Correction

CELL BIOLOGY

Correction for "Itch E3 ubiquitin ligase regulates large tumor suppressor 1 stability," by King Ching Ho, Zhonghua Zhou, Yi-Min She, Alex Chun, Terry D. Cyr, and Xiaolong Yang, which appeared in issue 12, March 22, 2011, of *Proc Natl Acad Sci USA* (108:4870–4875; first published March 7, 2011; 10.1073/pnas.1101273108).

The authors wish to note: "There is an error on the fourth panel of Fig. 2, the Ponceau S staining figure. We have now replaced it with a correct figure." As a result, Fig. 2 has been corrected. The corrected figure and its legend appear below.

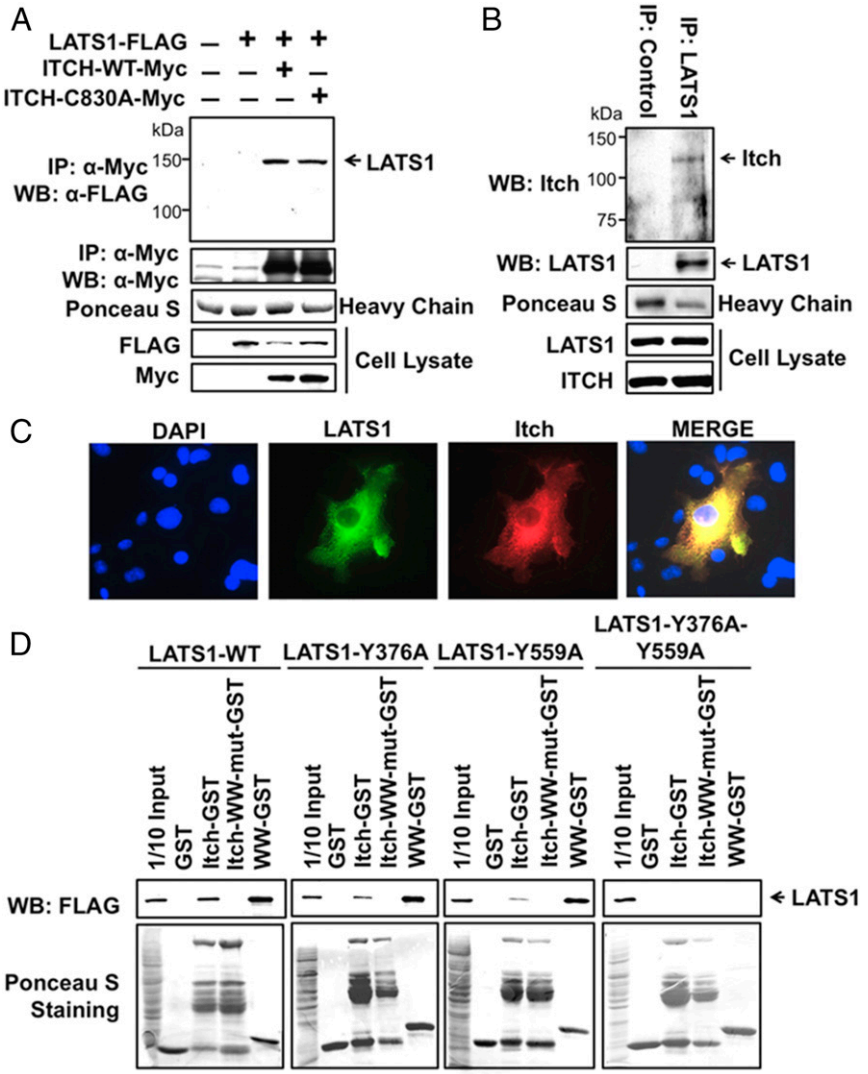

Fig. 2. Interaction of LATS1 and Itch in vivo and in vitro. (*A*) Interaction of ectopically expressed LATS1 and Itch. COS7 lysates expressing either LATS1-FLAG alone or together with Itch-Myc or Itch-C830A-Myc were immunoprecipitated with anti-Myc antibody, followed by Western blotting with anti-FLAG antibody. Ponceau S staining of antibody heavy chain indicates equal amounts of anti-Myc antibody were used. (*B*) Interaction of endogenous LATS1 and Itch. Protein lysates from MDA-MB-231 cells were immunoprecipitated with either control anti-FLAG antibody or anti-LATS1 antibody, followed by Western blotting with anti-Itch antibody. (*C*) Immunostaining analysis of LATS1 and Itch. LATS1-FLAG and Itch-Myc were cotransfected into COS7 cells, followed by immunostaining with anti-FLAG and anti-Myc primary antibodies and AF488 anti-mouse IgG and AF555 anti-rabbit IgG secondary antibodies. (*D*) GST-pulldown analysis of interaction of LATS1 and Itch in vitro. COS7 lysates expressing wild-type (LATS1-WT-FLAG), single-PPxY mutants (LATS1-Y376F-FLAG or LATS1-Y559F-FLAG), or double PPxY mutant (LATS1-Y376F-Y559F-FLAG) of LATS1 were pulled down with GST, GST-Itch, GST-Itch-WW-mutant, or GST-WW, followed by Western blotting for LATS1-FLAG using anti-FLAG antibody. 1/10 input (10 μg) represents 1/10 of protein lysate (100 μg) used for GST pulldown.

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