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Quantifying the LAMP1 positive puncta

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Protocol status: Working

We use this protocol and it's working

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Abstract

This describes quantifying the LAMP1 positive puncta.

Attachments



[724-1817.docx](#)

14KB

Materials

Tools required

- ImageJ/Fiji



Quantifying the LAMP1 positive puncta

- 1 Select cells of interest with similar shapes and without saturation, and isolate single ROI.
- 2 Apply auto threshold to total stacks of the images.
- 3 Estimate the number of LAMP1 positive vesicles in each ROI by using "Analyze particle" in ImageJ/Fiji.
- 4 Plot the results by calculating the mean number of LAMP1 positive puncta per each cell.