Stem Cell Reports, Volume 9

## **Supplemental Information**

# BACE1 Deficiency Causes Abnormal Neuronal Clustering in the Dentate Gyrus

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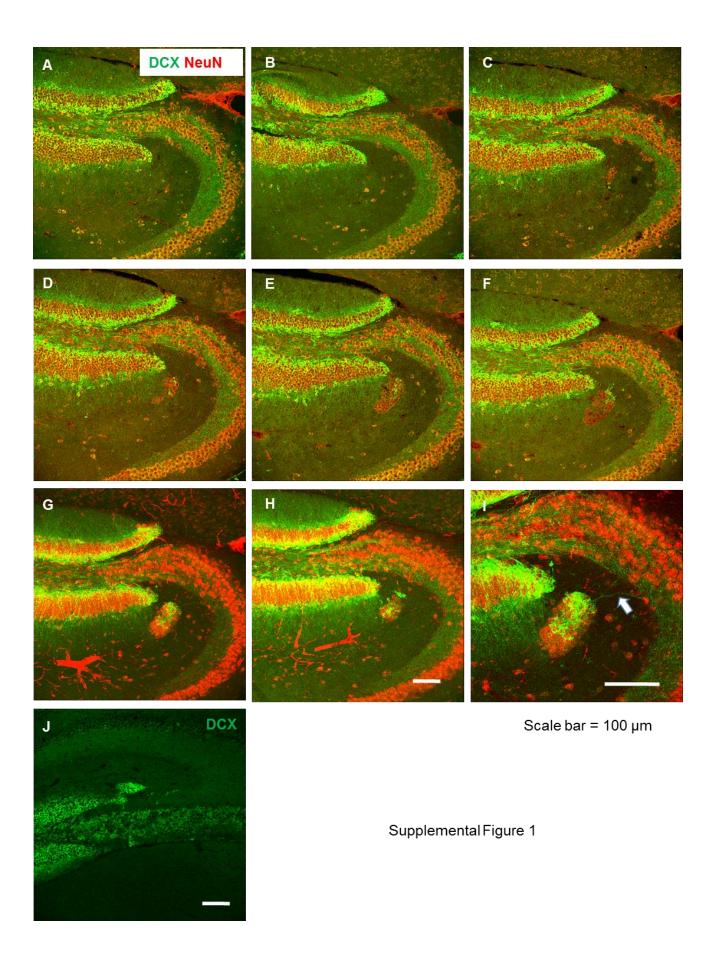
#### **Supplemental Figure and Text**

## BACE1 deficiency causes abnormal neuronal clustering in the dentate gyrus

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Supplemental Figure 1: The presence of neuronal clusters near the mossy fiber side of the DG molecular layer. Continuous brain sections [14- $\mu$ m (A-F) and 50- $\mu$ m (G-H) thick] from one-month-old BACE1-null mice were labeled with DCX and NeuN antibodies. DCX+ clusters were sporadically found in the DG molecular layer, which is close to mossy fibers. The inset in G is shown in I. In another case, clusters are shown in connection with the GCL (J), likely due to a break from GCL or incomplete fusion with GCL. Scale bar is 100  $\mu$ m.

# Supplemental Table 1: Summary of doublecortin-positive clusters seen in BACE1-null mice and related analyses

Phenotype	BACE1 <sup>+/+</sup>	BACE1 <sup>-/-</sup>
Doublecortin-positive neuronal clusters in SPZ/MZ of dentate gyrus	Positive: 6 of 49 BACE1 <sup>+/+</sup> mice. (ages from P6 to P11) The number of clusters in BACE1 <sup>+/+</sup> mice was small and mostly small in size. Clusters have not been detected in mice older than 1 month (12 examined).	Positive: 65 out of a total of 68 BACE1 <sup>-/-</sup> brains (ages P6 to 11M). Negative: 1 from 11M and 2 from P3 brains. Easy to detect in sagittal sections: for example, lateral 0.36 mm-1.92 mm in P20 mice. Some immature clusters in SPZ/MZ can differentiate into mature neurons and may exist throughout life (age of the oldest mice examined in this study was 11M).
Pseudo dentate gyrus	Never detected.	Detected in 6 out of 29 BACE1 <sup>-/-</sup> brains. The age range examined was P11 to 6M (1 in 7 P11, 1 at P20, 2 at P30, 1 at P60, and 1 at P180). The pseudo dentate gyrus was most likely found in brain sections between lateral 0.84 mm to lateral 1.92 mm*.
Reelin-positive cells in the dentate gyrus	The majority of reelin-positive cells resides in the SPZ/MZ region. The number decreased sharply during postnatal development.	The expression and distribution pattern of reelin-positive cells were similar to those in WT mice. The number of reelin-positive cells was similar between WT and BACE1 <sup>-/-</sup> mice in 3 different age groups. DCX-positive clusters were always adjacent to reelin-positive cells, and no reelin-positive cells in clusters were labeled.