

SIRT1 deacetylates ROR γ t and enhances Th17 cell generation

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The authors regret that the labels “Input” and “IP: α -Flag” were switched in the original publication of Fig. 2 (E and F). This has now been corrected in the HTML and PDF versions of the paper. The corrected image also appears below.

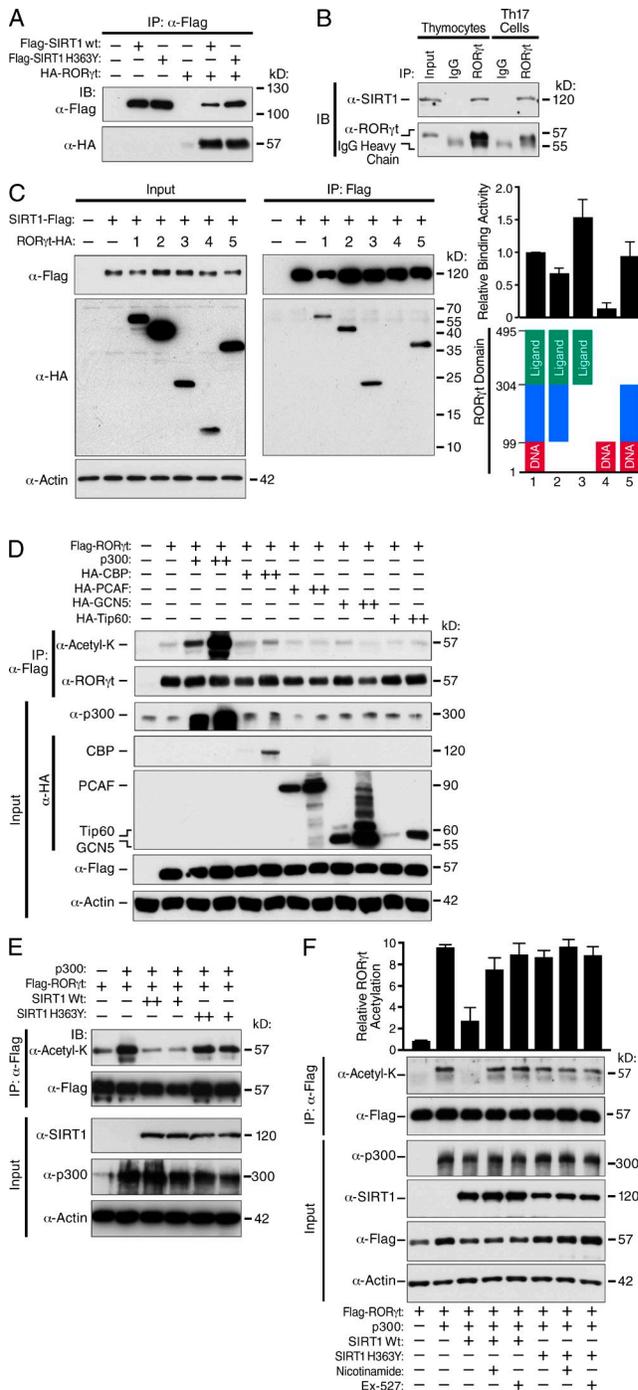


Figure 2. SIRT1 interacts with ROR γ t. (A) Flag-tagged WT or H363Y mutant SIRT1 was immunoprecipitated from transfected 293T cells and probed as indicated. (B) ROR γ t was immunoprecipitated from thymocytes and Th17 cells, and probed with antibody against SIRT1. (C) Immunoprecipitation using lysates of 293T cells co-transfected with constructs encoding SIRT1 and various deletion mutants of ROR γ t. Relative binding was calculated by normalizing the ratio of immunoprecipitated ROR γ t/SIRT1 to the ratio of input ROR γ t/SIRT1. (D) Acetylation of Flag-tagged ROR γ t immunoprecipitated from 293T cells transfected with various acetyltransferases and Flag-ROR γ t. (E and F) Acetylation of Flag-ROR γ t co-transfected with p300 and WT or H363Y mutant SIRT1, in the absence (E) or in the presence (F) of nicotinamide and Ex-527. Equal amounts of Flag-ROR γ t were loaded (D–F). Representative data are shown from four (A), three (B and E), and two (D) independent experiments, and combined data are shown from three (C and F) independent experiments with error bars representing \pm SEM.