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containing 0.1% saponin or 0.5% formaldehyde for 20 min. The cells were
incubated in 0.4% Triton X-100 for 30 min and then incubated in blocking buffer
(PBS containing 1% BSA) for 30 min, and FITC (green) or TRITC (red) conjugated
secondary antibody in blocking buffer was added to the cells for 30 min. The cells
were mounted on a slide with mounting medium and the images were captured
using a fluorescence microscope (Nikon, Tokyo, Japan). 4.8. Immunofluorescence
Staining {#sec4dot8-ijms-17-00817} ----- HepG2 and L-02 cells
cultured on coverslips were washed twice with ice-cold PBS and fixed in 4%
paraformaldehyde for 15 min. The fixed cells were permeabilized with 0.1% Triton
X-100 in PBS for 30 min and then washed with PBS for 15 min at room
temperature. The cells c6a93da74d

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