





Figure S6. Levels of overexpression of Cenp-A/Cid and Cenp-C in *Drosophila* embryos.

(A) Total extracts of 5-8 h old w^l embryos and embryos overexpressing paternally derived UAS-Cenp-A/cid driven by maternal $\alpha 4tub$ -GAL4-VP16 were probed with anti-Cenp-A/Cid (α -Cenp-A) and anti- α -Tubulin (α -Tub), which served as loading control. The numbers indicate loading in embryo equivalents (above the lanes) or the position of molecular weight marker (kDa; left side). The asterisk marks a prominent, unspecific band.

(B and C) Paternally derived UAS-Cenp-A/cid, UAS-Cenp-C and UAS-cal1-EGFP or UAS-cal1 were expressed individually or in combinations using maternal $\alpha 4tub$ -GAL4-VP16. Total extracts were prepared and analyzed by immunoblotting as illustrated in panel A. The band intensities obtained with anti-Cenp-A/Cid (B) and anti-Cenp-C (C) were quantified (see Materials and Methods). The band intensity observed in the w^I control embryos was set to 100%. The type(s) of UAS transgene expressed is indicated below the bars.