

Supplementary Material

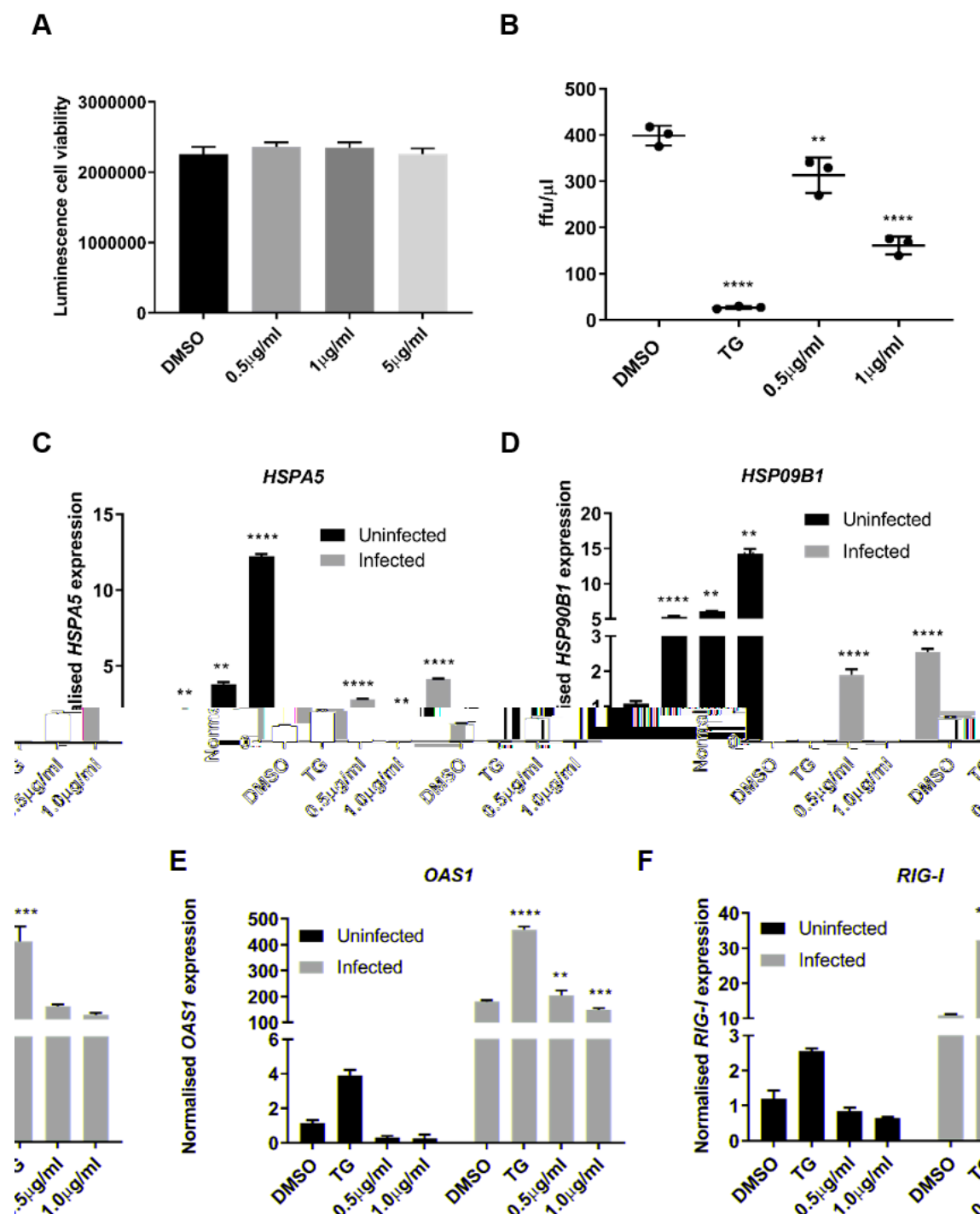


Figure S1. TG and tunicamycin mediated ER stress. (A) NPTr cells were incubated in indicated concentration of tunicamycin or DMSO control for 30 min and cell viability assay (CellTiter-Glo luminescent cell viability assay) performed 24 h later. Significance determined by one-way ANOVA, relative to DMSO control. (B–F) NPTr cells were primed for 30 min with 0.5 µM TG, tunicamycin (0.5 or 1.0 µg/mL) or DMSO, and subsequently infected with USSR H1N1 at 0.5 MOI for 24 h. (B) Spun sns were used in 6 h FFAs on MDCK cells. Significance determined by one-way ANOVA relative to the DMSO control. (C–E) Total RNA was extracted from each sample

to detect expression of ER stress markers (*HSPA5* and *HSP90B1*) and type I IFN associated (*OAS1* and *RIG-I*) genes, normalised to *18S* rRNA. Significance determined by two-way ANOVA, relative to corresponding DMSO control. * $p < 0.05$ ** $p < 0.01$ *** $p < 0.001$ **** $p < 0.0001$).