Figure S1. Generation of *Cdh1* gene-trap mice. *a.* A diagram depicting the gene-trapped allele of Cdh1. Arrowheads indicating the position of genotyping primers. *b.* PCR genotyping of embryos from an intercross of $Cdh1^{+/gt}$ mice. *c.* Western blotting analysis of Cdh1 in E9.5 embryo. *d* and *e.* LacZ staining of an E9.5 $Cdh1^{+/gt}$ embryo (*d*) and placenta (*e*). Scale bar, 10 μm.

Figure S2. Mitotic defects in $Cdh1^{gt/gt}$ MEFs. a. Distribution of mitotic phases. b. Micrographs of MEFs stained for tubulin (red) and DNA (blue). Arrows indicate binucleated cells. Scale bar, 40 μ m. c. Quantitation of binucleated cells in b. Results in a and c were from three independent experiments. Error bars are standard deviations.

Figure S3. Suppression of growth defects of *Cdh1* null MEFs by depleting *Ets2*. *a*. Western blot analysis of Ets2 and p16. Full scan of the gel is presented in supplemental figure 5. *b*. Quantitation of the results in *a*. *c*. Growth curve analysis of control- and *Ets2*-shRNA treated *Cdh1* null MEFs at passage 6. Data were obtained from triplicates at each time point. Error bars represent standard deviations.

Figure S3. Potential function of Cdh1 in the brain. **a.** β-gal staining of adult $Cdh1^{+/gt}$ brain. Scale bar, 20 μm. **b.** Immunoblotting of Cdh1 in adult hippocampus. **c.** Plot of fEPSP slope versus fiber volley amplitude. Error bars indicate SEM for ten determinations. **d.** Paired-pulse facilitation in wildtype and $Cdh1^{+/gt}$ mice. Shown were responses to paired pulses in which the fEPSP slope of the response to the second stimulus was expressed as a percentage of the fEPSP slope of the response to the first

stimulus plotted against the interpulse interval of the paired pulses. Error bars indicate SEM for 17 determinations.

Figure S5. Full gel scans of selected Western blots.

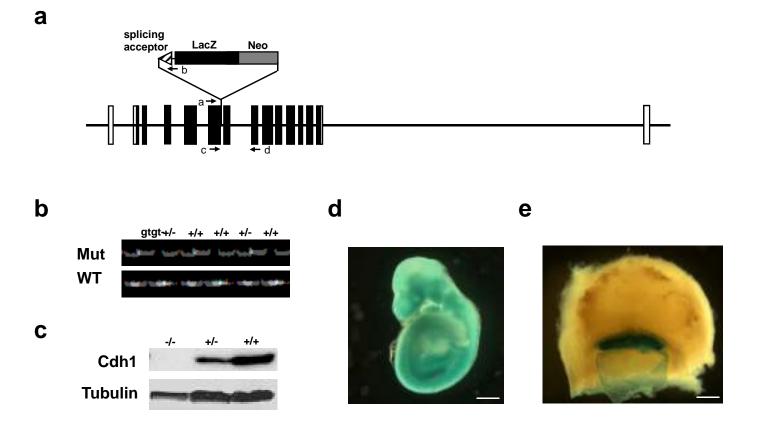


Figure S1

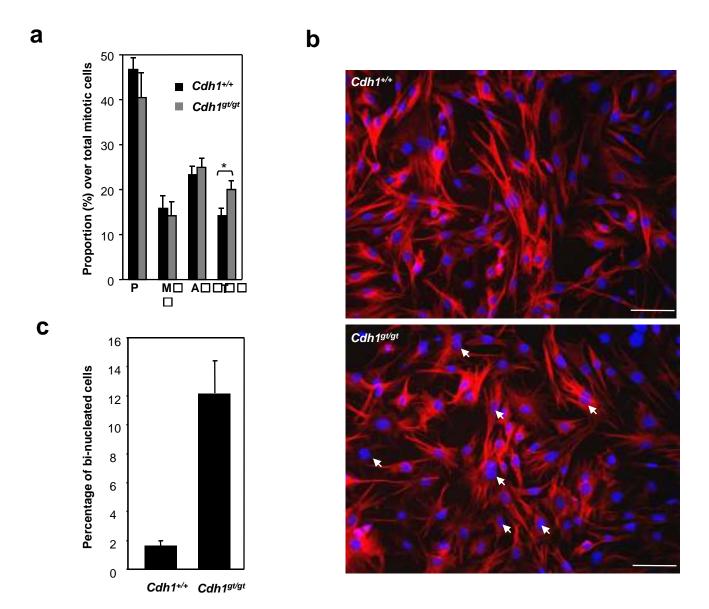
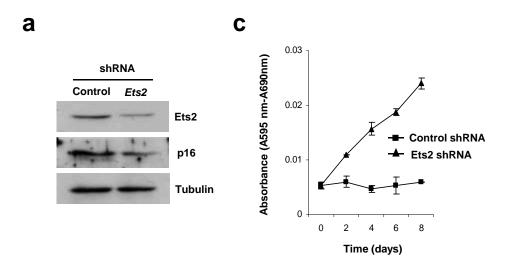


Figure S2



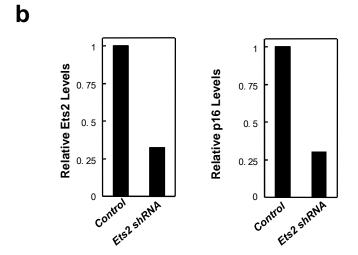


Figure S3

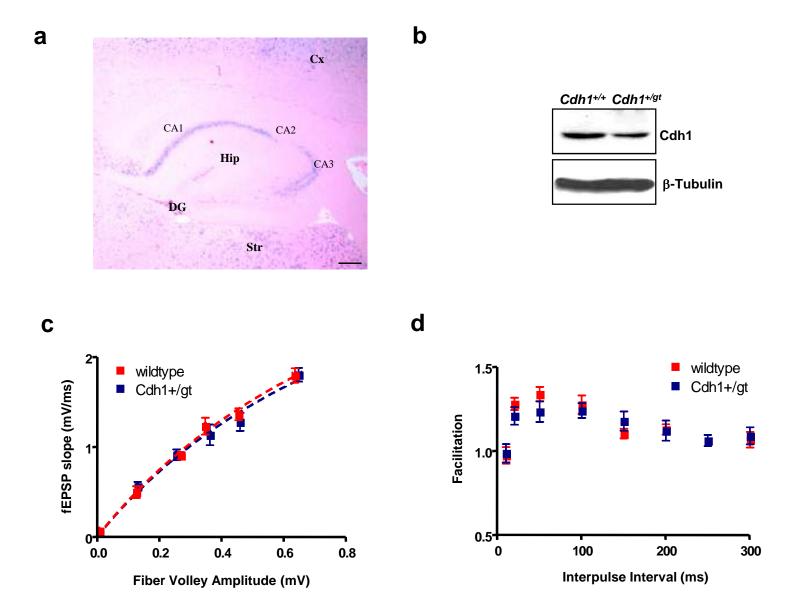


Figure S4

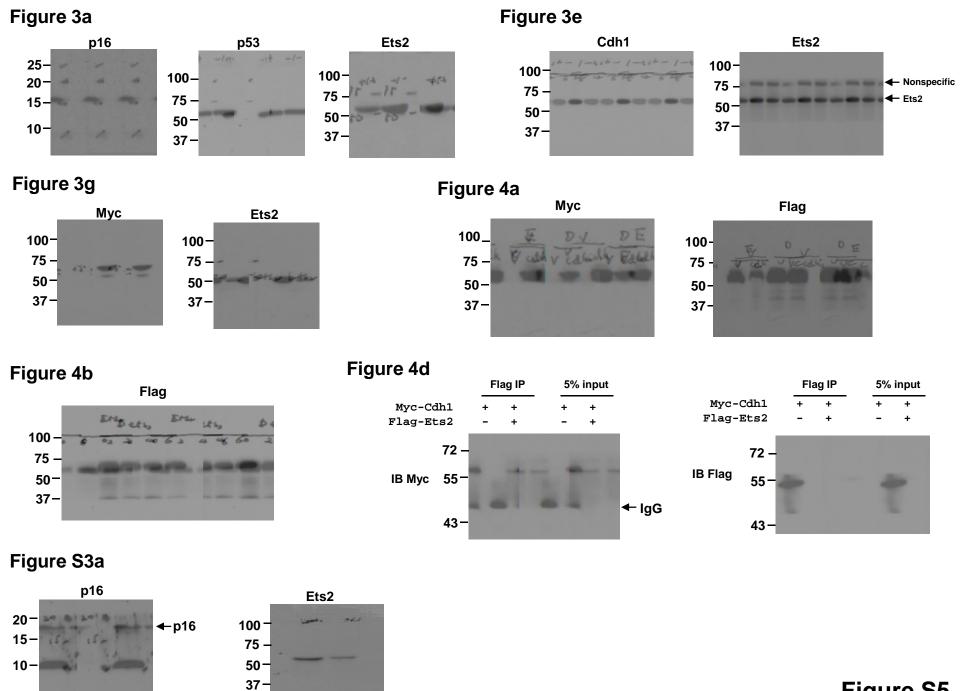


Figure S5