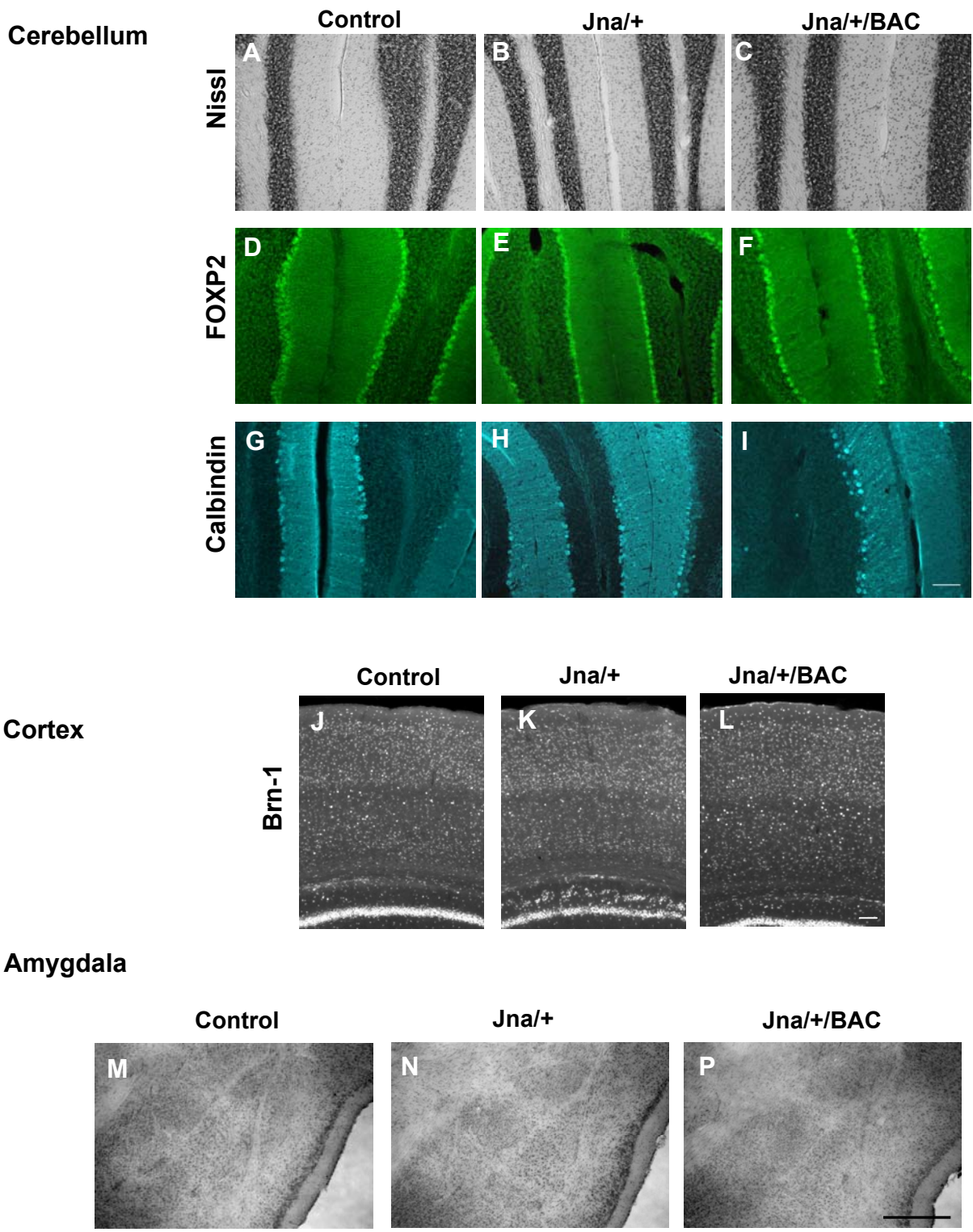


Figure S4: Panels A-I show cresyl violet, FOXP2 and calbindin staining of the cerebellum. No abnormalities in the laminar structure of the cerebellum were observed in Jna/+ mice . Scale bar shows 200  $\mu$ m. Panels J-K show Brn-1 staining in the cortex, confirming an intact laminar structure. Scale bar shows 100  $\mu$ m. Panels M to P show cresyl violet staining of the amygdala (M-P). The amygdala appears normal in mutant animals. Scale bar shows 500 $\mu$ m.