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Vibrio cholerae motility exerts drag force to impede attack by the bacterial predator Bdellovibrio bacteriovorus

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The bacterial predator Bdellovibrio bacteriovorus is evolved to attack and kill other bacteria, including the human intestinal pathogen Vibrio cholerae. Although B. bacteriovorus exhibit a broad prey range, little is known about the genetic determinants of prey resistance and sensitivity. Here we perform a genetic screen on V. cholerae and identify five pathways contributing to predation susceptibility. We find that the essential virulence regulators ToxR/S increase susceptibility to predation, as mutants of these genes are more resistant to predation. We observe by flow cytometry that lipopolysaccharide is a critical defense, as mutants lacking O-antigen are rapidly attacked by predatory B. bacteriovorus. Using polymer solutions to alter media viscosity, we find that when B. bacteriovorus attacks motile V. cholerae, increased drag forces slow its ability to prey. These results provide insights into key prey resistance mechanisms, and may be useful in the application of B. bacteriovorus in treating infections.
B
dellovibrio bacteriovorus is a predatory bacterium capable
of preying on a wide array of Gram-negative bacteria
including human pathogens and is often described as a
living antibiotic. These small bacteria first hunt and then attach
to their prey via mechanisms thought to involve motility and type
IV pili (T4P) respectively. Within 10 to 20 min of attachment, B.
bacteriovorus invades the prey periplasm where it kills the host
and remodels host peptidoglycan to create a spherical bdello-
plast. Inside this protected niche, it degrades the host’s cytosolic
proteins and nucleic acids, using these products to fuel its own
replication before it lyses its prey and releases several daughter
cells. This lifecycle is usually completed within 2–3 h, underscoring B. bacteriovorus’ potential as an anti-infective therapy.
Owing to the alarming rise of antibiotic-resistant bacteria and
dearth of new treatments entering the clinic, B. bacteriovorus is
being evaluated for therapeutic purposes and has recently been
shown to attenuate Klebsiella pneumoniae and Shigella flexneri
in vivo.

Another susceptible prey species is Vibrio cholerae, the
causative agent of the severe diarrheal disease cholera. The global
cholera burden is substantial, with 3–5 million annual cases
leading to over 100,000 deaths each year. The disease is
demic in parts of the Americas, Africa, and South Asia, and recent
epidemics have occurred following natural disasters in Haiti.
Disrupted water infrastructure can increase disease burden as the
pathogen lives in marine and brackish waters in many temperate
parts of the world. Considering that B. bacteriovorus is ubiquitous
in water and soil, it likely preys on V. cholerae in its natural
environment.

While B. bacteriovorus’ basic lifecycle is well established, the
genetic mechanisms of predation for prey and predator remain
poorly understood. One study, focusing on the predation of
Erwinia caratova and Pseudomonas syringae, found that resis-
tance was plastic rather than genetic and could be quickly
reversed. However, a recent study demonstrated that under certain
conditions Chromobacterium piscinae produces cyanide to
protect itself from B. bacteriovorus predation. To further
explore potential genetic differences in susceptibility, we created
saturated transposon libraries in V. cholerae and Escherichia coli,
subjected them to predation by B. bacteriovorus, and used
transposon-insertion sequencing (Tn-seq) of the surviving prey to
identify susceptibility and resistance genes. We identified very few
such genes in E. coli, but many in V. cholerae, suggesting that prey
species may vary widely in their evolved strategies to resist pre-
dation by B. bacteriovorus.

Here we report five pathways important for increased or
reduced sensitivity of V. cholerae to B. bacteriovorus predation.
Two pathways sensitize V. cholerae to predation, as these mutants
are more resistant to B. bacteriovorus killing. A key resistance
mechanism is lipopolysaccharide (LPS), as mutants lacking O-
antigen are attached to and killed more rapidly by B. bacterio-
vorus. Additionally, we find that drag exerted by the motion of
V. cholerae slows the predator’s ability to kill its prey.

Results

Tn-seq for prey genes involved in predation. To identify prey
pathways playing a role in predation, we created mTn10 trans-
poson libraries in V. cholerae and E. coli (Supplementary Data 1)
and subjected them to predation by B. bacteriovorus. The V.
cholerae library contained 50,000 unique insertions while the E.
coli library contained 80,000 unique insertions, such that both
libraries approached saturation. For each of three biological
replicates, we prepared two parallel culture tubes with 10⁶ colony-
forming units (CFU)/ml prey, infecting one with B. bacteriovorus
at a multiplicity of infection (MOI) of 0.1 for V. cholerae and
MOI 0.5 for E. coli. For V. cholerae, we observed 0.6% survival
after 14 h of predation, while E. coli was more sensitive to B.
bacteriovorus and showed 0.4% survival after just 3 h. To avoid
sequestering contaminating DNA from dead prey, we outgrew all
of the prey that survived the infection (≈5 x 10⁶ CFU) in 200 ml
LB broth overnight, while in parallel outgrowing 5 x 10⁶ CFU of
the uninfected prey in 200 ml LB broth as well.

We next isolated genomic DNA (gDNA) from each sample and
processed the DNA for massively parallel sequencing of the
mTn10 junctions using Illumina sequencing. A new method
termed Nextera Tn-seq was devised for sample processing, which
uses Illumina Nextera Tagmentation to place a defined sequence
adjacent to mTn10 insertions to allow for subsequent PCR
amplification of the transposon junctions (see Methods for
details). Each PCR product was comprised of a unique 8 base pair
index adjacent to the mTn10 inverted repeat, an unknown gDNA
sequence of variable length flanking the transposon, and
Illumina-specific sequence necessary for sequencing (Supplemen-
tary Data 2). We combined the DNA samples to allow multiplexing
and sequenced the DNA using the Illumina HiSeq 2500 to determine the frequency of each transposon insertion in
the infected and uninfected populations. We calculated the fitness
contribution of each gene using the bioinformatics software as
previously described, where fitness represents the net survival
of that gene disruption mutant relative to the bulk population
(Supplementary Data 3 and 4).

Using gene ontology terms, we categorized the 151 insertions
that reduced V. cholerae fitness <0.4 (Fig. 1a), indicating that
these transposon mutants had a survival defect during B.
bacteriovorus predation. Over one third of the reduced fitness
mutants had insertions in flagellar motility genes, in part a
reflection of the many genes required to construct and regulate a
flagellum. These genes included the master transcriptional
regulator FlrA and the flagellar motor protein MotY. Other
prominent gene categories included metabolism, general regula-
tory genes, cell envelope genes, and LPS genes. The majority of
genes required for O-antigen biosynthesis and transport were
hits. Seventy-nine V. cholerae gene insertions demonstrated
improved survival during predation with fitness values >2.5,
including the virulence regulators ToxR and ToxS (Supple-
mentary Data 3).

The E. coli reduced-fitness hits were less promising as there
were only 14 gene insertions producing fitness values <0.4
(Supplementary Data 4). Furthermore, the smallest fitness value
was 0.26, whereas the V. cholerae screen demonstrated 71 hits
with fitness values <0.1 (Supplementary Data 3). The E. coli Tn-
seq screen resulted in 69 gene insertions with fitness values >2.5,
but none of these hits validated in later experiments.

Validation of V. cholerae gene hits. To validate the V. cholerae
hits from the Tn-seq screen, we created a small library of
representative marked gene deletion mutants and performed a
mini Tn-seq screen on the pooled mutants. We used the FRT/FLP
recombinase system to create 32 V. cholerae mutants, each with
an identical 81 bp FRT scar in place of the gene of interest. As
controls, this library included several mutants with initial fitness
values between our cutoffs of 0.4 and 2.5 (VC2298, AmpG, PutR,
VCA1097), as well as two neutral mutants with fitness values
close to 1.0 in the initial Tn-seq. We performed the selection for
the screen as described above, with five biological replicates for
each pair of infected and uninfected prey V. cholerae. The FRT
scar served as a pseudo-transposon for junctional sequencing
and, thus the sample preparation, sequencing, and bioinformatics
analysis were repeated as described above for the ten new samples.
Following the mini Tn-seq screen, 9 of the 26 mutants tested for validation showed significantly increased or decreased fitness values when compared to the neutral genes (Fig. 1b). Both non-motile mutants ΔflrA and ΔmotY showed reduced fitness, while of the two only the ΔmotY mutant makes a flagellum (Supplementary Figure 1). This suggests flagellum presence is irrelevant to predation and it is motility of wild-type (WT) prey that alters resistance to *B. bacteriovorus*. Other genes required for resistance to predation included OppC, a transmembrane component of the ABC transporter permease system for small peptides, and ManC, a type II phosphomannose isomerase required for O-antigen biosynthesis (Supplementary Figure 2). Genes that in WT prey increase sensitivity to predation (these mutants were more resistant) included the virulence regulator ToxR/S and the response regulator VC1087. The majority of these mutants showed similar motility to WT *V. cholerae* in soft agar chemotaxis assays (Supplementary Figure 3). For follow-up experiments, we decided to focus on the 5 prey pathways that demonstrated fitness values <0.4 or >2.5 in the mini Tn-seq.

To confirm these mutants did not harbor off target mutations, we complemented each representative strain by cloning their respective open reading frames into the plasmid pMAMB67EH and transferring these plasmids to each mutant parent strain. Additionally, we created a mutant with a tumble-biased version of the chemotaxis protein CheY (CheY**) to determine whether prey swimming speed or style affects susceptibility to predation.

We confirmed the CheY** mutant both had a severe tumble bias compared to the smooth swimming phenotype of the WT by microscopy and defective chemotaxis in the soft agar assay (Supplementary Figure 3). We also created a mutant of the outer membrane porin OmpU and a complemented strain. ToxR/S positively regulates *ompU* expression, thus we aimed to test whether reduced OmpU expression could explain predation resistance by the ΔtoxS strain. Next we competed these strains 1:1 with an isogenic ΔlacZ *V. cholerae* strain during *B. bacteriovorus* predation and plated the survivors on 5-bromo-4-chloro-3-indolyl-b-D-galactopyranoside (X-gal) plates to enumerate blue and white CFUs. Complementation restored WT sensitivity to predation (Fig. 2). In addition, the tumble-biased CheY** mutant showed an intermediate phenotype, as it was more sensitive to predation than WT *V. cholerae* but less sensitive than the non-motile ΔmotY mutant. The ΔompU strain was also more resistant to predation, similar to the ΔtoxS mutant.

B. bacteriovorus rapidly attach to O-antigen minus prey. We hypothesized differences in mutant *V. cholerae* susceptibility to *B. bacteriovorus* might be due to attachment rates. We used flow cytometry on fluorescent bacteria to gauge direct predator-prey interaction. We first generated a constitutive green fluorescent protein (GFP) construct and transformed it into the *V. cholerae* strains. For *B. bacteriovorus*, we used a previously created strain expressing a green fluorescent protein (GFP) construct and transformed it into the *V. cholerae* strains.

![Graph showing fitness values for different mutants](image-url)
harboring a tdTomato-expressing plasmid. Following 30 min of infection at an MOI of 1, we fixed the samples before diluting into phosphate-buffered saline (PBS) and analyzing by flow cytometry. We reasoned that any events deemed double positive (green and red) would indicate attachment or interaction between \textit{B. bacteriovorus} and \textit{V. cholerae}. The presence of such double-positive complexes was confirmed by sorting the two populations and using standard fluorescence microscopy (Supplementary Figure 4). To account for false-positive events, we fixed one set of samples immediately upon mixing, which was well before \textit{B. bacteriovorus} could attach to \textit{V. cholerae}.

We collected 10,000 events for each sample and applied a gating strategy to limit false positives (Fig. 3a, Supplementary Figure 5). Surprisingly, the majority of \textit{V. cholerae} mutants did not show a significant change in \textit{B. bacteriovorus} attachment compared to WT prey (Fig. 3a–c). The exception was the \textit{ΔmanC} mutant, which does not produce O-antigen and showed a two-fold higher attachment rate. This suggests that normal \textit{V. cholerae} O-antigen might interfere with predator attachment. We observed the same results at 30 min (Fig. 3b) or 60 min (Fig. 3c) postinfection.

The \textit{ΔmotY} and \textit{ΔmanC} strains are rounded more than WT prey. Since the majority of \textit{V. cholerae} mutants did not show alterations in predator attachment, we hypothesized that the predation cycle of \textit{B. bacteriovorus} is altered in the mutant prey. If true, we would predict prey rounding, the first step following \textit{B. bacteriovorus} attachment and entry, might be altered for the mutants. To test this, we infected GFP-expressing WT, \textit{ΔmanC}, and \textit{ΔmotY} \textit{V. cholerae} with \textit{B. bacteriovorus} at an MOI of 1 for 1 h. We then fixed the cells and imaged the \textit{V. cholerae} by fluorescence microscopy (Fig. 3d). Next we used a custom Matlab script to analyze each bacterium’s circularity by calculating its eccentricity. Eccentricity scores can range from zero to one, with perfect circles scored as zero. For our Matlab script, we used a cutoff of 0.4, where any bacteria below that number were considered bdelloplasts and any bacteria >0.4 were considered unrounded vibroid-shaped cells.

After analyzing a minimum of 3300 cells across three biological replicates, our script identified 6% of uninfected cells as rounded and 30% of infected WT \textit{V. cholerae} cells as rounded (Fig. 3e, Supplementary Figure 6). These ratios matched our visual observations of the images, validating our eccentricity cutoff of 0.4. Both \textit{ΔmotY} and \textit{ΔmanC} \textit{V. cholerae} were significantly more rounded than WT, with >40% of these mutant cells scoring <0.4 for eccentricity. This \textit{ΔmanC} data is consistent with the higher predator attachment shown in Fig. 3b, c. However, the greater rounding of the non-motile \textit{ΔmotY} mutant suggests that its increased susceptibility to predation manifests post-attachment and that swimming motility by WT \textit{V. cholerae} slows progression of the \textit{B. bacteriovorus} lifecycle.

**Drag forces slow \textit{B. bacteriovorus} predation of \textit{V. cholerae}**. The attachment phase of \textit{B. bacteriovorus} is reported to last about 10–20 min, before the predator is able to invade its host’s periplasm. However, we suspected this process may be extended or interrupted for highly motile prey-like \textit{V. cholerae}, whose motion should exert a strong drag force on the attached predator. Prey bdelloplast formation occurs shortly after predator invasion, so this might explain why non-motile mutant prey are rounded and killed more than motile prey (Fig. 3d, e). To calculate the difference in force on \textit{B. bacteriovorus} attached to the two types of prey, we employed a variation of Stokes’ law (Eq. 2), which is used for calculating drag at low Reynolds numbers, with a modification to account for the non-spherical shape of the bacteria:

\[
F_{\text{drag}} = 3\pi \eta v (d_a)(1/3 + (2/3)d_d/d_a)
\]

Here \(\eta\) represents the medium’s dynamic viscosity, \(v\) is velocity, and \(d_a\) and \(d_d\) represent the equivalent spherical diameters of, respectively, the object’s cross-sectional area along the direction of motion and the object’s surface area. Using this equation, we find that, when attached to \textit{V. cholerae} traveling at 80 \(\mu\text{m/s}\) (Supplementary Figure 7), \textit{B. bacteriovorus} experiences 0.32 piconewtons (pN) of drag force. Comparatively, non-motile \textit{V. cholerae} travel at approximately 1 \(\mu\text{m/s}\) due to Brownian motion, meaning any attached \textit{B. bacteriovorus} would only experience 0.004 pN of drag force. We also observed \textit{V. cholerae} dragging \textit{B. bacteriovorus}, at an average speed of 29.4 \(\mu\text{m/s}\), under a \(\times90\) objective by phase-contrast microscopy (Supplementary Movies 1–4). Using this \textit{V. cholerae} speed with an attached predator, we calculate that the \textit{B. bacteriovorus} would face 0.12 pN of drag force.

To further explore the role of drag forces on \textit{B. bacteriovorus} predation, we used ficoll and methylcellulose, polymers that increase medium viscosity to a greater extent than they slow bacterial swimming speed. We used ficoll at 10%, which increases the medium viscosity from 0.77 centipoise (cP) to 4.03 cP at 30 °C. This 5.23-fold viscosity increase was accompanied by a 2.5-fold decrease in \textit{V. cholerae} velocity, calculated by tracking individual bacteria over time by time-lapse microscopy (Supplementary Figure 7). Since the drag force is proportional to the product of velocity and viscosity, we expect that \textit{B. bacteriovorus} attached to \textit{V. cholerae} will experience roughly a factor of 2.1 greater drag in the high-viscosity condition, all else being equal. We also used the polymer methylcellulose at 1%, which increased viscosity to 3.06 cP.

We then performed 1:1 competitions between a \textit{ΔlacZ} mutant and isogenic WT or \textit{ΔmotY} \textit{V. cholerae} in plain HEPES medium or medium supplemented with ficoll or methylcellulose.
Consistent with our hypothesis, the WT strain was four- and six-fold more resistant to \textit{B. bacteriovorus} killing at these higher viscosities, respectively, while the non-motile strain Δ\textit{motY} showed the same survival rate regardless of medium viscosity (\(P = 0.999\), Fig. 4a). The increased viscosity led to a small, but not significant, decrease in predator–prey attachment for both WT and Δ\textit{motY} prey (Supplementary Figure 8). We also tested Δ\textit{cheY3}, a strain locked in the smooth-swimming state that does not pause to tumble, in the higher viscosity predation conditions (Fig. 4b). As expected, in ficolli the Δ\textit{cheY3} strain showed an even
greater competitive advantage over WT prey. We hypothesize these mutants exert more consistent drag force on attached *B. bacteriovorus* as they do not stop to tumble like WT *V. cholerae*. The increased viscosity did not alter the competitive index for the ΔlacZ *V. cholerae* used in the competition experiments (Supplementary Figure 9).

Finally, we used fluorescent *V. cholerae* and *B. bacteriovorus* to observe whether higher viscosity alters predator invasion of prey (Fig. 4c). We infected WT *V. cholerae* with *B. bacteriovorus* at MOI 0.1 for 30 min or 1 h and then fixed the cells for fluorescence microscopy. For each of the three biological replicates, we scored 200 *B. bacteriovorus*, double blind, for invasion or external attachment to *V. cholerae* in HEPES or 10% ficoll medium (Fig. 4d). While at 30 min, we only noticed a trend, at 1 h we observed significantly less predator attachments that had progressed to prey invasion. These data are consistent with the finding that non-motile prey are more rounded (Fig. 3e) than WT we observed significantly less predator attachments that had progressed to prey invasion. These data are consistent with the finding that non-motile prey are more rounded (Fig. 3e) than WT and suggests that high drag forces slow *B. bacteriovorus* prey invasion and killing.

**Discussion**

First identified by Hans Stolp in 1962, *B. bacteriovorus* has long been studied for its unique predatory lifestyle and therapeutic potential against drug-resistant pathogens. Several years after its discovery, Varon and Shilo reported cell wall mutant prey with varied *B. bacteriovorus* attachment rates. However, in the intervening years little research has focused on prey genes involved in the predation process, nor has a specific receptor for *B. bacteriovorus* been identified in any prey species. To address these questions, we used Tn-seq to identify prey genes with positive or negative roles in susceptibility to predation. Although we found no evidence for a specific receptor, we did find five prey pathways that influence susceptibility to *B. bacteriovorus* predation. Using flow cytometry, we show that *V. cholerae* mutants lacking O-antigen are attached to more easily by *B. bacteriovorus*. We also find that both O-antigen mutant and non-motile mutant *V. cholerae* are rounded into spherical bdelloplasts more than WT prey at 1 h postinfection. Additionally, we propose that motile prey exert increased drag forces on attached *B. bacteriovorus*, slowing predator entry and killing compared to non-motile prey.

In our attachment assay, only the O-antigen-defective Δ*manC* mutant showed significantly increased interaction between *V. cholerae* and *B. bacteriovorus*. It is interesting to note that, while LPS might serve as a defense against *B. bacteriovorus*, it is often a receptor for bacteriophages that prey on *V. cholerae*. These opposing selective pressures likely play out in the aquatic environment and possibly even in the human host. While none of the other *V. cholerae* mutants showed significantly altered attachment, the non-motile Δ*motY* strain did trend toward less *B. bacteriovorus* interactions than WT. This is consistent with kinetic theory whereby motile prey should collide with motile *B. bacteriovorus* more frequently.

Despite standard predator attachment, the Δ*motY* mutants are rounded into bdelloplasts more than motile WT prey. We propose that this is due to the drag force exerted by motile *V. cholerae*. 

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**Fig. 4** Increasing viscosity reduces killing and invasion of motile *V. cholerae*. **a, b** The survival percentage of motile and non-motile *V. cholerae* strains in HEPES medium alone or supplemented with ficoll or methylcellulose to increase viscosity. Infections were carried out for 14 h with an MOI of 0.1. Survival percentage and SEM is shown for three to six biological replicates. **c** Fluorescence microscopy of *B. bacteriovorus* (magenta) and *V. cholerae* (cyan), following 1 h of infection at MOI 0.1 in HEPES or Ficoll-supplemented medium. Scale bar indicates 10 μm. **d** Images from **c** were scored for externally attached or internalized *B. bacteriovorus* at 30 and 60 min. The percentage of internalized attachments and SEM is shown for three biological replicates. Error bars represent standard error of the mean. Statistics were analyzed by ANOVA with Dunnett’s multiple comparisons test.
cholerae on attached B. bacteriovorus, as increasing this drag force with ficoll or methylcellulose improved survival rates for motile V. cholerae and reduced predator invasion. Currently, it is not known what B. bacteriovorus proteins or mechanisms mediate attachment34,35 or the force this attachment can withstand. V. cholerae motility may exert sufficient drag force to shear the predators off of their prey, and though we did not observe this process in action, we did observe B. bacteriovorus attached to still-moving V. cholerae. An alternative hypothesis is that the angle of attachment is crucial for B. bacteriovorus invasion, and drag force prevents the predator from crossing the prey cell’s outer membrane. The ratio of runs to tumbles in motility appears to play a role in prey defense as well. For instance, the smooth-swimming-biased Δchey3 strain demonstrated nearly two-fold increased resistance to predation, whereas the tumble-biased ΔcheY** strain showed decreased resistance to predation. These data suggest that B. bacteriovorus may opportunistically invade prey when they slow down and there is a concomitant drop in shear force.

It is unclear why genes of the oligopeptide permease Opp were identified in our screen, but mutants of this transporter were significantly more sensitive to predation. This permease is involved in peptidoglycan recycling, so it is possible that B. bacteriovorus is better able to remodel or cross the host peptidoglycan layer upon entry into the periplasm20,36. The critical virulence regulators ToxR/S were also involved, as these mutants were more resistant to predation. It is noteworthy that environmental V. cholerae may face evolutionary pressure against ToxR/S from B. bacteriovorus, but losing these regulators would see them lose the ability to cause disease in the human host37. Similarly, mutants of the response regulator VC1087 were more resistant to predation, and this gene has been linked to V. cholerae virulence22. ToxR/S regulates over 20 genes in V. cholerae, yet all of these genes, except for ompU, had neutral fitness values in our Tn-seq screen (Supplementary Data 3). Our data suggest that OmpU may promote predation, as ΔompU and ΔtoxS strains, which produce less OmpU, were both resistant to predation. This result contrasts with previous work in E. coli, where Schelling and Conti found that outer membrane porins do not effect prey susceptibility38. OmpU is the most abundant outer membrane protein in V. cholerae39 and could have effects on early predator–prey interactions. B. bacteriovorus encodes 150 putative proteases40 and may use these enzymes to degrade outer membrane proteins and invade the prey cells. Thus it is possible that the less proteaceous membrane of the ΔompU strain may take longer to penetrate.

B. bacteriovorus has the potential to treat infections caused by many multidrug-resistant bacteria, including World Health Organization-designated priority pathogens and most of the ESKAPE pathogens1. The genetic mechanisms of prey resistance identified in this study may inform how B. bacteriovorus is applied to patients alone or used in combination therapies. For instance, we have shown that highly motile prey are more resistant to B. bacteriovorus predation. However, gut microbes are known to dampen flagella expression to avoid immune detection by the human host41, likely making them more susceptible to B. bacteriovorus. Additionally, this may bode well for B. bacteriovorus’ use in disrupting the non-motile bacteria in biofilms, as has been shown previously32.

In this study, we developed what we believe is the first flow cytometric assay to measure bacterial predator–prey interactions, as well as the software to analyze microscopic images for prey rounding by B. bacteriovorus. We hope that these two tools will be useful for research of both mutant prey and mutant predator phenotypes. Future studies may provide further insights into the mechanisms of resistance and sensitization for VC1087 and Opp permease genes.

Methods

Pooled transposon screens and growth conditions. A new V. cholerae strain was used in this study, HC1037 (GenBank accession: CP026647 and CP026648). This strain is of the O1 serogroup, Ogawa serotype, El Tor biotype, and was isolated from an adult patient at the Cholera Treatment Center in Jacmel, Haiti, on January 8, 2014. We deleted the endonuclease ideA33 in strain HC1037 to increase natural competence. Single-gene knockouts were constructed using the FRT/FLP recombinase method of deletion18. Complemented strains were created by cloning deleted genes and their native promoters into pMMB67EH21 and grown with ampicillin at 50 µg/ml to maintain the plasmid. Constitutive GFP-expressing V. cholerae was constructed via splicing-by-overlap extension (SOE) PCR and chitin-induced natural transformation. Briefly, three-piece SOE PCR was performed by amplifying an upstream region of homology with P172 and P61 (F1R), a central region of homology with P176 and P171 (F3R), and a middle product with P178 and P188 (F2R), which are designed to contain overlapping sequences with P61 and P176, respectively. The FR2 product includes the short half-life GFPSV while FR3 product contains a kanamycin-resistant marker. These SOE-PCR products were pre-warmed LB and serve as a template for PCR using P172 and Th71. The final PCR product was added to V. cholerae, grown overnight on chitin to induce natural competence, and the next day transformants were selected for on antibiotic plates. Strains used in this study are shown in Supplementary Data 1. Primer sequences for gene deletions are listed in Supplementary Data 2. V. cholerae were grown overnight at 30 °C with shaking in LB Miller broth. B. bacteriovorus 109D and a red fluorescent strain expressing tdTomato on pMQ414 were used as predators in this study. Prey E. coli WM3064, a diaminopimelic acid (DAP) auxotroph, was grown overnight at 30 °C with 0.6 mM DAP and shaking. Predator stock lysates were prepared by co-culturing prey cells with predators in 25 mM HEPES buffer supplemented with 3 mM MgCl2, 2 mM CaCl2, For the prey suspension, 1 ml of E. coli overnight, resuspended in HEPES, and 1 ml of predator lysate were added. The coculture of prey and predator was incubated at 30 °C overnight with shaking for 24 h. Any remaining prey were removed by passing the lysate through a 0.45 µm filter. Gentamicin at 10 µg/ml was used to propagate the fluorescent B. bacteriovorus strain.

Transposon library construction. Transposon libraries were created in V. cholerae and E. coli MG1655 using the temperature-sensitive mTn10 delivery vector pDL1093 (Supplementary Figure 1) as described for pDL109819. In brief, overnight cultures of V. cholerae or E. coli were grown in LB Miller broth, with shaking at 30 °C, with chloramphenicol at 10 µg/ml and kanamycin at 50 µg/ml to maintain the plasmid. The next day, 40 µl of the overnight culture was diluted into 400 ml of pre-warmed LB miller broth at 40 °C with kanamycin at 50 µg/ml. The temperature shift blocks plasmid replication and promotes transposition of mTn10 from the plasmid into the genome. The following day, stationary-phase cultures were collected and frozen as glycerol stocks for use in future Tn-seq experiments.

Tn-seq screens during B. bacteriovorus predation. For the initial Tn-seq, V. cholerae and E. coli transposon libraries, previously frozen in glycerol, were outgrown overnight in LB broth at 30 °C with shaking. The next day, these prey bacteria were washed and resuspended in 25 mM HEPES buffer at 1×109 CFU/ml. For the mini Tn-seq, each strain was grown individually overnight and pooled for the following day. For each library outgrowth, 1 ml of prey was left uninfected, while another 1 ml tube was infected with fresh B. bacteriovorus at an MOI of 0.1 for V. cholerae and MOI 0.5 for E. coli. The infections proceeded at 30 °C with shaking for 3 h (E. coli) or 14 h (V. cholerae) sufficient for prey killing. Following the co-incubation, the full 1 ml of surviving prey (≈5×10⁹ CFU) and 3 µl of the uninfected prey (≈5×10⁶ CFU) were outgrown separately overnight in 200 ml LB broth supplemented with kanamycin at 10 µg/ml at 30 °C with shaking. This outgrowth was necessary to recover enough cells for DNA isolation from both conditions. Genomic DNA was isolated using the DNeasy Blood and Tissue Kit (Qiagen). The Tn-seq DNA was prepared for sequencing by a new method, NEXTera Tn-seq, employing Nextera tagmentation (Illumina). In this method, the DNA was first tagmented using Tagment DNA enzyme 1 from the Nextera DNA Library Preparation Kit (FC-121-1030, Illumina). Next, nested PCR was carried out, first amplifying mTn10 transposon (for FRT screen for the mini Tn-seq) and one of the Nextera transposomes. The second PCR added specific indices and Illumina end sequences needed for sequencing on the Illumina platform. The final PCR products were then pooled and purified using the QiAquick PCR Purification Kit (Qiagen). Primers for amplifying and indexing the samples are listed in Supplementary Data 2.

Sequencing and fitness calculation. Pooled and indexed DNA samples were sequenced on the Illumina HiSeq 2500 using an mTn10 or FRT scar-specific sequencing primer. The sequence reads of transposon or FRT junctions were analyzed using the Tufts University Core Facility Galaxy server. Reads were mapped to V. cholerae HC1037 (accession: CP026647 and CP026648) using Bowtie. The default Bowtie parameters were used, and the Tufts University Core Facility Galaxy server custom “adapter removal” and “no ‘hopcount’” settings were used. The reads were then mapped on the read-mapped reads. Hopcount tallies the number of sequencing reads at every insertion site, which was used to determine the complexity of the input
library, as well as the severity of the genetic selection. Aggregate hop table totals the number of sequencing reads in each gene. Included in the aggregate hop table output is a standardized frequency of transposon insertions for every gene (stan-
dardized read frequency), corrected for the gene size and calculated by the equa-
tion: standardized read frequency = (number of reads in gene X/total number of reads in Illumina library)/(size of gene X/total genome size).

**Competition assays.** Following overnight growth, WT, deletion, or complemented V. cholerae strains were washed twice and resuspended at 10^8 CFU/ml in HEPES buffer. Each strain was then mixed 1:1 with an isogenic ΔΔZ strain and split into two tubes. One tube was left uninfected, while the other was infected with B. bacteriovorus at MOI 0.1. Following 14 h of infection, the bacteria were serially diluted and plated for CFU/ml on LB with 80 µg/ml X-gal agar plates. For the increased viscosity conditions, Ficoll 400 (Sigma F8016) or 14% KD methylcellulose (Sigma M7140) were added at 10% and 1% final weight/volume, respectively. The competitive index was calculated as the ratio of the infected mutant and control strains normalized to the uninfected ratio of each strain.

**Flow cytometry.** Overnight cultures of GFP-expressing V. cholerae were resus-
pended in HEPES to 10^8 CFU/ml. Fresh cultures of tdtomato-expressing B. bacteriovorus were added to the V. cholerae at an MOI of 1. At 30 or 60 min postinfection, the bacteria were fixed in 1% formaldehyde for 10 min and diluted to 10^7 CFU/ml in PBS. Three biological replicates were carried out for each sample. Attachment/colocalization was measured for the fluorescent B. bacteriovorus and V. cholerae using an S3e Cell Sorter (Bio-Rad). We used the FL1 and FL3 filters to detect GFP and tdtomato fluorescence, respectively. To confirm that our gating strategy correctly identified green-only (V. cholerae) and double-
positive (V. cholerae + B. bacteriovorus) events, we sorted the populations into separate tubes and observed the cells by fluorescence microscopy (Supplementary Figure 4).

**Fluorescence microscopy.** For the prey rounding experiment, overnight cultures of GFP-expressing V. cholerae were resuspended in HEPES to 10^8 CFU/ml and infected with WT non-fluorescent B. bacteriovorus at MOI 1. Following 1 h of infection, the bacteria were fixed in 1% formaldehyde for 10 min. The bacteria were adhered to poly-L-lysine-coated slides using a Cytospin 3 (Shandon) at 450 x g for 1 min. Slides were imaged using a Nikon Eclipse 80i microscope. Three images were taken per slide at x40 magnification, each with approximately 500 bacteria present. In all, 3300 images were taken per slide at ×40 magni-

**Viscosity measurements.** We measured the kinematic viscosity of HEPES buffer alone, HEPES buffer with 10% ficoll, and HEPES buffer with 1% methylcellulose using Cannon–Fenske viscometers (Fungilab). All measurements were conducted with viscometers submerged in a water bath at 30 °C. The kinematic viscosity was approximated as a prolate spheroid with an equatorial radius of 0.15 µm and a sphericity of 1.000 for the viscosity calculations. All measurements were conducted with the Fenske viscometers (Fungilab). All measurements were conducted with viscometers submerged in a water bath at 30 °C. The kinematic viscosity was approximated as a prolate spheroid with an equatorial radius of 0.15 µm and a sphericity of 1.000 for the viscosity calculations. The raw data that support the findings of this study are available from the cor-
responding author upon request.

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M.C.D., J.C.F., D.W.L., R.M.Q.S., D.E.K. and A.C. conceived the study. M.C.D. and R.K.G. carried out the experiments. M.C.D. and Y.N. constructed the new strains. A.A., R.M.Q.S. and D.E.K. provided critical strains and reagents. L.M.S. wrote the code and aided M.C.D. with image analysis. M.C.D. and A.C. wrote the manuscript with comments and input from the other authors.

Additional information

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