

EDEM1. Immunoprecipitation (IP) and immunoblotting (IB) for detecting proteins interacted with TRAP complex. In the upper six panels, proteins from HEK293 were extracted using lysis buffer containing 3% digitonin, and 1 mg of cell lysate was used for IP for detecting endogenous proteins interacted with TRAP complex. IPs were performed with (+) or without (-) anti-TRAPα antibody. Immunodetections were performed using antibodies indicated. 5% of cell lysates were loaded on input lanes. In the bottom panel, IP and IB for detecting overexpressed EDEM1-HA interacted with TRAP complex. HEK293 cells were transfected with mock plasmids or plasmids encoding EDEM1-HA, and IP-IB experiment was performed. Asterisks indicated non-specific signals.