



Figure S4. Downregulation of *tube* by *opus* insertion does not depend on an alternative transcription start site.

A. Schematic of primer locations for quantitative RT-PCR and 5' RACE to assay for use of an alternative transcription start site in embryos from *tub^{ste}/tub^{null}* females. Quantitative RT-PCR primer sets are labeled D-H. Products were detected from primer set H, but not from primer sets D-G.

B. Results from 5' RACE using RNA prepared from embryos from *tub^{ste}/tub^{null}* females. Expected size of *tube* transcript originating from wild-type transcription start site was 830 bp.