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Cover photo. Mechanical permeabilization and RhoDex entrapment in cultured hepatocytes. Cultured hepatocytes were loaded with green-fluorescing MTG and incubated in ICB (A) RhoDex ( $400~\mu\text{M}$ ) was then added followed by mechanical perturbation using a micropipette. Afterwards, RhoDex penetrated the cytoplasm and nuclei (B) After 120 s, the medium was replaced with ICB containing both RhoDex and DIDS ( $30~\mu\text{M}$ ), the latter a VDAC inhibitor to entrap RhoDex in the mitochondrial intermembrane space. After another 60~s, the hepatocytes were washed with ICB containing only DIDS to remove unbound RhoDex. After another 300~s, images of MTG and RhoDex fluorescence were collected to identify RhoDex remaining within mitochondria (C) Higher magnification (lower panels) shows retention of red-fluorescencing RhoDex (E) in MTG-labeled mitochondria (D) as evident in the overlay (F) White arrows identify strongly RhoDex-labeled structures that do not co-label with MTG. For more information please see the article by E. Holmuhamedov, J.J. Lemasters in this issue.



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