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Zebrafish Embryo Infection Model to Investigate Pseudomonas aeruginosa Interaction With Innate Immunity and Validate New Therapeutics

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The opportunistic human pathogen Pseudomonas aeruginosa is responsible for a variety of acute infections and is a major cause of mortality in chronically infected patients with cystic fibrosis (CF). Considering the intrinsic and acquired resistance of P. aeruginosa to currently used antibiotics, new therapeutic strategies against this pathogen are urgently needed. Whereas virulence factors of P. aeruginosa are well characterized, the interplay between P. aeruginosa and the innate immune response during infection remains unclear. Zebrafish embryo is now firmly established as a potent vertebrate model for the study of infectious human diseases, due to strong similarities of its innate immune system with that of humans and the unprecedented possibilities of non-invasive real-time imaging. This model has been successfully developed to investigate the contribution of bacterial and host factors involved in P. aeruginosa pathogenesis, as well as rapidly assess the efficacy of anti-Pseudomonas molecules. Importantly, zebrafish embryo appears as the state-of-the-art model to address in vivo the contribution of innate immunity in the outcome of P. aeruginosa infection. Of interest, is the finding that the zebrafish encodes a CFTR channel closely related to human CFTR, which allowed to develop a model to address P. aeruginosa pathogenesis, innate immune response, and treatment evaluation in a CF context.

Keywords: Pseudomonas aeruginosa, zebrafish, innate immunity, host–pathogen interactions, drug screening

INTRODUCTION

Pseudomonas aeruginosa is a mesophilic Gram-negative bacterium able to thrive in very diverse habitats, which is linked to its metabolic versatility, impressive number of regulators, and two-component systems (Arai, 2011). P. aeruginosa can colonize a broad range of hosts, from plants to animals including humans (de Bentzmann and Plesiat, 2011), where it is considered as an opportunistic pathogen as it mainly affects compromised patients. This pathogen is a leading cause of nosocomial contaminations capable of causing a myriad of infection types like bacteremia, pneumonia, keratitis, wound, and urinary tract infection, in immunodeficient subjects (e.g., neutropenic, burn, or oncological patients). In addition, most individuals with cystic fibrosis (CF), a genetic disorder caused by mutations of the cystic fibrosis transmembrane conductance
Infection in zebrafish embryos as CF model is discussed and future promising studies using this model are underlined.

**INFECTION ROUTES FOR PSEUDOMONAS AERUGINOSA IN ZEBRAFISH EMBRYOS**

Infection in zebrafish embryos is most often established through microinjection of bacteria into the embryo, either in the circulation or in closed compartments (Figure 1). The site of microinjection determines whether the infection will rapidly become systemic or will initially remain localized. In addition,
a bath immersion method has been developed with *P. aeruginosa* (Figure 1).

### Microinjection in the Circulation or in Closed Compartments

In initial experiments, *P. aeruginosa* infections were established by microinjecting bacteria into the bloodstream (injection in caudal vein or duct of Cuvier) of 1 or 2 days post-fertilization (dpf) old embryos, which causes a rapid systemic infection. The detailed handling for injection in the caudal vein has been described (Llamas and van der Sar, 2014).

Microinjection in a closed compartment such as the hindbrain ventricle (HBV), otic vesicle, or tail muscle provides a local infection that facilitates live imaging. Notably, injection in HBV or muscle allowed to visualize in real time macrophages that migrate at the injection site to phagocytose bacteria (Moussouni et al., 2021) (Figure 2). Neutrophil recruitment has also been monitored upon HBV injection (McCarthy et al., 2017). Interestingly, lines of evidence of the development of *P. aeruginosa* microcolonies were reported upon HBV injection 24 hpi, which have been proposed as precursors of biofilm (Rocker et al., 2015). Accordingly, microcolonies were less frequent and of lower size and volume for a *psl* mutant, which is defective in the production of Psl exopolysaccharide and thus biofilm formation (Rocker et al., 2015). This study was, however, conducted on fixed and not live embryos. To further evaluate biofilm initiation *in vivo*, it would be of great interest to follow the formation of microcolonies in real time on live embryos over a period longer than 24 h.

The swim bladder shares similarities with the lung, with an air–epithelial interface that produces epithelial mucus, and localized microinjection in the swim bladder has been used to model mucosal mixed infection with *Candida albicans* and *P. aeruginosa* (Bergeron et al., 2017). Infection within the swim bladder is usually performed at 4 dpf, when the organ starts to inflate and is visible for microinjection.

### Bath Immersion

Microinjection of bacterial cells into zebrafish embryos requires specific expertise and is time consuming. Bath immersion, which is easier to handle, is a natural way of infection for aquatic bacterial pathogens (Rowe et al., 2014), but not for other pathogens. A lethal infection after bath immersion of 3 dpf healthy embryos with concentrated *P. aeruginosa* culture was reported in one report (Diaz-Pascual et al., 2017), but was not reproduced in other studies (Clatworthy et al., 2009; van Soest et al., 2011; Nogaret et al., 2021). Proteomic analysis indicated induction of hypoxia response, but no inflammatory response, in zebrafish embryos exposed to the immersion method, suggesting that healthy larvae suffered from a lack of oxygen when exposed to *P. aeruginosa* by static immersion (Diaz-Pascual et al., 2017). Moreover, the host response was also likely corresponding to a response against bacteria that are outside the fish or in contact with its skin (Diaz-Pascual et al., 2017).

An alternative infection route has been recently proposed with the bath immersion of tail-injured embryos (Poplimont et al., 2020; Nogaret et al., 2021). A rapid (within 24 h) bacterial dose-dependent mortality was observed when 2 dpf embryos were injured at the tail fin before static immersion with *P. aeruginosa* PAO1 strain. Moreover, a mutant strain known to be attenuated upon microinjection in the caudal vein is similarly attenuated upon bath infection of injured embryos (Nogaret et al., 2021), supporting the reliability of the bath model in assessing *P. aeruginosa* virulence. This infection mode, which could be considered as a model of wound infection, is of interest for drug screening (see below).
INTERPLAY OF PSEUDOMONAS AERUGINOSA WITH THE ZEBRAFISH INNATE IMMUNE SYSTEM

Infection of zebrafish embryos by *P. aeruginosa* induces an acute infection and lethality of embryos, mostly within 24–30 hpi, suggesting that infection is cleared or controlled in surviving embryos. The embryo response to *P. aeruginosa* infection is strongly related to the initial concentration of infection bacteria. The zebrafish is relatively resistant to *Pseudomonas*, and injection of large inocula (above 1,000 bacteria per embryo) is required to induce host killing. Macrophages and neutrophils can rapidly phagocytose *P. aeruginosa*, suggesting that both phagocytic cell types play a role in protection against infection (Brannon et al., 2009; Clatworthy et al., 2009; Cafora et al., 2019).

A *P. aeruginosa* clinical isolate from a CF patient microinjected into the duct of Cuvier was also predominantly found to be associated or engulfed by macrophages within 6 hpi (Kumar et al., 2018). **Figure 2** recapitulates some advantages of the zebrafish embryo model to address the role of phagocytic cells during *P. aeruginosa* infection.

Phagocytosis of *P. aeruginosa*, injected in the muscle or HBV, by recruited macrophages has been visualized in real time (Cafora et al., 2019; Moussouni et al., 2021) (**Figure 2**). Depletion of macrophages, through the use of *pu.1* morphants or lipochlodronate, increased the susceptibility of larvae to *P. aeruginosa* (Brannon et al., 2009; Belon et al., 2015; Moussouni et al., 2021) (**Figure 2**), supporting the role of macrophages in the clearance of *P. aeruginosa* during acute infection. Survival of infected embryos is reduced, in association with increased bacterial burden, in embryos defective for the mitochondrial superoxide dismutase 2 (*sod2* morphant), which contributes to generating oxidative stress in phagocytes (Peterman et al., 2015).

In addition, the survival of infected embryos is largely reduced in the presence of baflomycin, which inhibits host vacuolar ATPase, supporting the idea that acid stress within phagocytic cells is important for host defense (Moussouni et al., 2021).

The recruitment of neutrophils at the site of infection has also been observed (Diaz-Pascual et al., 2017; McCarthy et al., 2017). The effect of neutrophil depletion in the outcome of *P. aeruginosa* in zebrafish embryos has not been investigated, but defects in neutrophil migration greatly sensitize embryos to *P. aeruginosa* infections (Rosowski et al., 2016; Houseright et al., 2020). More precisely, embryos defective for Rac2 function, which have neutrophils that are unable to migrate, are highly susceptible to *P. aeruginosa* (Rosowski et al., 2016). Upon localized otic infection with *P. aeruginosa*, systemic activation and mobilization of neutrophils from hematopoietic tissues is also mediated by Cxcr2 signaling, a receptor enabling neutrophils to sense the IL8 chemokine (Deng et al., 2013). In addition, the increased susceptibility to *P. aeruginosa* infection observed in zebrafish deficient for *ctxa1*, which is homologous to the C3 component of human complement, is likely due to a neutrophil-intrinsic function of C3, possibly its ability to recruit neutrophils at the infection site (Houseright et al., 2020).

In parallel to the recruitment of phagocytic cells, *P. aeruginosa* triggers a potent proinflammatory response in zebrafish, with an increased expression of cytokines TNF-α.
and IL-β that was monitored by RT-qPCR (Clatworthy et al., 2009; Cafora et al., 2019). However, even though a correlation has been made between the intensity of the TNF-α response and the virulence of strains (Clatworthy et al., 2009), it is still unclear whether the mortality of embryos is due to a cytokine storm. An inflammatory response is also supported by global expression profile (see below).

Interestingly, the diguanylate cyclase SadC and the methyltransferase WarA, which interact with the LPS biosynthesis machinery of P. aeruginosa to modify the distribution of LPS O antigen, have been involved in neutrophil recruitment (McCarthy et al., 2017). SadC and warA mutants are slightly but significantly attenuated during zebrafish infection at early time post-infection (12 hpi), without being associated with a reduced bacterial load (McCarthy et al., 2017). During infection with sadC or warA mutant in HBV, more neutrophils were recruited to the site of infection, and a higher expression of the LPS-associated proinflammatory cytokine TNF-α was measured as compared to larvae infected with the wild-type strain, suggesting that SadC/WarA modifications of LPS mediate immune evasion in vivo.

Taken together, these studies show the strong contribution of zebrafish’s innate immune system in the outcome of P. aeruginosa infection and highlight the powerful tools available to decipher the interplay between pathogen and host components. These findings corroborate the role of innate immunity in humans with the increased sensibility of immunocompromised patients, such as neutropenic patients, to P. aeruginosa infection (Sadikot et al., 2005).

**PSEUDOMONAS AERUGINOSA FACTORS INVOLVED IN PATHOGENESIS IN ZEBRAFISH**

The zebrafish infection model has been used to assess the virulence of various P. aeruginosa mutant strains, thus allowing us to evaluate the role of different bacterial factors in this model. The mutants identified as attenuated in this model are summarized in Table 1. Of specific interest, several mutants are as virulent as a wild-type strain in macrophage-depleted embryos, indicating an interplay between these bacterial factors and phagocytes.

### T3SS, QS, and Other Classical Virulence Factors

A PA14 mutant lacking T3SS-structural protein PscD is attenuated upon microinjection into the yolk circulation valley at 2 dpf (Clatworthy et al., 2009). Accordingly, a PAK mutant strain lacking ExsA, which positively regulates T3SS, is also attenuated upon microinjection in the caudal vein of 2 dpf embryos (Brannon et al., 2009). The first effect of T3SS was reported between 4 and 8 hpi, when the PAK strain, but not an exsA mutant, began to proliferate despite an initial phase of bacterial clearance, probably by phagocytes (Brannon et al., 2009).

The virulence of P. aeruginosa strains lacking a functional T3SS can be restored upon phagocyte depletion, suggesting that T3SS influences virulence through its effects on phagocytes (Brannon et al., 2009). The dramatic dependence of P. aeruginosa on the T3SS to overcome normal phagocyte defenses in the absence of adaptive immunity suggests that the zebrafish may be a useful and relevant model to understand the details of T3SS–phagocyte interactions. The specific involvement of T3SS effectors during infection in the zebrafish model still remains to be investigated.

Two bacterial factors, the inner membrane protein MgtC and the outer membrane porin OprF, which are involved in the intramacrophage survival of P. aeruginosa, have been involved in the regulation of T3SS genes (Garai et al., 2019). Similar to T3SS mutants, mgtC and oprF mutants (in the PA01 background) have been shown to be attenuated after microinjection in the caudal vein of 2 dpf embryos in a macrophage-dependent manner (Belon et al., 2015; Moussouni et al., 2021) (**Figure 2**). Attenuation of oprF mutant has also been reported upon infection by immersion of tail-injured embryos (Nogaret et al., 2021). Interestingly, the infection defect of oprF mutant in zebrafish embryo can be suppressed upon the addition of bafomycin, an inhibitor of phagosomal acidification, which correlates with a preferential association of oprF mutant with acidified compartments in cultured macrophages (Moussouni et al., 2021).

LasR and MvfR are transcriptional regulators of QS that control many genes encoding virulence factors. P. aeruginosa lasR and mvfR mutants (PA14 background) are attenuated in zebrafish embryos infected at 2 dpf into the yolk circulation valley (Clatworthy et al., 2009).

### Table 1 | Pseudomonas aeruginosa mutants attenuated in the zebrafish embryo infection model.

| Gene name | Site of injection | References |
|-----------|------------------|------------|
| exsA<sup>a</sup>, pscD (T3SS) | CV, duct of Cuvier | Brannon et al. (2009); Clatworthy et al. (2009) |
| oprF<sup>a</sup> | CV, HBV, immersion | Moussouni et al. (2021); Nogaret et al. (2021) |
| mgtC<sup>a</sup> | CV | Belon et al. (2015) |
| mvfR, lasR (QS) | Duct of Cuvier | Clatworthy et al. (2009) |
| phz1/2 | Duct of Cuvier | Chand et al. (2011) |
| sarAwarA | HBV | McCarthy et al. (2017) |
| retS, gapC, phoR, copS, bgpS, kindB (TCS) | Duct of Cuvier | Chand et al. (2011) |
| PA2206 (LysR regulator) | CV | Reen et al. (2013) |
| vnrR | CV | Llamas et al. (2009) |

<sup>a</sup>Attenuation in a macrophage-dependent manner.
Among other putative virulence genes tested, a phz1/2 mutant defective in phenazine-1-carboxylic acid biosynthesis (whose main derivative is pyocyanin) was attenuated for virulence in the zebrafish embryo infection model, suggesting that pyocyanin is a critical virulence factor in this model (Chand et al., 2011), which may interfere with bacterial clearance by phagocytes. In contrast, mutants deficient for elastase production (lasB), flagellar motility (flgK), alginate (algD), or Pel exopolysaccharide (pelA) production were not attenuated during acute infection in zebrafish embryos (Chand et al., 2011).

**Two-Component Systems and Transcriptional Regulators**

The role of 60 two-component sensors (PA14 genetic background) has been systematically tested with individual mutants in 50 hpf zebrafish embryos microinjected into the bloodstream through the yolk circulation valley. Six sensors that are required for *P. aeruginosa* virulence were identified (GasS, RetS, PhoR, CogS, BqsS, KinB) (Chand et al., 2011). A deeper analysis of kinB mutant showed that KinB is required for acute infection in zebrafish embryos and regulates a number of virulence-associated phenotypes, including QS, pyocyanin production, biofilm formation, and motility (Chand et al., 2011).

Extracytoplasmic function sigma factors are important signal-responsive regulatory proteins in *P. aeruginosa*, and a member of this family, σ^Vrel^, promotes the transcription of secretion systems and secreted proteins. The VreR anti-sigma factor, involved in the regulation of σ^Vrel^, is required for *P. aeruginosa* virulence in zebrafish embryos, possibly through the modulation of bacterial toxicity toward host cells (Llamas et al., 2009; Otero-Asman et al., 2020).

The non-classical LysR-type transcriptional regulator PA2206 is required for an effective oxidative stress response in *P. aeruginosa* (required for tolerance to H_2O_2 in vitro) and was found to be important for the lethality of zebrafish embryos injected at 26 hpf in the blood island (PAO1 derivative) (Reen et al., 2013). Whether the attenuated phenotype of PA2206 mutant in the zebrafish infection model is directly linked to its reduced tolerance to oxidative stress remains to be determined. In addition, the implication of phagocytic cells in the attenuated phenotype has not been investigated.

Cumulatively, the use of diverse mutated strains indicates that the relevance of *P. aeruginosa* factors, such as T3SS and QS, during pathogenesis in zebrafish embryo model is consistent with the results reported in acute models of mice infection (Rumbaugh et al., 1999; Vance et al., 2005; Kumar et al., 2009). In addition, virulence phenotypes in zebrafish corroborate clinical observations in humans, since T3SS has been shown to increase the infection risk and is associated with poor clinical outcomes (Hauser et al., 2002; Ledizet et al., 2012), similar to pyocyanin production upon bloodstream infections (Gupte et al., 2021). On the other hand, similar to what is observed in zebrafish, swimming motility and protease secretion have not been associated with increased pathogenicity and disease severity in humans (Ledizet et al., 2012; Gupta et al., 2021).

**Expression of Host and *Pseudomonas aeruginosa* Genes During Infection**

The transparency of the zebrafish embryos can be used to monitor bacterial gene expression in vivo using strains with transcriptional fusions with fluorescent reporter genes. Expression in vivo of a *P. aeruginosa* gene regulated by the σ^Vrel^ transcriptional factor was observed upon bacterial injection in HBV (Otero-Asman et al., 2020). How activation of σ^Vrel^ in response to the host occurs remains to be investigated.

The use of a zebrafish embryo model also allows us to assess the global gene expression of both host and microbe in parallel. This provides a unique opportunity to investigate the molecular mechanisms underlying the interaction between the host’s innate immune system and the pathogen.

A global proteomic approach was used to track simultaneously in vivo the pathogen response and host immune response at 22 hpi using 3 dpf zebrafish larvae infected with *P. aeruginosa* PAO1 by immersion (without tail injury) or microinjection (Diaz-Pascual et al., 2017). Some zebrafish metabolic pathways, such as hypoxia response, as well as the integrin signaling pathway and angiogenesis, were exclusively enriched in the larvae exposed by static immersion. In contrast, inflammation mediated by chemokine and cytokine signaling pathways was exclusively enriched in the larvae exposed by injection. Important virulence factors from *P. aeruginosa*, involved in toxin production, T3SS, QS, and the production of extracellular polymeric substances, were enriched only after exposure by injection, which is consistent with the role of these factors during acute infection in this model (Table 1).

A dual host–pathogen transcriptomic analysis was also conducted at a later infection time (3 dpi) on embryos surviving infection using a CF clinical isolate (PASS1) microinjected into the duct of Cuvier at 2 dpf (Kumar et al., 2018). PASS1 displayed increased expression of an array of genes shown previously to be important in pathogenesis, including genes encoding pyoverdine biosynthesis, flagellin, non-hemolytic phospholipase C, proteases, superoxide dismutase, and fimbrial subunits. In addition, phosphate and iron acquisition genes are significantly upregulated in PASS1, suggesting that phosphate and iron are limiting nutrients within the zebrafish host. Regarding the host, proinflammatory genes as chemokine receptors, IL-1β, Toll-like receptors, and TNF receptor signaling family were activated. Transcriptional regulators of neutrophil and macrophage phagocytosis were also upregulated, highlighting phagocytosis as a key response mechanism to *P. aeruginosa* infection.

Real-time imaging using zebrafish transgenic lines carrying reporters of inflammatory genes and a cohort of clinical isolates could increase the relevance of these observations, notably by shedding light on the ability of strains from different origins to escape immune detection. In humans, host responses to chronic *P. aeruginosa* infections are indeed complex, ranging from vigorous inflammation ineffective at eradicating infecting bacteria, to relative host tolerance through a dampened activation of host immunity (Faure et al., 2018).
ROLE OF HOST FACTORS DURING PSEUDOMONAS AERUGINOSA INFECTION: THE CASE OF CFTR

As indicated above, *P. aeruginosa* infections are a major cause of mortality and morbidity in patients with CF. The CFTR channel has a broad cellular distribution and CF affects multiple organs in humans including the lung, gastrointestinal tract, liver, male reproductive tract, and pancreas.

**CFTR in Zebrafish**

The amino acid sequence of zebrafish CFTR (zCFTR) is 55% identical to human CFTR (hCFTR), and 42 out of 46 sites of mutations found in patients with CF are conserved in zCFTR. The structure of zCFTR was the first CFTR structure to be solved (Zhang and Chen, 2016; Zhang et al., 2017) and is essentially identical to the structure of hCFTR (Liu et al., 2017). Zebrafish *cftr* is expressed in the liver, kidney, spleen, and intestine, and *cftr* transcripts are detected in cells of the myeloid lineage that includes macrophages and neutrophils (Phennicie et al., 2010).

The role of CFTR in zebrafish was addressed by generating a fish line with frameshift mutation in the *cftr* gene (Navis et al., 2013; Liao et al., 2018). In zebrafish, left–right asymmetry requires cilia-driven fluid flow within the lumen of Kupffer’s vesicle (KV), and a reduced fluid secretion in *cftr* mutant impairs KV lumen expansion leading to defects in organ laterality (Navis et al., 2013). Due to male infertility (Liao et al., 2018), the lines have to be maintained at heterozygous stage, and homozygous embryos are screened based on the altered KV morphogenesis (Navis et al., 2013). Interestingly, deregulation of *cftr* function in zebrafish causes a phenotype that mirrors other defects present in the human disease such as severe pancreatic dysfunction (Navis and Bagnat, 2015) and hematopoietic defects (Sun et al., 2018), which may correlate with anemia presented by patients with CF.

Zebrafish embryos with *cftr*-loss-of-function (also called “CF embryos”) thus represent a promising model to study the implication of CFTR in innate immune response and mucin secretion. Another way to generate a CF zebrafish model, which has been used in the context of *P. aeruginosa* infection, is through the injection of morpholino that transiently knocks down *cftr* gene expression.

**Pseudomonas aeruginosa Infection in CF Zebrafish Embryos**

CF zebrafish embryos (*cftr* morphants) have a 3.5-fold higher number of *P. aeruginosa* PA14 bacterial cells after 8 h of infection than control embryos (Phennicie et al., 2010). A similar pattern of differences in bacterial burden at early time post-infection was observed with a *P. aeruginosa* CF clinical isolate. However, no effect was reported at later times, and this difference early in the course of infection did not result in higher mortality in the *cftr* morphants compared with that of the control. On the other hand, in another study carried out with PAO1 strain microinjected in the duct of Cuvier at 48 hpf, the mortality of the CF embryos was slightly, but significantly, increased at 20 hpi comparative to control embryos (Cafora et al., 2019). Time laps analyses after HBV injection by confocal microscopy showed several microcolonies at 18 hpi (Rocker et al., 2015), and interestingly, the area of microcolonies formed in CF embryos was reported to be higher than in wild-type fish (Cafora et al., 2019). A deeper analysis of the formation of these microcolonies over a long period would be of interest to determine if they persist, increase in size and harbor biofilm characteristics, or are eliminated.

The production of reactive oxygen species is significantly dampened in *cftr* morphants compared with control embryos, and a reduction of neutrophil migration toward the injection site is observed in the case of local injection (HBV), supporting a link between CFTR and innate immune response (Phennicie et al., 2010). Moreover, CF embryos present a reduced proinflammatory immune response following bacterial infection in comparison with the wild-type, as shown by significantly reduced TNF-α and IL-1β responses (Phennicie et al., 2010; Cafora et al., 2019).

Taken together, these results indicate that zCFTR contributes, to a moderate extent, to the resistance against *P. aeruginosa* infection. This is likely linked to an alteration of the inflammatory response and could also be related to an alteration of the bactericidal action of innate immune cells (Bernut et al., 2019). Such findings corroborate a reduced clearance of *P. aeruginosa* in the CF mouse model and in patients with CF, which is in part linked to the altered ability of CF macrophages to properly control the inflammatory response and kill bacteria (Hartl et al., 2012). The results in zebrafish rely on a morphant-based CF model and it would be of interest to carry out *P. aeruginosa* infection with a mutated fish line, to strengthen the invalidation of the *cftr* gene.

**VALIDATION OF ANTI-PSEUDOMONAS STRATEGIES IN ZEBRAFISH**

Aside from being of interest as an infection model, the zebrafish embryo is also suitable for *in vivo* chemical screening (Zon and Peterson, 2005; Rennekamp and Peterson, 2015), with the advantage that permeability of the larvae allows the entry of small compounds added directly to the fish water. This model, which allows also to address drug toxicity (Eimon and Rubinstein, 2009), has been successfully used for drug testing in the context of infectious diseases (Bernut et al., 2014b). The mode of infection with bath immersion of cut-tailed embryos in 96-well plates is of particular interest for the screening of anti-infectious compounds.

**Test of Clinically Used Antibiotics**

Treatment of infected embryos, microinjected at 50 hpf with PA14 in the circulation, with ciprofloxacin (50 µg/ml) or imipenem (50 µg/ml) added in the bath medium, could rescue embryos from lethality (Claworthy et al., 2009). Protection with ciprofloxacin (50 µg/ml) was also observed with the infection mode of tail-injured embryos (Nogaret et al., 2021). In this case, an antibiotic was added in the bath 2 h after injury and bacterial
immersion, at a time when the wound was closed. Interestingly, a significant protective effect, of lower amplitude, was also shown at a much lower dose of 1 μg/ml, supporting the pertinence of the model for drug testing.

Validation of Novel Antibacterial Strategies
Phage therapy using a phage cocktail, found to be efficient to treat P. aeruginosa acute infections in mouse and Galleria mellonella larvae, was used in zebrafish embryos (Cafora et al., 2019; Cafora et al., 2020b). Phage therapy, done by injecting a phage cocktail in the yolk sac of PAO1-infected embryos, was shown to reduce lethality, bacterial burden, and the proinflammatory response caused by PAO1 infection at 20 hpi both in wild-type and CF zebrafish (Cafora et al., 2019). In addition, phages alone mitigate inflammation in wild-type and CF zebrafish by reducing the expression levels of proinflammatory cytokines and the neutrophilic recruitment to the infection site (Cafora et al., 2020a).

Anti-virulence strategies have emerged as attractive novel therapeutic approaches that would apply less selective pressure to develop resistance and better preserve microbiota than traditional antimicrobial therapy (Muhlen and Dersch, 2016). In this context, the zebrafish embryo is suitable for testing the efficacy of specific inhibitors of virulence factors that are important for P. aeruginosa infection in this model (Table 1). This is the case of QS, which has been proposed as an attractive target to fight P. aeruginosa infections using alternative therapies (Smith and Iglewski, 2003). The bath immersion model of injured embryos was used to show the anti-infective potential of a novel anti-QS compound, without exhibiting any toxicity (Nogaret et al., 2021). This compound is an antagonistic analog of C4-HSL (Furiga et al., 2016) that was not previously tested in an animal model. In vitro, C11 was shown to inhibit P. aeruginosa biofilm formation and reduce the expression of QS regulatory/regulated genes (Furiga et al., 2016). Further studies are required to determine how C11 addition in vivo reduces embryo mortality, which can be due to its ability to downregulate QS gene expression and impair virulence and/or to the fact that initiation of infection at the injured tail shares properties with the initiation of biofilm formation.

Taken together, these studies support the relevance of zebrafish as a first-intention vertebrate model to validate molecules that reduce P. aeruginosa pathogenicity and to perform drug screening. In addition, therapeutic molecules can also be tested in a CF context.

CONCLUSION AND PERSPECTIVES
In the present review, we have recapitulated the multiple points of interest of using zebrafish embryos as an infection model for P. aeruginosa (Figure 3). This model appears as a potent first-intention vertebrate model to monitor P. aeruginosa pathogenesis and test novel therapeutic strategies. Importantly, this model offers unprecedented opportunities to investigate the role of phagocytic cells and inflammatory response during infection. Whereas the role of alveolar macrophages in internalization and early clearance of P. aeruginosa in mouse lung remains controversial (Kooguchi et al., 1998; Cheung et al., 2000), macrophages play a key role during P. aeruginosa infection in the zebrafish model, thus allowing to investigate how intraphagocytic stage contributes to dissemination, persistence, and susceptibility to drug treatment.

In addition to these multiple points of interest, zebrafish embryos also offer perspectives to study P. aeruginosa clinical isolates and polymicrobial infections (Figure 3). While reference laboratory strains causing acute infections (PAO1, PA14, and PAK) have been widely used in the zebrafish model to gain insights on the host–P. aeruginosa interaction in vivo, investigation of the pathogenesis of clinical isolates in this vertebrate model remains very scarce, with only two CF isolates used (Phennicie et al., 2010; Kumar et al., 2018). In the future, a broader and deeper analysis of the behavior of clinical acute or chronic strains would shed light on their pathogenesis and on the pertinence of the model to assess persistence strategies used by CF isolates, as well as investigate in vivo their ability to induce or reduce the production of proinflammatory cytokines.

Zebrafish embryo has been proposed as a novel vertebrate CF model, but the increased sensitivity of cftr morphants to

FIGURE 3 | Insights from infecting zebrafish embryo with Pseudomonas aeruginosa (GFP-expressing bacteria are shown in the inserted picture). Insights presented in this review, using diverse infection routes, are represented in green, while new appealing perspectives that are clinically relevant are highlighted in red. Created with BioRender.com.
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