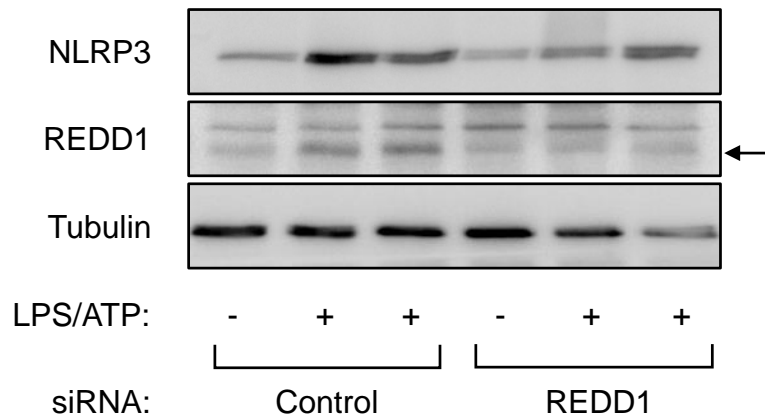
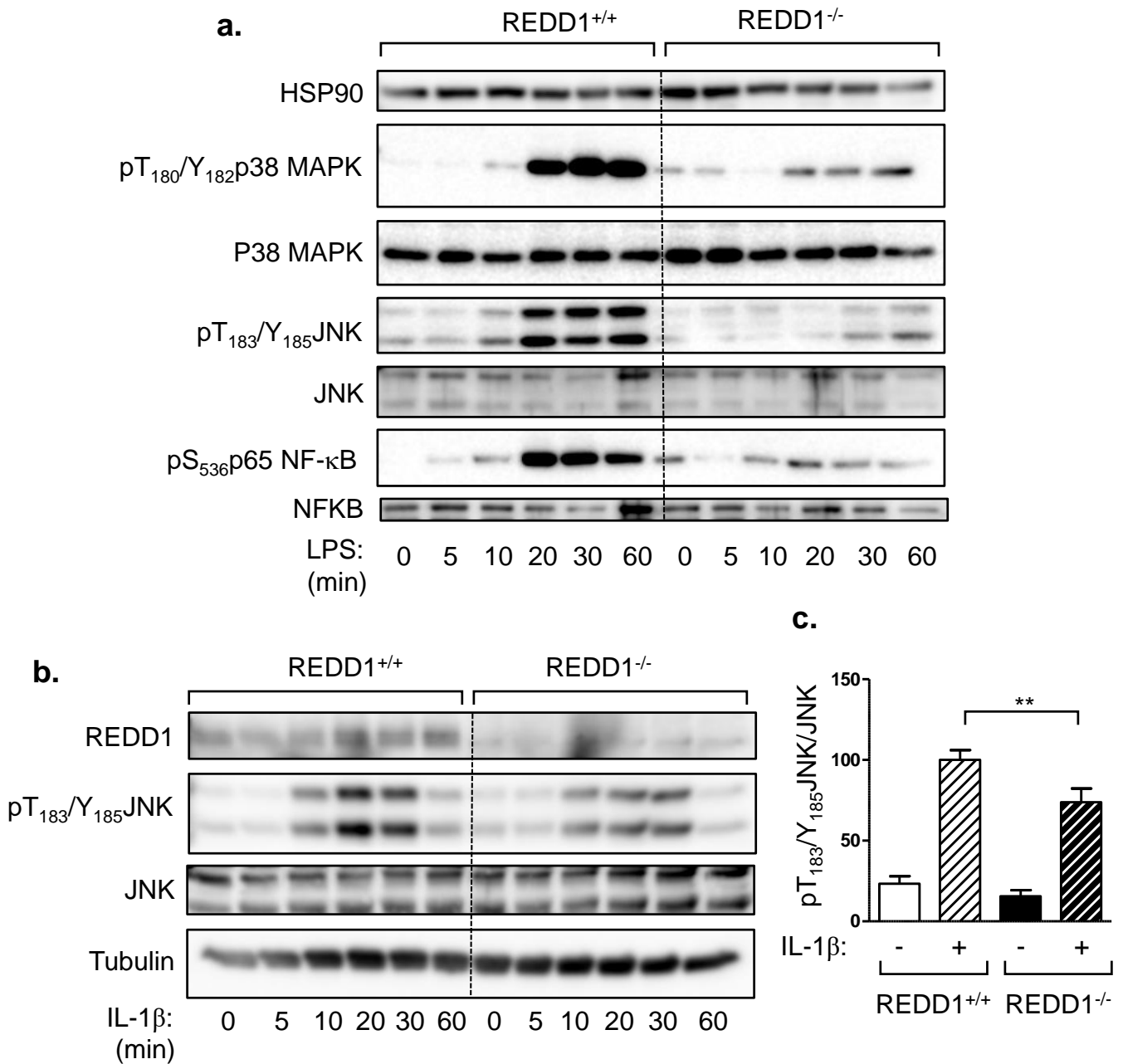


Implication of REDD1 in the activation of inflammatory pathways

Faustine Pastor, Karine Dumas, Marie-Astrid Barthélémy, Claire Regazzetti, Noémie Druelle, Pascal Peraldi, Mireille Cormont, Jean-François Tanti, Sophie Giorgetti-Peraldi

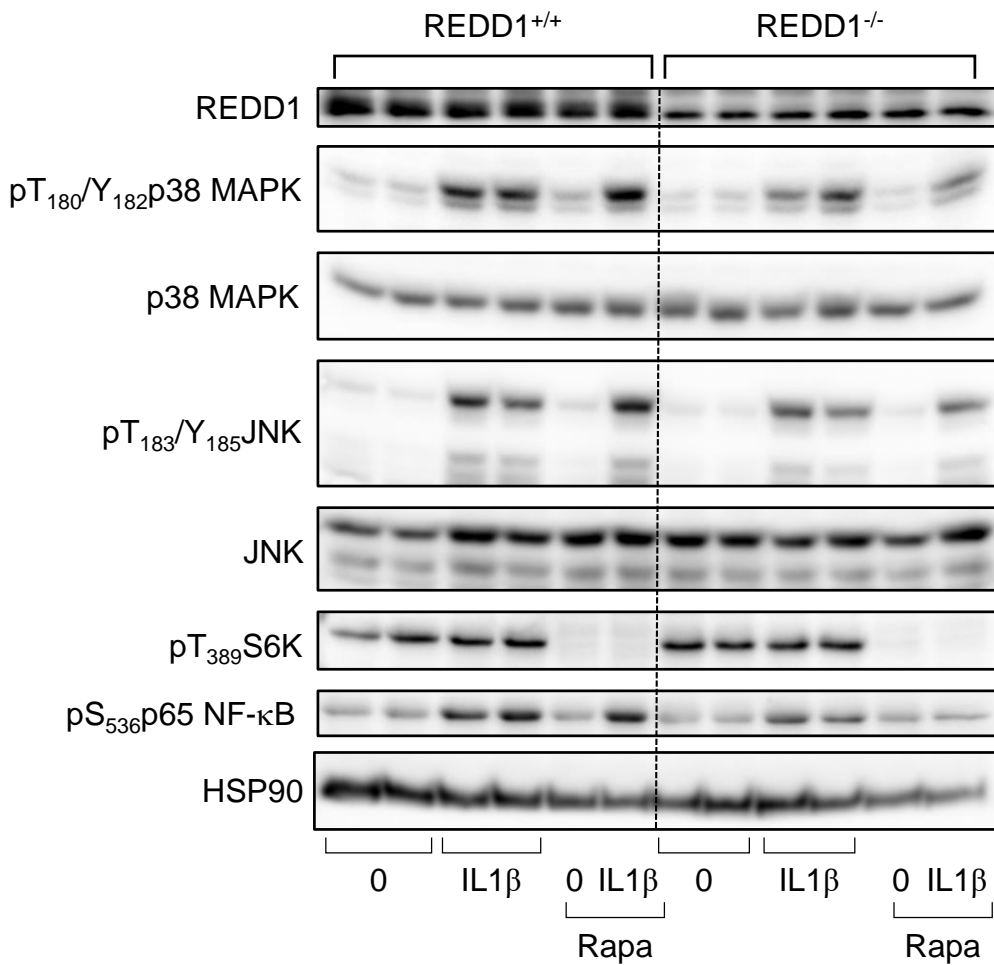


Supplementary Figure S1: J774 macrophages were transfected with siRNA against REDD1 and analyzed by immunoblots with indicated antibodies



Supplementary Figure S2: Activation of p38 MAPK, JNK and NF-κB was impaired in MEF REDD1^{-/-} in response to LPS and IL-1β

REDD1^{+/+} and REDD1^{-/-} MEF were stimulated with LPS (100 ng/ml) (a) or IL-1β (20 ng/ml) (b) for indicated periods of time. Cell lysates were analyzed by immunoblots with indicated antibodies. c: Quantification of pT₁₈₃/Y₁₈₅ JNK in response to IL-1β stimulation for 20 minutes (n=4 independent experiments in duplicate, ** p<0.01).



Supplementary Figure S3: Inhibition of inflammation in REDD1^{-/-} cells is mTORC1 independent

REDD1^{+/+} and REDD1^{-/-} MEF were treated with rapamycin (rapa) 40 nM for 45 minutes before being stimulated with IL-1 β for 20 minutes. Cell lysates were analyzed by immunoblots with indicated antibodies.