
Casanova Iota Idm Patch

In 3D confocal imaging, cd13-containing vesicles formed a loose apical membrane with a single, long tube-like structure. This was consistent with the tracking of individual vesicles that established that cd13-containing vesicles were associated with the apical membrane before lumen formation (supplementary Fig. 3f). At the lumen initiation site, cd13 was distributed throughout the apical membrane, resulting in a diffuse distribution, and had little vesicular association (supplementary Fig. 3g). Although cd13 was observed to colocalize with rab11 at the cell-cell adhesion, this was lost as the apical membrane expanded, indicating that cd13 is not associated with recycling vesicles at this stage (supplementary Fig. 3g). Together, these results indicate that cd13-containing vesicles are recruited to the apical membrane prior to lumen formation. We next examined the relationship between cd13 and rab11. To visualize the relationship between rab11, cd13, and e-cadherin in control and cd13-depleted caco-2 cells, we observed rab11 localization to e-cadherin patches in control caco-2 cells, interestingly, these rab11 patches were not coincident with cd13-containing vesicles in control cells (Fig. 6, i). In cd13-depleted cells, rab11 localized in patches with cd13 and e-cadherin, the rab11 patches were not coincident with cd13-containing vesicles, and were also not coincident with e-cadherin (Fig. 6, j). These data indicate that rab11 localizes to apical sites that contain cd13 prior to lumen formation. In contrast to rab11, we did not observe cd13-containing vesicles that colocalize with rab11 at the apical.

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