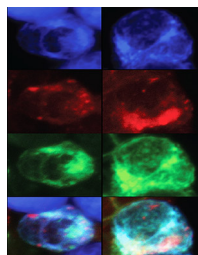


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COVER IMAGE



*Cover photograph:* Live cell fluorescent confocal microscopy of wild-type (left panels) and *SURF4*-deleted (right panels) reporter cells expressing green fluorescent protein-tagged erythropoietin (EPO), mCherry-tagged alpha1-antitrypsin, and the endoplasmic reticulum (ER) marker ERoxBFP. Findings demonstrate accumulation of EPO in the ER of *SURF4*-deleted cells, consistent with the dependence of EPO on *SURF4* for exit from the ER. (See related article in December 2020, vol. 40, no. 23, e00180-20.) (Copyright © 2020 American Society for Microbiology. All Rights Reserved.)

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**ERRATUM**

**Erratum for Caslini et al., “MLL Associates with Telomeres and Regulates Telomeric Repeat-Containing RNA Transcription”** e00566-20

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**AUTHOR CORRECTION**

**Correction for McMullen et al., “Deletion of Ribosomal S6 Kinases Does Not Attenuate Pathological, Physiological, or Insulin-Like Growth Factor 1 Receptor–Phosphoinositide 3-Kinase-Induced Cardiac Hypertrophy”** e00635-20

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