



Supplementary Fig. 1. Current-voltage (I–V) relationship of mosapride on 5-hydroxytryptamine (5-HT)<sub>3</sub> receptor current inhibition. (A) Representative traces of 5-HT<sub>3</sub> receptor currents induced by 3  $\mu$ M 5-HT alone at holding potentials of –50–+30 mV (*left*) and 5-HT<sub>3</sub> receptor currents induced by the co-application of 3  $\mu$ M 5-HT with 10  $\mu$ M mosapride at the same holding potentials (*right*). The open horizontal bars indicate the drug application period. (B) I–V plots of averaged normalized peak amplitudes induced by 5-HT with ( $\bullet$ ) or without ( $\circ$ ) mosapride. The reversal potentials were 7.05  $\pm$  0.37 mV (n = 11) for 5-HT alone, and 7.75  $\pm$  0.53 mV (n = 11) for 5-HT with mosapride (The reversal potential was not significantly changed by co-application of mosapride (p = 0.2964, unpaired t-test). (C) Fractional block of mosapride (ratio of peak amplitude of 5-HT with mosapride / peak amplitude of 5-HT alone) as a function of the holding potentials (V<sub>Holding</sub>). The data are expressed as means  $\pm$  standard error of the mean.