Supplement Figures and legends

Fig. S1. The activation of TRPC6 reporter by OAG is dependent on the extracellular Ca²⁺.

Representative time courses of normalized FRET/CFP emission ratio of TRPC6 reporter in HEK293T cultured in Ca^{2+} free medium with 5 μ M BAPTA under the stimulation of 100 μ M OAG. The data represent the means±SD of each time point from multi-samples by setting the average FRET/CFP ratio of time points before stimulation to 1.0. "n" represents the cell number in each group.

Fig. S2. TRPC6 responded to OAG stimulation after TG pretreatment.

Representative time courses of normalized FRET/CFP emission ratio of TRPC6 reporter in HEK293T with 1 μ M TG pretreatment for 1 hr before stimulated by 100 μ M OAG. The data represents the means±SD from multi-samples as described in Fig. S1. "n" represents the cell number in each group.

Fig. S3. The effect of TG on the PDGF induced ER Ca²⁺ release.

(A) Representative time courses of normalized FRET/CFP emission ratio of ER Ca $^{2+}$ reporter in response to 10 ng/ml PDGF stimulation (time 0) in MEFs. (B) Representative time courses of normalized FRET/CFP emission ratio of D3cpv reporter in MEFs. 1 μ M TG and 10 μ M ATP were added at time 0 and 40 min as indicated, respectively. The data represents the means \pm SD from multi-samples as described in Fig. S1. "n" represents the cell number in each group.

Fig. S4. The activation of TRPC6 reporter by PDGF is dependent on the

extracellular Ca²⁺.

Representative time courses of normalized FRET/CFP emission ratio of TRPC6 reporter in 1 μ M TG pretreated MEFs cultured in Ca²⁺ free medium with 5 μ M BAPTA under the stimulation of 10 ng/ml PDGF. The data represents the means±SD from multi-samples as described in Fig. S1. "n" represents the cell number in each group.

Movie legends

Movie 1. OAG activates TRPC6 reporter but not its mutant in HEK293T cells.

FRET/CFP emission ratio of TRPC6 reporter (left) and its mutant (right) in response to $100~\mu M$ OAG in HEK293T cells with cold and hot colors representing low and high activities of reporter during 12 min imaging. The FRET/CFP emission ratio range is from 1.7 to 3.2.

Movie 2. PDGF activates TRPC6 reporter but not its mutant in MEFs.

FRET/CFP emission ratio of TRPC6 reporter (left) and its mutant (right) in response to 10 ng/ml PDGF in MEFs with cold and hot colors representing low and high activities of reporter during 12 min imaging. MEFs were pretreated with 1 μ M TG for 1 hr before imaging. The FRET/CFP emission ratio range is from 1.6 to 3.0.

Figure S1

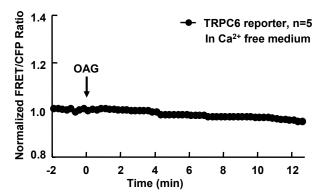


Figure S2

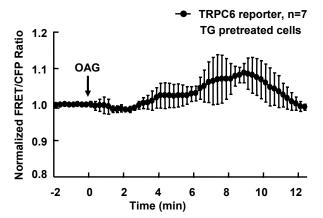
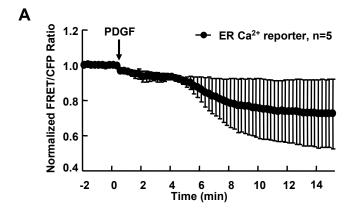


Figure S3



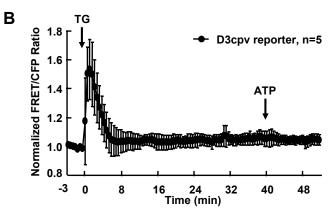


Figure S4

