LEGENDS TO SUPPLEMENTAL DATA

Figure S1. Subcellular redistribution of HIP14 protein in response to magnesium using COS-7 epithelial cells. Immunofluorescence staining of HIP14-GFP fusion protein of transiently expressing COS-7 cells. A, Golgi localization of HIP14. Cells were cultured in normal magnesium, fixed and incubated with GFP antibody (left panel) and the Golgi marker, GM130, (center panel). The merged image is given in right panel. B, Golgi localization of HIP14 in cells cultured in low magnesium media. Cells fixed and incubated with GFP antibody (left panel) and the Golgi marker, GM130, (center panel). The merged image is given in *right panel*. C, submembrane location of post-Golgi HIP14 protein. Cells were cultured in normal magnesium. HIP14 (left panel), phaloidin (center panel), and the merged image (right panel) are shown. Note the predominant submembrane localization of post-Golgi HIP14 protein. D, additional cell images from independent cell preparations at 2-fold digital enlargement of S2C. E, increase in HIP14 protein in post-Golgi vesicles and plasma membrane of cells cultured in low magnesium media. HIP14 staining (left panel), HIP14 merged with phaloidin (center panel), and graphic summary of increase in subplasma membrane HIP14 staining with low magnesium (right panel); values are means ± S.E. in arbitrary units, n=76 cells. Note the increase in post-Golgi vesicle and plasma membrane HIP14 protein (indicated by arrows). F, Additional cell images from independent cell preparations at 2-fold magnification of Fig. S2E. G, absence of HIP14 protein in early recycling endosomes. COS-7 cells were cultured in normal media with normal magnesium concentration. HIP14 staining (left panel), Rab5 (center panel), and the overlay image (right panel). H, absence of HIP14 protein in early recycling endosomes. COS-7 cells were cultured in media with low magnesium. HIP14 staining (left panel), Rab5 (center panel), and the overlay image (right panel).

Fig. S1

