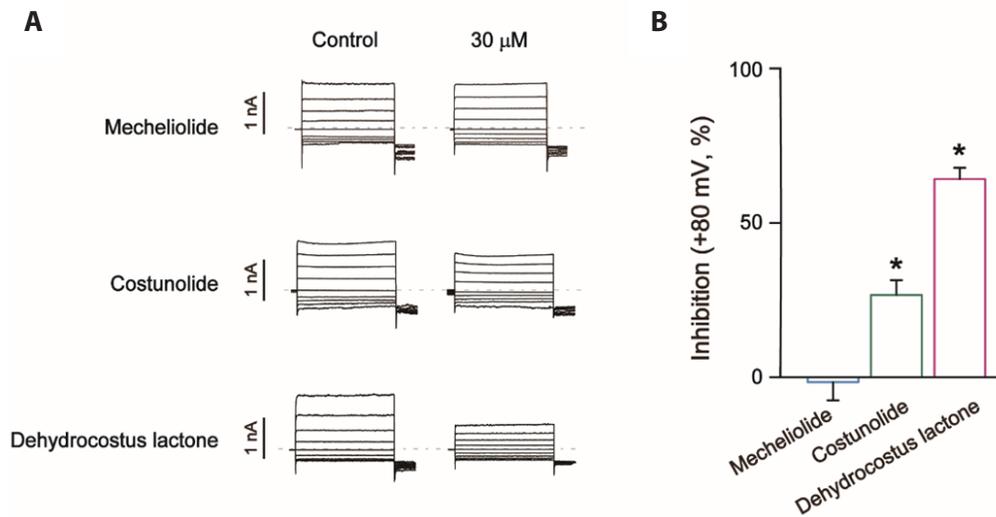


**Supplementary Fig. 1. Typical curve of intracellular  $Ca^{2+}$  fluorescence level after addition of 30  $\mu$ M mecheliolide (A), costunolide (B) and dehydrocostus lactone (C), positive control thapsigargin (TG, 2  $\mu$ M, a blocker of sarco-ER  $Ca^{2+}$ -ATPase) in TMEM16A transfected CHO cells. CHO, Chinese hamster ovary.**



**Supplementary Fig. 2. The inhibitory effects of mecheliolide, costunolide and dehydrocostus lactone on recombinant TMEM16B-mediated CaCC currents in CHO cells.** (A) Representative patch-clamp recordings of whole-cell TMEM16B-mediated CaCC currents evoked by 300 nM  $\text{Ca}^{2+}$  in the absence (control) or presence of 30  $\mu\text{M}$  mecheliolide, costunolide and dehydrocostus lactone in the bath medium. (B) Bar graph showing the inhibitory effects of mecheliolide, costunolide, and dehydrocostus lactone on TMEM16B-mediated CaCCs at +80 mV. The values between groups were statistically compared using the paired Student's t-test. CaCC,  $\text{Ca}^{2+}$ -activated  $\text{Cl}^-$  channel; CHO, Chinese hamster ovary. \* $p < 0.05$ , vs. the current amplitudes in the absence of drugs.