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data single click to view music videos, 3gp, mpeg, mp3, lyrics and on the go viewing! watermark free! :) !-1607-! I am very happy with the preview performance, the preview quality was good! This will definitely make an announcement more than this video. To sum up I think there are a few things that can be improved, but the video quality was very good. If you're the author of this preview, log in so we can remove this message.pone.0113717-Waligore1]. The number of GFP-positive cells was counted and quantified in a blinded fashion. *In vitro* immunostaining {#s4f} ----- Immunostaining was performed as previously described [pone.0113717-Zhou1]. In brief, frozen sections (10 μ m) were fixed in cold acetone, dried and rehydrated in PBS. For immunostaining, sections were blocked with 5% donkey serum for 1 h at room temperature and incubated with anti-pan-MAP2 antibody (Sigma, Cat. No. SAB4502227, 1:500) and anti-TuJ1 antibody (Covance, Cat. No. MMS-435P, 1:1000) overnight at 4°C. After washes, the sections were incubated with fluorophore-conjugated secondary antibodies (1:200, Jackson ImmunoResearch) and DAPI (Vector Laboratories) for 1 h at room temperature. Fluorescence images were taken with an Axiovert 200 M fluorescent microscope (Zeiss). For the BrdU labeling experiments, the tissues were incubated with 10 μ M BrdU in medium for 30 min at 37°C, and then fixed and processed. Immunostaining was performed as described above. *In situ* hybridization {#s4g} ----- *In situ* hybridization for the *SmNkx2.2* riboprobe was performed as described [pone.0113717-Gu1]. In brief, brain sections (10 μ m) were dewaxed and rehydrated, digested with proteinase K, acetylated with acetic anhydride and treated with 0.2 M triethanolamine, pH 8.0 for 10 min. After washes

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