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Supplemental Figure 2. *Localization of MARCH1 mutants*. Confocal microscopic analysis of DC2.4 cells stably expressing the indicated mutants. MARCH1 staining is shown *versus* the indicated markers: EEA-1 (early endosome) and Derlin-1 (ER). Note that a lower threshold was applied uniformly to each set of samples (imaged for a given marker) based on negative controls (background subtraction). The signal for MARCH1 (HA) and the marker protein in each set of images was normalized by setting the brightest pixel in each image (for each fluorescence channel) as the maximum signal (histogram-stretching) in order to facilitate visualization and comparison across fields.

Figure S1

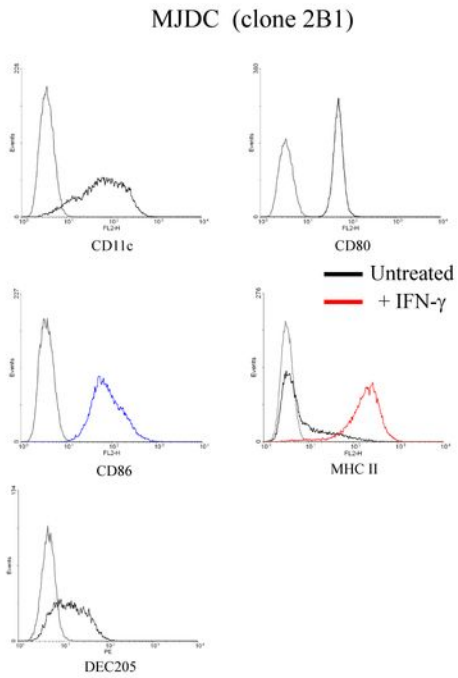
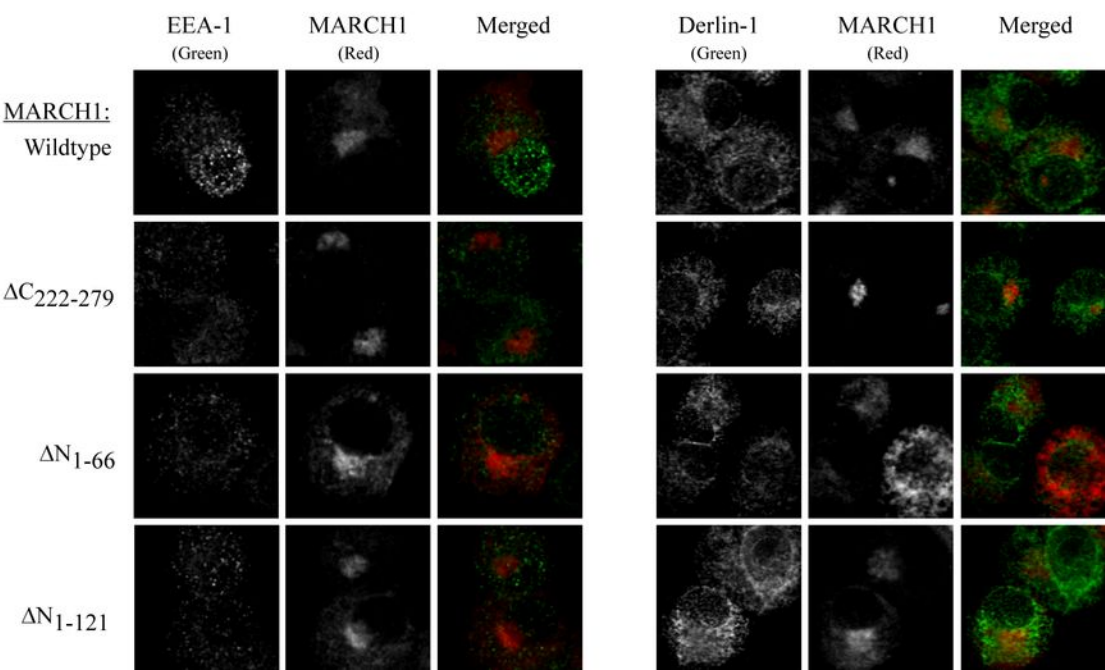


Figure S2



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