## **Supporting Information**

## Bowman et al. 10.1073/pnas.1305411110



**Fig. S1.** Progeny of tamoxifen-inducible Cre at Axin2 ( $Axin2^{CreERT2+}$ ) mouse strain cells labeled embryonically contribute to multiple structures. (A) Embryonic day 6.5 (E6.5)–E7.5 tracing of  $Axin2^{CreERT2+}$ ; *Rosa26* membrane tomato/membrane green fluorescent protein ( $R26R^{mTmG/+}$ ) embryos reveals green fluorescent protein (GFP<sup>+</sup>) cells in the ectoderm and mesoderm, longitudinal section. (B) E8.5–postnatal day 21 (P21) tracing reveals GFP<sup>+</sup> cells in the subgranular and granular zone of the dentate gyrus of the hippocampus. (Scale bar, 50 µm.)



**Fig. 52.** Postnatal subventricular zone (SVZ) cells are broadly Wnt/ $\beta$ -catenin responsive. Representative images of coronal brain sections from *Axin2*<sup>CreERT2/+</sup>; *R26R<sup>mTmG/+</sup>* mice dosed with tamoxifen at P14 and examined at P56. (A) When tamoxifen is administered postnatally GFP<sup>+</sup> cells are found around all surfaces of the ventricle. Postnatally labeled Axin2<sup>+</sup> SVZ neural stem cells (NSCs) are able to produce all classes of olfactory bulb neurons, including calbindin<sup>+</sup> cells (*B*), tyrosine hydroxylase<sup>+</sup> cells (C), and calretinin<sup>+</sup> cells (*D*). Arrowheads point to double labeled cells. DAPI is shown in blue, GFP is shown in green, and other markers in red. (Scale bar, 50 µm.)



**Fig. S3.** *Axin2<sup>CreERT2</sup>* marks multiple radial glial populations throughout the central nervous system. (A) Postnatal labeling reveals GFP<sup>+</sup> cells in the cerebellum that are calbindin<sup>-</sup> (arrowheads point to calbindin<sup>+</sup> Purkinje cells) and (*B*) GFAP<sup>+</sup> (arrowheads point to double labeled fibers), hallmarks of Bergmann glia. (C) GFP<sup>+</sup> Müller glial cells, arrowheads mark radial fiber. DAPI is shown in blue, GFP in green, and other markers in red. (Scale bar, 50 μm.)

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