Pseudomonas aeruginosa infection of zebrafish involves both host and pathogen determinants

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Abstract

Zebrafish (Danio rerio) have a number of strengths as a host model for infection, including genetic tractability, a vertebrate immune system similar to mammals, ease and scale of laboratory handling allowing analysis with reasonable throughput, and transparency, which facilitates visualization of the infection. With these advantages in mind, we examined whether zebrafish could be used to study Pseudomonas aeruginosa pathogenesis and found that infection of zebrafish embryos with live P. aeruginosa (PA14 or PAO1) by microinjection results in embryonic death, unlike E. coli or heat-killed P. aeruginosa, which have no effect. Similar to studies in mice, P. aeruginosa mutants deficient in type three secretion (pscD) or quorum sensing (lasR and mvfR) are attenuated in zebrafish embryos infected 50 hours post-fertilization (hpf), a developmental stage where both macrophages and neutrophils are present. In contrast, embryos infected 28 hpf, when only macrophages are initially present, succumb to lethal challenge with far fewer P. aeruginosa cells than embryos infected 50 hpf, are susceptible to infection with lasR and pscD deletion mutants, but are moderately resistant to infection with an mvfR mutant. Finally, we show that we can control the outcome of infection through the use of morpholinos, which allowed us to shift immune cell numbers, or small molecules (antibiotics), which rescue embryos from lethal challenge. Thus, zebrafish are a novel host model that is well suited for studying the interactions among individual pathogenic functions of P. aeruginosa, the role of individual components of host immune defense, and small molecule modulators of infection.
Introduction

*Pseudomonas aeruginosa*, one of the most common causes of nosocomial infections in the United States, typically infects injured, burned, and immunocompromised patients and is the primary cause of mortality among cystic fibrosis patients. It is a ubiquitous, Gram-negative bacterium adapted to a variety of niches including water, soil, and in association with other eukaryotic organisms. A number of evolutionarily divergent model hosts have been used to examine *P. aeruginosa* pathogenesis including amoebae, plants, nematodes, insects, and rodents (25, 36, 37). While much has been learned about *P. aeruginosa* pathogenesis from these models, each model has different strengths and weaknesses. Invertebrate model hosts such as *Caenorhabditis elegans* offer greater genetic tractability than rodent models. Moreover, the size and life cycle of organisms like *C. elegans* enable experiments such as comprehensive genetic screens that require large numbers of animals, in contrast to rodent models where such studies are often simply unfeasible due to cost and space requirements. The drawback to modeling human infections in invertebrate hosts is the dissimilarity between vertebrate and invertebrate immune responses. Invertebrate model hosts like *Drosophila melanogaster* and *C. elegans* do not possess adaptive immunity, a true complement system, or the immune cell multi-lineage complexity that is characteristic of humans, though *D. melanogaster* does possess phagocytic cells. Thus, a model host that combines the advantages of invertebrate and rodent models would be extremely powerful in efforts to further understand *P. aeruginosa* pathogenesis.

Zebrafish (*Danio rerio*) have a number of advantages as a model host and thus have been used to study infections with a number of pathogens including *Mycobacterium*
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*marine, Salmonella typhimurium, Edwardsiella tarda, Staphylococcus aureus and Strep-tococcus iniae* (12, 29, 34, 35, 52). Zebrafish are genetically tractable, both forward and reverse classical genetic approaches are possible in this organism (50) and sophisticated techniques using morpholinos and small molecules to precisely control spatiotemporal gene regulation in zebrafish have recently been developed (14, 44). In addition, chemical genetic approaches are feasible; chemical screens for small molecules that modulate a number of phenotypes including cell cycle progression and nervous and cardiovascular system development have been successfully performed in zebrafish (33, 47). The capacity to conduct large-scale classical and chemical genetic studies in zebrafish is possible due to their fecundity and small size; embryos/larvae may be kept in 96-well format during the first 5-6 days of development and a single adult pair of fish can generate ~200 embryos from a single mating. Furthermore, zebrafish embryos are optically transparent, which facilitates the visualization of development or infection progression in real-time. Finally, zebrafish are jawed vertebrates and thus possess both innate and adaptive immunity similar to mammals.

Zebrafish immunity resembles mammalian immunity in a number of ways, including the expression of Toll-like receptors, complement proteins, pro-inflammatory cytokines, and acute phase response proteins (5, 10, 23, 26). On a cellular level, zebrafish innate immunity includes a myeloid compartment comprised of both monocyte/macrophage and granulocytic lineages (3). Primitive macrophages have been shown to be capable of engulfing invading microorganisms when inoculated 28-30 hours post fertilization (hpf) (17, 52), while primitive neutrophil differentiation lags slightly behind with functional neutrophils appearing by 32-48 hpf (22, 34). In contrast, while T
cell progenitors do begin to populate the thymus by 3 dpf (days post-fertilization), functional maturity of lymphoid cells of the adaptive immune response has not been noted prior to 4-6 weeks post fertilization (50).

*P. aeruginosa* utilizes a number of different virulence mechanisms to combat host defense including a type three secretion system (T3SS) and the production of multiple exotoxins including elastase, exotoxin A, and phospholipase C (8, 54). A number of evolutionarily divergent host species, including *Dictyostelium discoideum, Arabidopsis thaliana, C. elegans, D. melanogaster, Galleria mellonella,* and rodents have been used to identify and define the role of these virulence determinants in *P. aeruginosa* pathogenesis (15, 25, 27, 36, 37). Remarkably, these studies have demonstrated that a number of virulence determinants are required for infection across these evolutionarily divergent host species. For example, MvfR, a transcriptional regulator of quorum sensing and virulence, was originally isolated and characterized using lettuce leaf and *Arabidopsis thaliana* host models (9, 38), but is also required in *C. elegans* and mouse burn models of *P. aeruginosa* infection (38, 48). Likewise, the quorum-sensing transcriptional regulator LasR, involved in the expression of virulence factors including elastase, phospholipase C, and exotoxin A, is required for full virulence in both murine (32, 42) and *C. elegans* (49) models. In contrast, T3S plays an important role in rodent and insect models (18, 27, 46), but not in *A. thaliana and C. elegans* host models (27). Thus, a model host that recapitulates the features of the mammalian (and ideally human) response to *P. aeruginosa* infection yet possesses advantages offered by invertebrate models would be invaluable.
Given the similarities between the zebrafish and mammalian immune response and the advantages offered by zebrafish as a model organism, we investigated whether we could establish a *P. aeruginosa* infection model in zebrafish embryos. Here we demonstrate that introduction of *P. aeruginosa* into the circulation of zebrafish embryos establishes a lethal infection that requires quorum sensing and T3S for full virulence in later stage embryos. We find that ciprofloxacin and imipenem can rescue embryos from lethal challenge. Finally we examine the contribution of both host and pathogen determinants important for the progression to lethality and place these observations in the context of other *P. aeruginosa* infection models.
Materials and Methods

Infection conditions

Zebrafish embryos derived from adults of the AB line were kept at 29°C and staged 28 hpf (or 50 hpf) according to previously described developmental criteria (19). Embryos were dechorionated manually or with pronase and then anesthetized with 0.015% ethyl 3-aminobenzoate methanesulfonate prior to injection. Bacterial cells (in a volume of 1 or 2 nL) were microinjected into the yolk circulation valley, as visually ascertained under the stereomicroscope. The inoculum size was determined by injecting an equal volume of bacterial cells into PBS in duplicate before and after injections for each needle and enumerating CFU on LB agar; the inoculum size stated throughout is the mean number of cells determined from these dilutions with the standard deviation on average ~15% of the inoculum size. Injected embryos were returned to embryo medium (E3) (30), incubated at 29°C, and monitored for survival at regular intervals under a stereomicroscope. Very small numbers of bacteria (<140 cells/mL), that had likely leaked from the micropipette during injection and been transferred along with the embryo, were determined to be present in the E3 media during monitoring. The scoring of living from dead embryos was ascertained by the presence of a heartbeat and circulating blood under a stereomicroscope. For antibiotic experiments, embryos were placed directly into E3 with or without ciprofloxacin and/or imipenem following infection by microinjection. All zebrafish experiments were performed with the approval of Massachusetts General Hospital’s Institutional Animal Care and Use Committee.

Imaging infection
Microscopy was performed with either a Zeiss SteREO Discovery.V12 stereomicroscope (Fig. 2) or a Zeiss Axio Imager.Z1 microscope equipped with differential interference contrast (DIC) and fluorescence optics using a 100X Plan-Apochromat oil immersion objective (N.A= 1.4). Living embryos were anesthetized in E3 media as described above prior to image acquisition using the stereomicroscope. Living embryos imaged using the Axio ImagerZ.1 were anesthetized as described above and mounted on glass depression slides in 1% low-melting agarose. Myeloperoxidase activity was detected using fluorescent tyramide (Cy3-TSA amplification reagent, Perkin Elmer, Waltham, MA) as a substrate for endogenous peroxidase activity as previously described (34) in embryos that had been fixed in 4% paraformaldehyde overnight at 4°C prior to staining. All images were collected using Axiovision (release 4.5) software.

Bacterial strains and growth conditions

Bacterial cells for microinjection were grown by streaking glycerol stocks to LB/agar plates (+/- antibiotics), incubated overnight at 37°C, and scraped into PBS the following day. The cell concentration was estimated by measuring the OD_{600} of the resulting suspension, which was then diluted to a stock concentration of either 1250, 2500, 3000, 4000, or 5000 cells/nl. Cells were passed through a 30 gauge needle 12 times to reduce clumping before loading the cells into a micropipette. The PA14, PA14ΔlasR, PA14ΔpscD (27), and PA14/GFP strains were obtained from Frederick Ausubel (Massachusetts General Hospital). PA14/GFP carries gfp encoded on pSMC21 (11). PA14ΔlasR is a clean deletion of lasR that was complemented by transforming PA14ΔlasR with a plasmid expressing the lasR gene in pUCP19 (43). The pscD deletion
was complemented by allelic exchange using plasmid pEX18pscD1 (27). The PA14ΔmvfR and mvfR complemented strains were obtained from Laurence Rahme (Massachusetts General Hospital, (9)), and PAO1 was obtained from Stephen Lory (Harvard Medical School).

Bacterial enumeration from infected embryos
Individual embryos were euthanized with ethyl 3-aminobenzoate methanesulfonate (0.4 mg/mL), washed twice in E3, and disrupted in 0.5 mL E3 using a 1 mL dounce homogenizer (Wheaton, 0.0035" - 0.0055") for 2 minutes. Homogenizers were then rinsed with an additional 0.5 mL E3, which was also added to the homogenate. Samples were supplemented with 200 µL PBS and 130 µL 1% Triton X-100, vortexed for 1 minute prior to bath sonication (Branson 1510) for 10 minutes, and then plated on LB medium supplemented with 15 µg/mL irgasan.

Zebrafish morphant generation
The pu.1 and gata1 morpholino oligos, previously described (16, 40), were obtained, along with the Gene Tools Standard Control morpholino, from Genetools (Philomath, OR). Morpholinos were injected into the embryo at the 1-2 cell stage at the following concentrations: gata1, 0.2mM; pu.1, 0.4 mM; and Standard Control, 0.4 mM. Elimination and expansion of the myeloid lineage in Pu.1 and Gata1 morphants, respectively, was confirmed by in situ hybridization for L-plastin expression as previously described (16, 40).
SYBR-Green Real-time quantitative (q)RT-PCR analysis

Embryos were infected with PA14, the lasR deletion mutant, heat-killed PA14, or DH5α cells or with PBS (sham-infected) and RNA was isolated from pools of 10 infected and PBS sham-infected, homogenized embryos in each condition at each time point using Trizol reagent (Invitrogen). The resulting RNA was used to template first strand cDNA synthesis reactions using oligo dT and Superscript III reverse transcriptase (both from Invitrogen) according to the manufacturer’s instructions. Primers were designed using Primer 3 (41). The sequences of the gene specific TNFα (GenBank Accession No: NM_212859) and IL-1β (GenBank Accession No: AY340959) and control (EF1α; GenBank Accession No: L47669) primers for detection are as follows: TNFaF: 5’-TGCTTCACGCTCCATAAGACC-3’, TNFaR: 5’-CAAGCCACCTGAAGAAAAGG-3’, IL1bF: 5’-TGGACTTTCGACGACAAAAATG-3’, IL1bR: 5’-CGTTCACTTCAGCTCTTGGATG-3’, EF1aF: 5’-AGAAGGAAGCCGCTGAGATG-3’, EF1aR: 5’-TGTCCAGGGGCATCAATAAT-3’; one primer from each primer pair overlapped an exon-exon junction. qRT-PCR reactions were carried out in a total volume of 25 µl with cDNA corresponding to 100 ng total RNA and 50 nM gene specific or control primers. Transcript abundance from each RNA preparation was assayed in triplicate using an ABI 7300 real-time PCR machine (Applied Biosystems). Resulting Ct values from qRT-PCR assays were analyzed by the relative standard curve method and normalized to ef1α expression. The fold change in TNFα and IL-1β transcript levels relative to PBS-sham infected embryos is the mean and SEM determined from at least three RNA preparations from each experimental condition that were each assayed in triplicate.
Statistical Analysis

Both survival curve and cytokine expression data were graphed and statistically analyzed using GraphPad Prism 4 software. Statistical differences in survival curves were analyzed using the logrank test. Statistically significant differences in TNFα and IL-1β expression between PA14 and either the lasR mutant, heat-killed-PA14 or DH5α infected embryos were determined from 3-5 biologic replicates by one-way ANOVA followed by Bonferroni’s multiple comparison test.
Results

Inoculation of embryos with *P. aeruginosa* is lethal

In an effort to develop a *P. aeruginosa* infection model in a genetically tractable vertebrate model host, we investigated whether *P. aeruginosa* could lethally infect zebrafish embryos at 28 hpf, a developmental stage where primitive macrophages are present and able to engulf invading microorganisms post infection (17, 52). Initial experiments to infect zebrafish with *P. aeruginosa* by immersing dechorionated embryos in a suspension of *P. aeruginosa* strain PA14 failed. Lethality required high concentrations of PA14 (1 x 10^9 colony forming units/mL (CFU/mL)) and was independent of the viability of the bacterial cells (data not shown), suggesting that the toxicity observed was due to a heat-stable component of the bacteria rather than from an active infection. While lower concentrations of *P. aeruginosa* (10^4 CFU/ml) have been shown to colonize the intestinal tract of 3 day post fertilization zebrafish larvae under similar static immersion conditions (39), we found that this concentration had no effect on embryo viability.

We next explored introducing PA14 into the zebrafish embryo bloodstream by microinjection into the yolk circulation valley (Fig. 1A), an area where venous blood returning from the trunk and tail is not contained within a vessel but instead flows freely over the lateral sides of the yolk before returning to the heart. Microinjection at 28 hpf of at least 1700 PA14 bacterial cells resulted in the death of all infected embryos by ~48 hours post infection (hpi) (Fig. 1B) while microinjection of equal or greater numbers of heat-killed PA14 or *Escherichia coli* strain DH5α resulted in complete survival of infected embryos (Fig. 1B). PA14 killing was dose dependent with microinjection of
fewer than 1500 cells resulting in incomplete lethality (data not shown). Finally, we examined whether *P. aeruginosa* lethality was specific to the PA14 strain, given that differences in virulence among common *P. aeruginosa* laboratory strains have been observed among different model hosts (37, 48), and found that the PAO1 strain was equally virulent to embryos infected 28 hpf (data not shown).

We next examined the ability of PA14 to cause a lethal infection in zebrafish embryos at a later developmental stage when both macrophages and neutrophils are present and functional (50 hpf). We found that, similar to 28 hpf embryos, microinjection of PA14 into 50 hpf embryos also elicits a lethal phenotype. However, a higher bacterial dose (>4500 CFU) was required to achieve 100% lethality in embryos infected at 50 hpf (Fig. 1C), suggesting embryos 50 hpf are more immunocompetent than 28 hpf embryos and can mount a more robust host defense.

**Expansion of the bacterial cell population**

Zebrafish embryo transparency allowed us to monitor the progression of infection using PA14 cells expressing GFP (PA14/GFP) episomally from a strong constitutive promoter (pSMC21, (11)). In the hours immediately following inoculation, GFP fluorescence was undetectable from background autofluorescence throughout the length of the embryo under the magnification offered by a stereomicroscope (Fig. 2A). As infection progressed, the first detectable change observed was a slowing of the embryo circulation and heartbeat, followed by the appearance of GFP fluorescence. At later stages of infection, a few hours prior to death, increasing GFP fluorescence localized to either the area around the eye (Fig. 2B) or the heart and pericardial cavity, with more diffuse GFP fluorescence.
fluorescence being detected along the length of the embryo until the time of death (Fig. 2B). Embryo death was often preceded by what appeared to be necrotic cell death in the tail (Fig. 2C). Fluorescence persisted after death for several hours. No fluorescence above background autofluorescence was observed in embryos infected with heat-killed PA14/GFP (Fig. 2D).

To gain further insight into the dynamics of PA14 replication during infection of older zebrafish embryos, we examined the overall health, fluorescence pattern, and bacterial load over time in embryos infected at 50 hpf with PA14/GFP. Embryos were infected with PA14/GFP and monitored at 0, 2, 6, 10, 24, and 48 hpi for health and fluorescence. Eight embryos were sacrificed at each time point, homogenized, and plated to examine the bacterial expansion in embryos over time. The inoculum size determined from enumerating bacteria from 8 embryos sacrificed immediately following microinjection (10,600 CFU) was similar to the inoculum size determined by plating the injection volume directly from the micropipette (10,200 CFU).

Similar to embryos infected 28 hpf, the appearance of fluorescence was only noted under a stereomicroscope several hours prior to death and was preceded by a slowing of the embryo heartbeat and a decrease in circulation in the embryo trunk and tail. In order to determine the correlation between the appearance of fluorescence and bacterial load in each embryo, bacteria were enumerated from 4 embryos displaying fluorescence (‘bright’) and 4 embryos that were not fluorescent (‘dim’), at 6 hpi and all subsequent time points; fluorescence was not observed prior to 6 hpi under a stereomicroscope. (Some living, ‘dim’ embryos were still present at 48 hpi, which is consistent with observed mild variations in time to death, with a mean of 48 hours.) In
general, GFP fluorescence correlated with a higher bacterial load, with the highest
bacterial loads observed reaching $3-4 \times 10^5$ cells within 24 hpi (Fig. 3). Conversely,
embryos scored as ‘dim’ generally had a lower bacterial burden with a few exceptions.
The overlapping bacterial burden among a few ‘bright’ and ‘dim’ embryos may have
resulted from both the subjective nature of the analysis as well as the limited sensitivity
of the stereomicroscope in comparing diffuse fluorescence across an entire organism and
intense, localized, fluorescence. In examining the correlation between bacterial burden
and fluorescence, there was partial plasmid loss as the bacteria divide in the absence of
selection within the embryo, despite the use of a GFP plasmid that is known to be
retained relatively stably in the absence of selection in vitro (4). This loss ranged from ~
26% at 7 hpi to ~ 50% at 25 hpi. However, this phenomenon is unlikely to have grossly
affected the outcome of the experiment as the differences in bacterial load between
fluorescent and non-fluorescent fish are on the log scale.

Interestingly, the majority of embryos scored as ‘dim’ between 6-48 hpi displayed
a decreased bacterial burden than in the original inoculum. There was variability in the
bacterial burden among ‘dim’ embryos, with about half of them displaying an order of
magnitude fewer bacterial cells than the initial inoculum size (Fig. 3). This finding
suggests that at least some embryos have the ability to initially control infection and clear
the majority of invading bacteria. At some point however, this immune control fails,
resulting in bacterial expansion and eventually death.

Quorum sensing and T3S are only required for full virulence in 50 hpf embryos
To further characterize *P. aeruginosa* infection in zebrafish embryos in the context of other *P. aeruginosa* infection models, we analyzed whether *P. aeruginosa* mutants that are attenuated in other animal models are also attenuated in zebrafish embryos. We examined the ability of PA14 mutants containing in-frame, clean deletions of genes involved in quorum sensing (*lasR*), quorum sensing and the transcriptional regulation of pyocyanin and hydrogen cyanide production (*mvfR*), and in T3S (*pscD*) to kill embryos infected at 28 and 50 hpf. We found that the survival curves from embryos infected 28 hpf with wildtype PA14 or either the isogenic *lasR* or *pscD* mutants were not statistically different from one another (Fig. 4A), indicating that the *lasR* and *pscD* mutants are not attenuated in embryos infected 28 hpf and that these genes are not required for full virulence at this early developmental stage. However, the survival curve from embryos infected with the *mvfR* mutant was statistically different from the curve generated from embryos infected with wildtype PA14 (p=0.01). Thus, the *mvfR* mutant is moderately attenuated in embryos infected 28 hpf.

We then infected 50 hpf embryos with wild-type PA14 and the same panel of deletion mutants to determine whether *P. aeruginosa*’s full virulence arsenal would be required for infection at a later developmental stage, when embryos might be capable of mounting a more robust immune response. In contrast to infection in earlier stage embryos, the *lasR* and *pscD* mutants were attenuated in the later stage embryos (Fig. 4B and D) and the level of attenuation with the *mvfR* mutant was even more pronounced than in embryos infected 28 hpf, with 60% survival in embryos infected 50 hpf (Fig. 4C) compared to 20% survival in embryos infected 28 hpf with the *mvfR* mutant (Fig. 4A). The survival curves for embryos infected at 50 hpf with the deletion mutants differed
significantly from those infected with the wildtype PA14 strain (the logrank test between PA14 and each mutant was $p<0.0001$ in each pairwise comparison). All mutant phenotypes could be complemented by reintroduction of the respective, deleted gene (Fig. 4B-D) with the logrank test between each mutant strain and its corresponding complemented strain determined to be $p<0.05$ in each pairwise comparison. Thus, while *lasR*-mediated quorum sensing and T3S are not required for full virulence in the infection of early-stage embryos, they are required for full virulence during infection of later-stage embryos. Likewise, the *mvfR* gene is also required for full virulence in later-stage embryos but it additionally contributes to some degree to virulence during infection of early-stage embryos. While *mvfR* and *lasR* are both involved in regulating quorum-sensing controlled genes, the genes they positively regulate only partially overlap (13) and *mvfR* has been reported to be a stronger determinant of virulence than *lasR*. Survival has been recorded to be slightly greater for mice infected with *mvfR* mutants than with the *lasR* mutant (9, 38, 49). Thus, it is not entirely surprising that there is a difference in phenotype between the *lasR* and *mvfR* mutant in embryos infected 28hpf even though they are both generally involved in the transcriptional regulation of genes involved in quorum sensing.

The myeloid cell lineage affects susceptibility to lethal infection

Since the susceptibility to infection with various PA14 mutant strains was dependent on embryo age and thus correlated with the development and function of the host immune defense, we sought to characterize the embryonic immune response to PA14 by examining the contribution of embryonic myeloid cells (macrophages and neutrophils) to
defense against *P. aeruginosa* infection. Taking advantage of zebrafish embryo transparency, we were able to visually confirm that PA14/GFP bacteria were indeed engulfed by both myeloperoxidase positive-neutrophils and macrophages (Fig. 5). We further examined the contribution of myeloid cells to defense against *P. aeruginosa* infection genetically, using morpholino knockdown of transcription factors that regulate myelo- and erythropoiesis. In zebrafish embryos, primitive myeloid and erythroid cells arise from a common myeloid-erythroid progenitor (40), whose fate is determined by the transcription factors Pu.1 and Gata1, which negatively regulate erythroid or myeloid development, respectively (16, 40). Knockdown of Pu.1 shifts progenitor cells to an erythroid cell fate, thus eliminating myeloid cells from the developing embryo and increasing the number of erythroid cells present (40). Gata1 inhibition antithetically commits progenitor cells to a myeloid cell fate, thereby effectively increasing the number of myeloid cells available to combat infection and eliminating erythroid cells (16, 40).

We first confirmed the elimination and expansion of the myeloid lineage in Pu.1 and Gata1 morphants, respectively, by *in situ* hybridization for L-plastin expression (Fig. 6A). We then found that Pu.1 morphants are exquisitely susceptible to infection with both wild-type PA14 and the *pscD* mutant in 50 hpf embryos, confirming that myeloid cells in the developing embryo are required to combat *P. aeruginosa* infection (Fig. 6B). We then determined whether susceptibility to infection with PA14 or the *pscD* mutant could be altered by increasing the number of myeloid cells by Gata1 knockdown. In later stage embryos (50 hpf), Gata-1 morphants were less susceptible to infection with PA14 compared with control embryos (Fig. 6C; logrank test between control and Gata-1
morphants infected with PA14 was p=0.0002), suggesting that the additional numbers of
myeloid cells (macrophages and neutrophils) present at this stage offered greater
protection from the lethality of PA14. Interestingly, there was no difference in the
survival of either control or Gata1 morphants infected with the pscD mutant, suggesting
that there is a mode of death independent of the T3SS that cannot be rescued with greater
numbers of myeloid lineage cells.

Cytokine response to infection

Since pro-inflammatory cytokine expression is an integral part of the vertebrate immune
response to infection and a clear advantage of modeling P. aeruginosa infection in
zebrafish rather than in invertebrate hosts, we examined zebrafish pro-inflammatory
cytokine expression in response to PA14 infection. We quantified relative transcript
levels of the pro-inflammatory cytokines TNFα and IL-1β by real-time RT-PCR in
embryos that were infected either 28 or 50 hpf with either PA14, the lasR mutant, heat-
killed PA14, or E. coli DH5α cells. Embryos that had been infected at 28 hpf
demonstrated similar levels of induction of both TNFα and IL-1β at 4 hpi regardless of
whether they were infected with PA14, the lasR mutant, heat-killed PA14, or DH5α cells
(Fig. 7). By 18 hpi however, TNFα and IL-1β levels in embryos infected with heat-killed
PA14 or DH5α cells had decreased or remained at similar levels compared with levels
observed 4 hpi. In contrast, in embryos infected with live PA14, both TNFα and IL-1β
transcript levels had increased significantly over levels observed 18 hpi in embryos
infected with either heat-killed or DH5α cells (TNFα, p<0.01; IL-1β, p<0.05; overall
data set: TNFα, p=0.0026 and IL-1β, p=0.0073) (Fig. 7).
The pro-inflammatory cytokine expression pattern observed in embryos infected 50 hpf with PA14 was even more striking. While the induction of TNFα at 4 hpi in embryos infected with PA14 was not statistically different from the levels of induction observed with any other strain, the level of TNFα at 18 hpi in embryos infected with PA14 was statistically higher from levels observed from embryos infected with either heat-killed PA14 (p<0.01) or DH5α cells (p<0.05) (overall data set, p=0.0068) (Fig. 7A). Upregulation of IL1β expression was even more dramatic, with expression levels significantly greater both at 4 hpi (p<0.01; overall data set, p=0.0058) and 18 hpi (p<0.05; overall data set, p=0.0024) compared to levels in embryos infected with either DH5α or heat-killed PA14 (Fig. 7B).

In addition to the difference in cytokine response to PA14 or heat-killed PA14/DH5α challenge, there was also an interesting difference in cytokine induction between wild-type PA14 and the lasR deletion mutant. While there was no statistically significant difference in either TNFα or IL-1β expression levels at either 4 or 18 hpi in embryos infected 28 hpf with either PA14 or the lasR mutant, there were notable differences in these cytokines in embryos infected 50 hpf. TNFα expression at 18 hpi in embryos infected 50 hpf with the lasR deletion mutant was significantly lower (p<0.05; overall data set, p=0.0068) than the level observed after infection with PA14 and more closely mirrored the expression levels observed following infection with either heat-killed PA14 or DH5α cells (Fig. 7A). However, there was no statistically significant difference in IL1β expression between lasR deletion mutant or wildtype PA14-infected embryos inoculated 50 hpf (Fig. 7B). Thus, it would appear that levels of TNFα and not IL-1β
expression late in PA14 infection correlate with death, as the lasR mutant is attenuated in embryos infected 50 hpf.

Small molecules are capable of rescuing embryos from lethal infection

One advantage of using zebrafish embryos to model human disease is the ability to conduct chemical screens for small molecules that perturb a given phenotype in a whole organism model (24). With this in mind, we examined whether treatment of infected embryos with known anti-Pseudomonal antibiotics could rescue zebrafish embryos from the lethality of PA14 infection. We found that either ciprofloxacin (50 µg/ml) or imipenem (50 µg/ml) could rescue 65-75% of embryos from lethal P. aeruginosa infection when embryos were inoculated at 50 hpf (Fig. 8B). In contrast, embryos infected 28 hpf required a cocktail of both imipenem (200 µg/ml) and ciprofloxacin (150 µg/ml) to rescue similar embryo numbers (Fig. 8A). It is possible that either the increased immunocompetence of 50 hpf embryos and/or potentially an increased ability to absorb antibiotics orally later in infection (as the zebrafish larval mouth opens and intestines become motile 72 hpf) accounts for the observation that lower concentrations of single antibiotic can rescue more embryos inoculated 50 hpf than embryos inoculated 28 hpf. Notably, the concentrations of antibiotic required for protection are much higher than the minimum inhibitory concentration (MIC) of either antibiotic for PA14 in axenic culture (ciprofloxacin, MIC= 0.8 µg/ml; imipenem, MIC=1.6 µg/ml), suggesting that pharmacokinetic and pharmacodynamic issues of antibiotic distribution in the host are dictating the required concentrations for rescue.
Discussion

We report that *P. aeruginosa* can establish a lethal infection in zebrafish embryos, thus establishing a new host model for studying *P. aeruginosa* pathogenesis that combines genetic tractability and vertebrate immunity. The outcome of infection can be influenced on the pathogen side by both the inoculum size and the presence of known virulence determinants (*lasR*, *mvfR*, and *pscD*) and on the host side by developmental stage and the presence of immune cells. The outcome of infection can also be modulated by the addition of small molecules to the embryo media. Using this model, one can examine the complex host-pathogen relationship while manipulating the pathogen and/or the host using classical or chemical genetics.

Notably, we find that the host response to infection is dependent upon the developmental stage of the embryo. We find that more bacterial cells are required to achieve 100% lethality in embryos inoculated 50 hpf than in embryos inoculated 28 hpf. While 28 hpf embryos are slightly smaller (~2.5mm) in length than 50 hpf embryos (~3.1mm), the difference in body mass between 28 and 50 hpf embryos is relatively small and unlikely account for the difference in *P. aeruginosa* lethal dose between these two developmental stages. If the toxicity of *P. aeruginosa* at 28 and 50 hpf was solely related to body mass, one would expect the pattern of susceptibility to infection with the *lasR*, *pscD*, and *mvfR* mutant strains examined to be the same between 28 and 50 hpf embryos. Instead, we find that embryos inoculated 28 hpf are equally susceptible to infection with the either the wildtype PA14 strain or the *lasR* and *pscD* mutant strains unlike embryos inoculated 50 hpf. The difference in susceptibility to mutant strains suggests that *P. aeruginosa* requires its full-virulence arsenal in 50 hpf embryos in order to create a niche
where is can survive and divide. In contrast, *P. aeruginosa*’s full virulence arsenal is not required to create a niche where it can survive and divide within 28 hpf embryos, suggesting that embryos 28 hpf are less immunocompetent than embryos 50 hpf to combat *P. aeruginosa* challenge.

The development of host immunity is a complex process that requires a series of coordinated events, including functional differentiation of immune cells, expansion of immune cell populations, and the expression of other immune functions such as complement proteins. Formally, any one or a combination of these possibilities may account for the difference in immunocompetence between 28 and 50 hpf embryos. Clearly, the presence of myeloid lineage cells is an important step for embryo survival following infection, as evidenced by the rapid death of Pu.1 morphants. Moreover, the expansion of the myeloid population renders them less susceptible to infection as evidenced by the Gata1 morphants, suggesting that the decreased susceptibility of the later embryos could be due to an increase in myeloid cell numbers. However, the difference in phenotype between the two stages of embryos can also be accounted for by differences in functional maturity of the myeloid cells present as embryos infected 28 hpf initially do not have functional neutrophils. Clearly, a more detailed understanding of the timing and development of immunologic competence in zebrafish is required before one could attribute a given immunologic function to the difference in phenotype between these two different embryonic stages.

Both macrophages and neutrophils, that we observed to engulf *P. aeruginosa*, are likely to be important in combating *P. aeruginosa* infection in embryos, as they are in human infection. However, the individual contributions of each cell type in controlling *P.
P. aeruginosa infection in zebrafish embryos cannot currently be determined genetically, as there is no known gene that one could specifically target that would disrupt either macrophage or neutrophil differentiation. While recent work suggests that zebrafish primitive macrophages are able to phagocytose microbes to a far greater extent than primitive neutrophils (20), the individual contribution of macrophages and neutrophils to defense against P. aeruginosa infection awaits further study.

While embryos inoculated 50 hpf are more capable of successfully mounting a defense against infection with P. aeruginosa strains mutated in different virulence mechanisms, they still succumb to infection with wild-type PA14 even though pro-inflammatory cytokine expression both early and late in infection is robust. It is possible that the consistently high levels of TNFα observed during PA14 infection may be detrimental to embryonic survival considering TNFα’s known effects on vascular permeability in mammals. Intraperitoneal injection of high doses of LPS alone in mammals are known to result in high levels of expression of TNFα that can result in dramatic increases in vascular permeability and death (51). While fish are thought to be more resistant to LPS toxicity than rodents or calves (2), zebrafish larvae are clearly sensitive to immersion in high concentrations of LPS and display pathophysiologic features characteristic of LPS intoxication similar to mammals (1). Here, the introduction of LPS in the form of either heat-killed PA14 or DH5α cells into the zebrafish embryo bloodstream does not result in high levels of TNFα transcripts at 18 hpi and is insufficient to result in lethality. Introduction of intact PA14 bacteria on the other hand does result in high levels of TNFα late in infection and this observation is correlated with
death. Whether high levels of TNFα induction in PA14-infection contribute to embryonic lethality or are simply a marker of PA14 infection awaits further study.

Comparison to other models

The virulence determinants required for infection in zebrafish embryos 50 hpf are more similar to those required in rodent models of acute *P. aeruginosa* infection (9, 18, 32, 42, 46, 49), than to invertebrate models like *C. elegans*. The quorum sensing mutants *lasR* and *mvfR* are attenuated in zebrafish and the mouse burn model to approximately the same degree, with ~50% survival for *lasR* and *mvfR* mutants in both hosts (38, 49). Here we also find that deletion of *pscD* attenuates infection in 50 hpf embryos and thus, T3S is required for full virulence in zebrafish, similar to burn, neutropenic, and acute pneumonia murine infection models (18, 46, 53) and unlike infection in *C. elegans*, where *pscD* is fully dispensable for infection (27). Since intravenous inoculation of zebrafish embryos elicits an acute, bacteremic infection that most closely resembles the mouse burn model of *P. aeruginosa* infection, based on the levels of attenuation of the mutants examined, this model will perhaps be more useful in modeling the systemic *P. aeruginosa* infections that occur in burned and immunocompromised patients than in chronically infected patients where mutations in *lasR* and defects in T3S have been noted to appear during the course of persistent infection (21, 45).

Advantageous features of a *P. aeruginosa* infection model in zebrafish

One advantage of this model is the ease with which various pathogen components can be analyzed in the context of varying host components, such as examining different *P.*
aeruginosa mutants infected at different developmental stages or examining infection while altering immune cell numbers using morpholinos to shift myeloid and erythroid cell populations. Another advantage, unlike most model host organisms, is that zebrafish are amenable to chemical genetics as well as classical genetics (24). Here we find that the outcome of infection can also be modulated by the addition of small molecules to the embryo media, thus this model can also be used to probe P. aeruginosa pathogenesis in the intact host using chemical genetics. We show that forward chemical genetic screens are feasible in this infection model by rescuing infected embryos with small molecules added to the surrounding water. Lower concentrations of antibiotics are sufficient to rescue embryos infected at 50 hpf than at 28 hpf from death. While we do not know the mechanisms for attaining adequate tissue and bloodstream concentrations, passive diffusion of the antibiotics likely occurs and accounts for the success of other reported chemical genetic screens in zebrafish embryos (33, 47).

Chemical screening of whole-organism infection models is an attractive approach for identifying next-generation antimicrobials, particularly given the current climate where antibiotic resistance is outpacing antibiotic discovery and development. Our current antibiotic stockpile is largely composed of variations of compounds discovered ~40-60 years ago for their ability to kill or inhibit the growth of logarithmically growing bacterial cells in vitro. Since that time, with the exceptions of the narrow spectrum drugs daptomycin and linezolid, no new classes of clinically relevant antibiotics have been discovered. More recent efforts using target-based approaches to identify inhibitors of gene products thought to be essential for bacterial viability have been largely unsuccessful (31). Unlike target-based assays, however, whole-organism screening has
the potential to directly identify compounds that are effective at eliciting the desired
phenotype (like attenuation of infection) and has the potential to leap-frog over some of
the major hurdles associated with drug development in that whole organism screening
inherently selects for compounds that are permeable to the cell, have little to no gross
toxic side effects, and have acceptable pharmacokinetic profiles (at least in the model
host) (24). While examples of whole organism screening for anti-infectives in rodent
models are extremely rare due to space, cost, and even ethical considerations, they have
historically resulted in the successful discovery of drugs like ivermectin, an anti-parasitic
therapeutic (7). Recent efforts to conduct whole organism screening of invertebrate
infection models have overcome many of the drawbacks associated with screening rodent
infection models and have been successfully used to identified compounds effective at
attenuating Enterococcus fecalis and Candida albicans infection in C. elegans (6, 28).
The drawback to screening whole-organism invertebrate infection models, of course, is
the relative dissimilarity between the invertebrate and mammalian immune response to
infection. Since zebrafish are vertebrates with an immune system similar to mammals,
whole organism screening of zebrafish infection models may be more effective at
identifying compounds useful in treating human infections.

We have demonstrated that zebrafish represent an effective new model for
examining P. aeruginosa pathogenesis that has many advantages, including ease of
manipulating the immune response in the setting of optical transparency. While
visualizing and manipulating the immune response in rodent models are technically
feasible using intravital microscopy and genetic tools, such studies will be technically
easier, faster, and cheaper in zebrafish. Thus, zebrafish, as a model host, may provide a
unique forum in which to conduct comprehensive studies of pathogen mutants in the
context of variations in host immunity and explore individual contributions of
macrophage and neutrophil lineages to host defense. Given the ability to conduct both
classical and chemical genetic studies, the ability to manipulate host immunity, and the
ability to examine the infection as it progresses in the living organism, the zebrafish
infection model is a useful complement to and combines the strengths of existing models
of *P. aeruginosa* infection.

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Figure Legends

Fig. 1. *P. aeruginosa* infection in zebrafish embryos. A) Landmarks in the 28 hpf embryo; arrows indicate the direction of blood flow in the yolk circulation valley. B) Kaplan-Meier embryo survival curves following infection of 28 hpf embryos with varying doses of PA14, heat-killed PA14, or *E. coli* DH5α; CFU, colony forming unit; HPI, hours post infection. Embryos were monitored for survival at 17 hpi and at regular intervals thereafter. The data are representative of 3 replicates with 30 embryos per condition per replicate. C) Kaplan-Meier embryo survival curves following infection of 50 hpf embryos with varying doses of PA14 or heat-killed PA14. Embryos were monitored for survival at 20 hpi and at regular intervals thereafter. The data are representative of 3 replicates with 30 embryos per condition per replicate.

Fig. 2. Visualization of *P. aeruginosa* PA14/GFP infection over time. Embryos staged 28 hpf were infected with PA14/GFP (~1,800 CFU) or heat-killed bacteria and imaged over time (all scale bars, 200 µm). A) Fluorescent image of an embryo infected with PA14/GFP at 4 hpi. B) Fluorescent image of the embryo in panel A at 22.5 hpi. C) Bright-field image of the embryo from panel A immediately after death, 24 hpi. D) Fluorescent and brightfield images of an embryo infected with heat-killed PA14/GFP at 22.5 hpi.

Fig. 3. Expansion of *P. aeruginosa* in embryos over time. Bacterial enumeration from embryos infected 50 hpf with PA14/GFP (inoculum size ~10,200 CFU; each point
represents CFU recovered from an individual embryo). From 6-48 hpi, each embryo was also scored for the presence of fluorescence (fluorescent (■), non-fluorescent (□)).

**Fig. 4.** Susceptibility to quorum sensing and T3S mutants is dependent on embryo developmental stage. A) Kaplan-Meier embryo survival curves following infection of 28 hpf embryos with PA14 (~2300 CFU; N=30), PA14ΔpscD (~1800 CFU; N=30), PA14ΔmvfR (~2400 CFU; N=30), PA14ΔlasR (~1800 CFU; N=30), or heat-killed PA14 (N=11). Embryos were monitored for survival at 17 hpi and at regular intervals thereafter; the data are representative of 3 replicates. B) Kaplan-Meier embryo survival curves following infection of 50 hpf embryos with PA14 (~8,000 CFU; N=32), PA14ΔlasR (~7,800 CFU; N=30), the PA14ΔlasR complemented strain (~6100 CFU; N=20), or heat-killed PA14 (N=12). C) Kaplan-Meier embryo survival curves following infection of 50 hpf embryos with PA14 (~6,300 CFU; N=30), PA14ΔmvfR (~5,900 CFU; N=30), the PA14ΔmvfR complemented strain (~8,100 CFU; N=20), or heat-killed PA14 (N=10). D) Kaplan-Meier embryo survival curves following infection of 50 hpf embryos with PA14 (~5,000 CFU; N=30), PA14ΔpscD (~5,500 CFU; N=30), the PA14ΔpscD complemented strain (~6,300 CFU; N=30), or heat-killed PA14 (N=5). Embryos in panels B-D were monitored for survival at 20 hpi and at regular intervals thereafter; the data are representative of 3 replicates.

**Fig. 5.** *P. aeruginosa* colocalizes with myeloid cells. A-D) DIC (A), Cy3-TSA (B), GFP-fluorescent (C), and merged (D) images of a neutrophil from an embryo inoculated 50 hpf with ~4300 PA14/GFP bacterial cells that was fixed 1 hpi and stained for
myeloperoxidase activity with Cy3-TSA. E-G) DIC (E), GFP-fluorescent (F), and merged (G) images of a macrophage obtained from a living embryo inoculated 50 hpf with ~8500 CFU PA14/GFP captured 3.75 hpi. All scale bars, 5 µm.

Fig 6. Myeloid lineage cells control *P. aeruginosa* infection. A) *In situ* hybridization for L-plastin expression, a marker for myeloid cells (20) in 24 hpf wildtype (top panel), *Pu.1* (middle panel) and *Gata1* (bottom panel) morphants. B) Kaplan-Meier embryo survival curves following inoculation of 50 hpf *PU.1* (PA14, ~5500 CFU; ΔpscD, ~6000 CFU) or control morphants (PA14, ~7000 CFU; ΔpscD, ~5000 CFU). C) Kaplan-Meier embryo survival curves following inoculation of 50 hpf *GATA1* (PA14, ~6000 CFU; ΔpscD, ~5900 CFU) or control morphants (PA14, ~6700 CFU; ΔpscD, ~4500 CFU). The data are representative of 3 replicates with 30 embryos per condition per replicate.

Fig 7. Pro-inflammatory cytokine expression following *P. aeruginosa* infection. Relative TNFα (A) and IL-1β (B) transcript levels 4 and 18 hpi determined by qRT-PCR analysis in embryos that were infected either 28 hpf with ~2200 CFU or 50 hpf with ~7000 CFU of either PA14, ΔlasR, heat-killed PA14, or DH5α cells. Fold changes in cytokine expression were determined relative to PBS-sham infected embryos. Data represent the mean and SEM of at least 3 biologic replicates.

Fig. 8. Antibiotics rescue embryos from lethal infection. A) Kaplan-Meier embryo survival curves following infection of 28 hpf embryos with PA14 and subsequent immersion in embryo medium containing either ciprofloxacin (150µg/ml; ~6500 CFU),
imipenem (200µg/ml; ~6000 CFU), a combination of both antibiotics (150µg/ml ciprofloxacin; 200µg/ml imipenem; ~6600 CFU), or no antibiotic (~6700 CFU). Embryos were monitored for survival at 17 hpi and at regular intervals thereafter. The data are representative of 3 replicates with 30 embryos per condition per replicate. B) Kaplan-Meier embryo survival curves following infection of 50 hpf embryos with PA14 and subsequent immersion in embryo medium containing ciprofloxacin (50µg/ml; ~8400 CFU), imipenem (50µg/ml; ~8500 CFU), or no antibiotic (~8200 CFU). Embryos were monitored for survival at 19 hpi and at regular intervals thereafter. The data are representative of 3 replicates with 30 embryos per condition per replicate.
Fig. 1

A) Schematic diagram of a developing embryo showing the heart, pericardial cavity, yolk sac circulation valley, and yolk sac.

B) Graph showing the survival of different bacterial strains over time, with labels for PA14 (6000 CFU), PA14 (3000 CFU), PA14 (1700 CFU), heat-killed PA14, and DH5α (7800 CFU).

C) Graph showing a different set of bacterial strains with labels for PA14 (5100 CFU), PA14 (3300 CFU), PA14 (1600 CFU), heat-killed PA14, and DH5α (7800 CFU).
Fig. 3
Fig. 4
Fig. 5
Fig. 6

A)

wild-type
Pu.1 morphant
Gata1 morphant

B)

C)

Percent survival vs. HPI for different conditions: PA14/ Control, ΔpscD/ Control, ΔpscD/ PU.1 MO, ΔpscD/ GATA1 MO, heat-killed PA14/ Control.
Fig. 7

A)  

B)
Fig. 8

A) [Graph showing survival rates with different treatments.]

B) [Graph showing survival rates with different treatments.]