

Supplemental Figure 20

Fig. S20. Pranlukast abolishes GPR17-mediated Ca^{2+} mobilization but does not blunt Ca^{2+} signaling induced by muscarinic M₃ receptors in primary rat oligodendrocytes. Primary rat oligodendrocytes were pre-treated with pranlukast or the muscarinic antagonist atropine for 30 min prior to stimulation with the indicated concentrations of MDL29,951 or carbachol and intracellular Ca^{2+} mobilization was quantified. Data are mean + S.E.M. of three independent experiments, each performed in triplicate.