

Invasion efficiency

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ment for T3SS1-mediated entry (A) Invasion efficiency of a T3SS1 mutant ( $\Delta prgI$ ) as determined by gentamicin protection assay in indicated cell lines at 1 h p.i., expressed as a percentage of the invasion efficiency for wild type bacteria in each cell line (set to 100%). HCT116, human colonic epithelial cells; HeLa, human cervical epithelial cells; Caco-2 C2BBe1, human colonic epithelial cells; J774A.1, mouse macrophage-like cells. (B) CHQ resistance assays in various cell lines. Bacterial subcultures were grown in the presence of 100 ng/ml ATc for 1.5 h prior to infection to induce synthesis of InvC-FLAG. MOIs were adjusted so that equivalent numbers of bacteria were internalized, and the proportion of cytosolic bacteria was determined at the indicated time points by CHQ resistance assay (mean  $\pm$  SD, n $\geq$ 3 independent experiments). Asterisks denote data significantly different from WT bacteria.

FIG S1: Effect of InvC destabilization in cell lines with differential require-