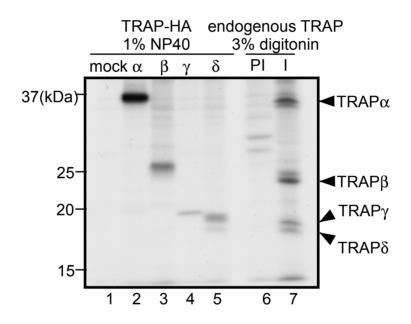
supplementary Fig 5



supplementary Fig 5. Detection of endogenous TRAP complex by immunoprecipitation.

C-terminally HA-tagged TRAP subunits were used as controls for identification of the endogenous TRAP subunits. Cells were labeled with [³⁵S]-Met/Cys for 4 h, then extracted in 1% NP40 lysis buffer and immunoprecipitated with anti-HA antibody (lanes 1-5). For detection of the endogenous TRAP complex, cells were radiolabeled with [³⁵S]-Met/Cys for 4 h and extracted with buffer containing 3% digitonin, then immunoprecipitated with preimmune rabbit serum (PI: lane 6) or anti-TRAPα antibody (I: lane 7).