



(1 column)

Fig. S2 **Pharmacological dissection of CB₁R function evoking corticofugal axon bundling and SCG10 localization in hippocampus.** (A) AM 251 (5 mg/kg) and WIN55,212-2 (5 mg/kg) were administered daily from E5.5-E17.5, fetuses harvested one day later and processed (Keimpema *et al.*, 2010). Corticofugal axon fasciculation was assessed by quantitative L1-NCAM histochemistry. The cross-section thickness (“diameter” (d)) of primary fascicles was determined in $n = 4-7$ male fetuses/group from independent pregnancies (> 300 measurements/group). AM 251 but not WIN55,212-2 induced the coalescing of axons into enlarged fascicles. (A₁) Quantitative morphometry including data on THC effects shown in Fig. 2A₁. (B) Cortical neurons exposed to THC or AM 251 recapitulated the *in vivo* axonal phenotype when cultured in the presence of the drugs indicated for 24h. (B₁) Note that AM 251 was particularly efficacious in promoting the alignment of multiple neurites ($n = 9-13$ observations/ group from $n > 2$ simultaneously processed samples in duplicate experiments). (C,C₁) SCG10 co-localized with CB₁R_s in the primordial hippocampus including migrating neurons (*open arrowheads*, c) on E14.5, and axons (*open arrowheads* in e.g. fimbria (f)) on E18.5. *Abbreviations:* Supplementary Information text. Data were expressed as means \pm s.e.m.; *** $p < 0.001$, ** $p < 0.01$. *Scale bars* = 350 μ m (C,C₁), 50 μ m (A), 5 μ m (B).

Keimpema E, Barabas K, Morozov YM, Tortoriello G, Torii M, Cameron G, Yanagawa Y, Watanabe M, Mackie K, and Harkany T (2010) Differential subcellular recruitment of monoacylglycerol lipase generates spatial specificity of 2-arachidonoyl glycerol signaling during axonal pathfinding. *J Neurosci* **30**: 13992-14007