Supplementary Information

Rab27A Regulates Transport of Cell Surface Receptors Modulating Multinucleation and Lysosome-Related Organelles in Osteoclasts

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Gene title	Log ratio
calcitonin receptor	3.84
cathepsin K	3.29
transmembrane 7 superfamily member 4 (DC-STAMP)	3.14
carbonic anhydrase 2	3.47
acid phosphatase 5, tartrate resistant (TRAP)	2.41
Rab27A	1.26
Rab27B	-1.09

(a)



Supplementary Figure 1

(a)	Rab27A	GM130	Merge
(b)	Rab27A	Calnexin	Merge
(C)	Rab27A	LAMP2	Merge

Supplementary Figure 2





Supplementary Figure 3



Supplementary Figure 1. Up-regulation of Rab27A during osteoclastogenesis. (a) List of up-regulated transcripts in rapid differentiating OCLs compared to slow differentiating OCLs. BMMs were cultured on plastic plate (rapid differentiating conditions) and dentin (slow differentiating conditions) for 72 h in the presence of M-CSF (30ng/mL) and RANKL (50ng/mL). Total RNA from these cells was analyzed by *Affymetrix Microarray* system. (b) Quantitative RT-PCR determination of mRNA expression levels of Rab27A and Rab27B in MC3T3, RAW-D, and RANKL-stimulated RAW-D cells (OCLs). The data are represented as mean \pm Standard Deviation (S.D.) of values from three independent experiments. **P* < 0.05 for the indicated comparisons.

Supplementary Figure 2 Subcellular localization of GFP-Rab27A expressed RAW-D cells. The cells on glass cover-slips were fixed, permeabilized with 0.2% Tween-20 in PBS, and then allowed to react with (a) anti-GM130 (marker for the Golgi), (b) anti-calnexin (marker for ER), (c) anti-LAMP2 (marker for late endosomes/lysosomes) antibodies on the glass cover slips. After being washed, the samples were incubated with a fluorescence-labeled secondary antibody and then were visualized by confocal laser microscopy. Bar: 10 μm