

2 3 4 Figure S1. The effects of virulence gene expression and quorum sensing on biofilm dispersal. A. Detachment of WT and $\Delta toxT$ mutant strains in the presence of TC or DC, in which remaining 5 biofilm mass is quantified by CV staining. **B.** Detachment of WT and $\Delta hapR$ mutant strains in 6 the presence of TC or DC, in which remaining biofilm mass is quantified by CV staining. As 7 $\Delta hapR$ strains form thicker biofilms, data are presented as fold change over the no supplement 8 condition. Statistical significance reported as NS: no significance. 9





Figure S2. Effects of chloramphenicol on protein syntheis. V. cholerae harboring pBAD-GFP

vector was inoculated into glass tubes and allowed to form biofilms for 24 hours at 22°C.

4 Biofilms were incubated with the protein-synthesis inhibitor chloramphenicol (Cm) and 0.1%

5 arabinose for two hours. Biofilms were washed with PBS, manually disrupted, and fluorescence

measured. Fluorescence was normalized to OD₆₀₀ and reported in relative fluorescence units

6 7 (RFU). Data are means and s.d. of three independent experiments.





Figure S3. Contribution of cell lysis to carbohydrate content measured. 1 mL of mid-log culture of wild type or $\Delta vpsA V$. *cholerae* was pelleted and resuspended in 1 mL of M9 minimal media + 0.2% glycerol then diluted 10, 100, or 1000-fold into 1mL of media. Aliquots were serially diluted and plated to quantify bacterial concentration. Samples were sterilized through 0.2 μ M filters and carbohydrate content of filtrate was estimated using a modified phenol-sulfuric acid method. Data are means and s.d. of three independent experiments. Statistical significance

9 reported as NS: no significance.

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Figure S4. The effects of Ca^{2+} on biofilm dispersal. Detachment of WT V. cholerae in the presence of Ca2+ and TC as measured by biofilm mass remaining. Data are means and s.d. of 2 3

4 three independent experiments. Statistical significance reported as NS: no significance; ****: P< 0.0001.

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Figure S5. The effect of taurine and TDC on *V. cholerae* biofilm dispersal. A. *V. cholerae* was
inoculated into glass tubes and allowed to form biofilms for 24 hours at 22°C. Biofilms of

4 cultures grown in the presence of taurodeoxycholate (TDC), or taurine (TR) were quantified with

5 crystal violet staining and data are presented in OD₅₇₀ values. **B and C.** Detached cells and

6 remaining biofilm mass when mature biofilms were incubated in the presence of 1mM TDC,

7 1mM TR or 10 mM TR for two hours. Data are means and s.d. of three independent experiments.