Supplemental Figure 1. Characterization of GODZ-mediated Ca^{2+} transport. A. summary of Ca²⁺-induced currents in oocytes expressing GODZ cRNA at a holding voltage-clamp of -60 mV. CaCl₂ was added at the concentrations indicated. B, GODZ mediates the transport of a number of other cations but not Mg²⁺. Where indicated SrCl₂, 0.2 mM; BaCl₂, 0.2 mM; or MgCl₂; 0.2 mM was added in place of 0.2 mM CaCl₂. Values are means \pm S.E.M. for 6 individual oocytes. * indicates significance, p<0.01, from calcium alone. C, GODZ does not mediate Mg²⁺ transport. Upper panel, current-voltage (I-V) relationships obtained from linear voltage steps in the presence of 2.0 mM MgCl₂ in control and GODZ expressing oocytes. Oocytes were clamped at a holding potential of -15 mV and then stepped from -150 to +25 mV in 25 mV increments for 2 s at each of the concentrations indicated. Values are means \pm S.E.M. of observations measured at the end of each voltage sweep. *Lower panel*, absence Mg²⁺ flux into GODZ-expressing oocytes. Mg²⁺ fluxes were determined with fluorescence using the Mg²⁺-sensitive dye, mag-fura-2. Results are presented as the 340/385 excitation ratio, which reflects changes in ionized divalent cation concentration. Mag-fura-2 fluorescence ratios were measured in control and GODZexpressing oocytes, at resting potentials, in solutions consisting of nominally magnesium-free solutions and then with 2.0 mM MgCl₂ with interruption and subsequently voltage-clamped at a holding potential of -70 mV, where indicated ¹, GODZ-mediated ⁴⁵Ca²⁺ uptake is inhibited by Mn²⁺, La³⁺, Ba²⁺, and Sr²⁺ but not Mg²⁺. The indicated divalent cations were present at 0.2 mM as the chloride salt in the presence of 0.2 mM CaCl₂. GODZ-mediated ⁴⁵Ca²⁺ uptake is not influenced by external Na⁺. Where indicated NaCl was replaced by choline chloride (*indicated by -Na*⁺). Values are means \pm S.E.M. for 6-25 oocytes. * indicates significance, p < 0.01, from calcium. E, H⁺ dependence of GODZ-mediated Ca²⁺ uptake. ⁴⁵Ca²⁺ uptakes performed at the given external pH values. Values are means \pm S.E.M. for five individual experiments. * indicates significance, p<0.01, from pH 7.5. F, GODZ-mediated Ca²⁺ currents were not inhibited by the calcium channel blocker, nifedipine. Nifedipine, 100 FM, (*indicated by* +*Nifed*) was added 30 min prior to voltage-clamp. Current measurements were determined as in Fig 1A and data are mean values of the largest observed currents \pm S.E.M for 3-5 oocytes. * indicates significance, p<0.01, from calcium alone.

Fig.S1

