

Figure S1 Cep57 is a PCM component. (A) Immunoblotting of HeLa cell lysates using anti-Cep57 antibody. (B) The immunofluorescence image shows the co-localization of Cep57 (green) and γ -tubulin (red). Boxed regions showing higher magnification images of the centrosome. Bar, 5 µm. (C) Immunostaining of HeLa cells at different mitosis stages for Cep57 (green) and α -tubulin (red). Bar, 5 µm. (D) The centrosome localization of Cep57 is microtubule-independent. HeLa cells were treated with nocodazole to depolymerize the microtubule network and then immunostained with Cep57 (green) and α -tubulin (red). Bar, 5 µm. (E) Immunoelectron microscopy images of HeLa cell centrosomes. Negative control, mouse normal IgG. Bar, 100 nm. (F) The graph shows the number of golden particles around the centriole within an area of $1.5\mu m^2$. A minimum of 15 cells was counted per sample. Error bars represent mean \pm SEM. ** *P*<0.001, unpaired Student's *t*-test.