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## **CFTR protects against *Mycobacterium abscessus* infection by fine-tuning host oxidative defences**

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## SUMMARY

Infection by rapidly-growing *Mycobacterium abscessus* is increasingly prevalent in cystic fibrosis (CF), a genetic disease caused by a defective CF transmembrane conductance regulator (CFTR). However, the potential link between a dysfunctional CFTR and vulnerability to *M. abscessus* infection remains unknown. Herein, we exploit a CFTR-depleted zebrafish model, recapitulating CF immuno-pathogenesis, to study the contribution of CFTR in innate immunity against *M. abscessus* infection. Loss of CFTR increases susceptibility to infection through impaired NADPH oxidase-dependent restriction of intracellular growth and reduced neutrophil chemotaxis, which together compromise granuloma formation and integrity. As a consequence, extracellular multiplication of *M. abscessus* expands rapidly, inducing abscess formation and causing lethal infections. Since these phenotypes are not observed with other mycobacteria, our findings highlight the crucial and specific role of CFTR in the immune control of *M. abscessus* by mounting effective oxidative responses.

## INTRODUCTION

Cystic fibrosis (CF) is a lethal genetic disorder caused by deleterious mutations in the CF transmembrane conductance regulator (CFTR) protein (Gadsby et al., 2006), resulting in compromised mucociliary clearance, chronic bacterial infections, and subsequent progressive inflammatory lung damages (Donalson and Boucher, 2003). CF-related lung infections are associated with a specific spectrum of colonizing microorganisms: highly prevalent bacteria such as *Staphylococcus aureus* or *Pseudomonas aeruginosa* (Lyczak et al., 2002) and also emerging bacterial pathogens, such as nontuberculous mycobacteria (NTM) (Olivier et al., 2003; Roux et al., 2009). Among the rapidly growing NTM, the *Mycobacterium abscessus* complex (MABSC) represents the most common species found in CF airways (Floto et al., 2016) and is emerging as a major CF pathogen due, in part, to indirect person-to-person transmission (Bryant et al., 2013), and progressing into severe pneumonia and accelerated inflammatory lung damages (Esther et al., 2010). Their presence is also a relative contraindication to lung transplantation (Orens et al., 2006). In particular, pulmonary infections with the multidrug-resistant *M. abscessus* subsp. *abscessus* (Mabs) (Nessar et al., 2012) are extremely challenging to treat, requiring aggressive and extended therapies with a high rate of therapeutic failure (DaCosta et al., 2017). To date, Mabs is widely considered to be the most significant rapidly growing NTM in CF with a worldwide prevalence rate of 5 to 20% (Floto et al., 2016).

Mabs exhibits two distinct morphotypes, relying on the presence or absence of surface-associated glycopeptidolipids (GPL): a smooth (S) high-GPL producer variant and a rough (R) low-GPL producer variant (Howard et al., 2006). Both morphotypes are recovered from the CF-airways during infection, but case reports indicate that the R form correlates with exacerbations of pulmonary disease and rapid decline of lung function in the patients (Jönsson et al., 2007; Catherinot et al., 2009). Nevertheless, understanding the specific vulnerability of the CF population to Mabs, the potential link with CFTR dysfunction and how these mycobacteria contribute to progression of lung disease remain unknown.

While it is assumed that susceptibility to infections in CF results from defective mucociliary activity, the CFTR dysfunction may also alter the inflammatory potential of innate immune cells contributing to the infectious pathology in this disease. Various hypotheses have attempted to explain the impairment of innate defences in CF, although these await definitive proof. Numerous mammalian models (Lavelle et al., 2016) have been generated to investigate the role of CFTR dysfunction in innate immunity and hyper-susceptibility to infections. However, these models are not suited for direct, real-time imaging of the early processes leading to disease development. Thus, new animal models that approximate the human altered immune phenotype and allowing direct visualization of host-pathogen interactions would provide the much-needed tools to establish *in vivo* how CFTR regulates innate immunity and controls Mabs infection.

Zebrafish (ZF) larval innate immunity is homologous to that of human (Renshaw and Trede, 2012) while their optical transparency allows non-invasive, real-time monitoring of infection outcomes and host-pathogen interactions. Thus, ZF innate immune cell behaviour and function can be visualized at sub-cellular resolution in the whole living animal, allowing dissecting the innate immune response during infectious diseases (Torraca et al., 2014). ZF led to new insights into the virulence of several CF bacteria, especially *P. aeruginosa* (Clatworthy et al., 2009), *S. aureus* (Prajsnar et al., 2008) and *Burkholderia cenocepacia* (Vergunst et al., 2010) and provide a useful model to study the pathophysiology of human Mabs infection (Bernut et al., 2014; Bernut et al., 2016). Importantly, from a structural perspective, ZF CFTR closely resembles the human protein (Zhang and Chen, 2016; Liu et al., 2017) and several reports suggest functional conservation of CFTR between ZF and human (Navis and Bagnat, 2015; Phennicie et al., 2010), making ZF larvae a clinically-relevant biological system. Indeed, several CF phenotypes that mirror human CF disease were reported in CFTR-defective ZF, including pancreatitis (Navis and Bagnat, 2015) or increased susceptibility to *P. aeruginosa* (Phennicie et al., 2010).

Herein, using CFTR-depleted ZF larvae as an innovative vertebrate model which recapitulates important aspects of the CF immuno-pathogenesis, we elucidated the role of CFTR in regulation of innate immunity to Mabs infections. We also interrogated the effects of CFTR ablation on host immunity, inflammation and infection independent of the overlapping infection and inflammation associated to CF-lung microenvironment. Importantly, our findings emphasize that differential CFTR-dependent ROS production allows the host to adjust inflammatory responses by modulating phagocyte bactericidal functions and their lifespan upon infection which, together, ensure the maintenance of a protective granulomatous structure to sequester and control Mabs infection.

## RESULTS

### Loss of CFTR function increases severity of *M. abscessus* infection

To address the role of CFTR in Mabs infection, *cftr* loss-of-function experiments were carried out in ZF using a specific morpholino-modified oligonucleotide (MO) (Figure S1A and S1B). While *cftr* is expressed and localized to the apical membrane or vesicular compartments of cells (Del Porto et al., 2011) (Figure S1C), *cftr*-MO injection abrogated production of native spliced *cftr* transcripts (Figure S1A and S1B) and altered *cftr* expression (Figure S1D). Embryos injected with *cftr*-MO survived similarly to control-MO injected animals and appeared morphologically similar throughout the observed periods (data not shown). Additionally, to support the knock-down results, we took advantage of the *cftr* ZF mutant (Navis et al., 2013). Upon intravenous infection, both *cftr* morphants and *cftr* mutants displayed hypersusceptibility to R or S Mabs morphotypes, correlating with increased larval mortality (Figure 1A) and higher bacterial burdens, as demonstrated by determination of the fluorescent pixel counts (FPC, Figure 1B) and whole larvae imaging (Figure 1C). The pronounced increase in bacterial loads in CFTR-deficient animals correlates with replicating extracellular bacteria (Figure 1D-H), translating into higher numbers of larvae with abscesses and with increased number of abscesses per larva in the central nervous system (Figure 1D and 1E). Mabs abscesses represent a marker of disease severity and uncontrolled infection caused by extracellular replicating mycobacteria that are often associated with cellular debris, tissue destruction and acute infection in zebrafish (Bernut et al., 2014). Whereas the S form induces abscesses only rarely in WT fish (Bernut et al., 2014), 30% of Mabs S-infected larvae exhibited abscesses at 3 dpi in absence of CFTR (Figure 1D). Electron microscopy (EM) analysis revealed that Mabs S abscesses in *cftr* morphant show enhanced replication of extracellular bacilli, promoting rapid bacterial expansion and tissue destruction (Figure 1F), similar to those reported in Mabs R abscesses found in WT fish (Bernut et al., 2014). Moreover, hypersusceptibility to Mabs infection in CF-fish is accompanied by increased bacterial cording in Mabs R-infected animals compared to WT larvae (Figure 1G and 1H). Collectively, these results indicate that *cftr* mutants recapitulate phenotypes induced by *cftr*-MO, implying that hypersusceptibility to Mabs in *cftr* morphants is not ascribed to off-target effects but to the direct consequences of *cftr* loss, thus validating the use of *cftr*-MO to further investigate the role of CFTR in innate immunity to Mabs.

Other MABSC subspecies or the rapid-growing NTM *Mycobacterium chelonae*, which is closely related to Mabs, can be isolated from CF expectorated sputum (Harris and Kenna, 2014). Infection of *cftr* morphants with *M. abscessus* subsp. *massiliense* (Figure S2A), *M. abscessus* subsp. *bolletii* (Figure S2B) or *M. chelonae* (Figure S2C) led to increased susceptibility to infections and larval killing, similarly to Mabs-infected CFTR-defective animals (Figure 1). In contrast, neither the non-pathogenic *Mycobacterium smegmatis* (Figure S2D) nor *Mycobacterium marinum*, one of the strict pathogenic NTM and closely related to *Mycobacterium tuberculosis* (Figure S2E), induced

increased larval killing in the absence of CFTR. These results indicate that the susceptibility to mycobacterial infections in CFTR-deficient embryos is specific and restricted to particular NTM species, such as those belonging to the *M. chelonae* complex (MCC), comprising MABSC and *M. chelonae*, and emphasize the protective role of CFTR in response to MCC infection by restricting bacterial pathogenesis and extracellular multiplication.

### **CFTR deficiency compromises *M. abscessus* granuloma maintenance by permitting rapid mycobacterial extracellular expansion**

Having previously demonstrated the importance of Mabs-induced granuloma formation and maintenance to prevent extracellular bacterial expansion and ensure the host defence (Bernut et al., 2014; Bernut et al., 2016), the increased mortality and extracellular mycobacterial growth in Mabs-infected CFTR-depleted animals prompted us to *i)* further characterize the granulomatous response to Mabs in the absence of CFTR and *ii)* ask if CFTR influences the course of granuloma formation and/or granuloma composition. Mabs-infected WT, *cftr* mutants and *cftr* morphants were compared and monitored over time for granuloma formation by fluorescent microscopy. In agreement with previous observations, Mabs-granulomatous lesions were found in both the presence and absence of CFTR (Figure 2A-G) (Tomashefski et al., 1996). However, while nascent granuloma appeared at 2 dpi and expanded in most infected WT embryos at 4 dpi (Figure 2A) (Bernut et al., 2014, Bernut et al., 2016), time-lapse microscopy revealed that the granuloma formation in CFTR-defective larvae remained unchanged with fewer granulomas observed in around 20-30% of the animals. No differences in the proportion of S or R granulomas were noticed in the context of CFTR impairment (data not shown). Additionally, confocal imaging showed that early granuloma formation and cellular aggregation events are maintained in the absence of CFTR function (Figure 2C), suggesting that granuloma elaboration proceeds regardless whether CFTR is present or not. Although this difference was not statistically significant, interestingly at all-time points, there was a trend towards increased average size of granuloma lesions in *cftr* morphants, which appeared more heavily infected than WT granulomas (Figure 2C and 2D) and continued expanding concomitantly with a time-dependent increased bacterial burden. This suggests that dysfunction of CFTR leads to a persistent acceleration in Mabs-granuloma growth beyond the initial aggregation event. Next, the late granuloma-like structure in absence of CFTR was explored by confocal imaging (Figure 2E and 2G) and EM (Figure 2F and 2H). Whilst the structures characterizing WT granulomas contained organized aggregate of phagocytes (Figure 2E) consisting of infected and uninfected cells surrounding a central necrotic region (Figure 2F), which efficiently control and sequester Mabs, we confirmed that impaired-CFTR granulomas are degenerated (Figure 2G and 2H), supporting the hypothesis that CFTR deficiency compromises granuloma maintenance. Overall, late CF-granulomas appear poorly delimited and contain dissociated cellular aggregates (Figure 2G and 2H) with abundant extracellularly replicating bacilli forming abscesses within tissues, presumably

responsible for the pronounced increase in phagocyte death observed in the granulomas (Figure 2H). In sharp contrast to WT Mabs-granuloma, microscopy revealed the profusion of highly infected phagocytes in CF-granulomatous lesions (Figure 2C), suggestive of impaired bacterial killing (Bernut et al., 2014; Bernut et al., 2016).

Together, these results indicate that a CFTR defect triggers the breakdown of Mabs-granulomas typified by extracellularly-growing bacteria released from dying phagocytes. This implies that CFTR is indispensable for normal granuloma structure and maintenance by controlling the phagocyte bactericidal functions and lifespan and preventing extracellular multiplication.

### **Dysfunction of CFTR impairs killing of *M. abscessus* in macrophages**

Our previous studies using Mabs-infected ZF highlighted the crucial role of macrophages (MΦ) for Mabs killing and infection control (Bernut et al., 2014; Bernut et al., 2016). In CF pathophysiology, the role of MΦ has been largely overlooked: scant evidence suggests altered MΦ properties in uncontrolled infection in CF lungs (Döring and Gulbins, 2009). To elucidate the cellular basis linking CFTR-deficiency with susceptibility to Mabs infection and alteration of the MΦ functions, expression of *cftr* was knocked-down in reporter lines harbouring labelled MΦ. We first examined if the lack of CFTR affects chemoattraction of MΦ to the invading bacteria and/or activation of these cells, as potential mechanisms promoting extracellular bacterial growth. Leucocyte mobilization was assessed by injecting fluorescent Mabs into the hindbrain ventricle (HBV) and their phagocytic capacity was monitored after intravenous infection of the bacilli. Deficiency of *cftr* compromised neither mycobacterial-induced migration to the infected HBV (Figure 3A) nor phagocytosis (Figure 3B) at early time points, suggesting that CFTR is not required for early interactions between Mabs and MΦ. Previous studies reported that dysfunction of CFTR is associated with reduced microbicidal capacities of immune cells (Assani et al., 2017; 2011; Di et al., 2006; Duranton et al., 2012). However, recent studies have shown comparable intracellular Mabs growth in murine MΦ carrying the CFTR<sup>ΔF508</sup> mutation as in control MΦ, suggesting that, in mice, functional CFTR is not required for the control of Mabs in MΦ infected *ex vivo* (Roux et al., 2016). Nevertheless, our microscopic observations show that CFTR ablation leads to hyperinfected phagocytes in granulomas, presumably relying on altered immune bacterial killing mechanisms (Figure 2C). Thus, to interrogate whether CFTR contributes to the MΦ mycobactericidal capacity *in vivo*, the number of Mabs in ZF MΦ was evaluated by confocal microscopy. The proportion of slightly infected (<5 bacilli), moderately infected (5–10 bacilli) or heavily infected (>10 bacilli) phagocytes was enumerated at 1 dpi. Compared to the control embryos, the *cftr* morphants infected by both R and S variants displayed a greater percentage of MΦ in the high burden category (Figure 3C). This is consistent with a reduced bactericidal ability and supports the hypothesis that CFTR controls intracellular growth and killing of Mabs. Because death of Mabs-infected phagocytes releases and propagates free bacilli in the extracellular milieu (Bernut et al., 2014), we examined the extent of



MΦ death in infected larvae. Combined confocal observations and quantification of acridine orange (AO)-positive infected MΦ shows that infection in *cftr* morphants is characterized by the presence of heavily infected phagocytes with an impaired ability to restrict bacterial growth, occurring prior to cell death (Figure 3D). While the basal levels of dead MΦ were equal between the PBS-injected control embryos and *cftr* morphants (data not shown), enumeration of AO-labelled MΦ infected by either R or S confirms higher yields of dead phagocytes in *cftr* morphants at 2 dpi as compared to the control embryos at 2 dpi (Figure 3E). This agrees with the presence of apoptotic immune cells seen in CF-granulomatous lesions (Figure 2). Importantly, the proportion of dead MΦ was lower in WT embryos infected with S- as compared to R-infected embryos (Bernut et al., 2014; Roux et al., 2016) but remains equal in absence of CFTR (Figure 3E), substantiating the crucial role of CFTR in containing intracellular *Mabs*.

### Modulation of the neutrophilic response to *M. abscessus* by CFTR

The chronic infections with a neutrophilic inflammation are a hallmark of CF lung pathophysiology (Cantin, 1995). Having reported that *Mabs* lesions are characterised by an influx of neutrophils and that these cells are critical in the host defence against *Mabs* infections (Bernut et al., 2014; Bernut et al., 2016; Malcolm et al., 2018), we next addressed whether CFTR ablation influences the behaviour of neutrophils by examining and comparing the dynamic of leucocyte mobilization in WT and CFTR-deficient larvae using the *Tg(mpx:GFP)i114* transgenic line labelling neutrophils with GFP (Renshaw et al., 2006). Surprisingly, CFTR deficiency strongly reduced neutrophil mobilization toward the infection sites, upon local injection of either R or S variants, as revealed by microscopy observations and quantitative analysis of the number of neutrophils at the site of infection at 4 hpi in *cftr* morphant (Figure 4A-D), and despite a higher baseline number of neutrophils in absence of CFTR (Figure S3A and S3B). These phenotypes are in line with the impaired neutrophil trafficking in *cftr* morphants infected with *P. aeruginosa* (Phennicie et al., 2010). Additionally, confocal microscopy unravelled an impaired neutrophil mobilization around and into *Mabs*-granulomatous lesions (Figure 4E and 4F) and abscesses (Figure 4G) in the *cftr* morphants. We previously established the linear relationship between the number of recruited neutrophils and the *Mabs*-granuloma volume in WT ZF (Bernut et al., 2016). Whilst WT granulomas contained numerous neutrophils distributed into and on the periphery of these cellular structures, CF-granulomas were mostly dominated by MΦ and contain fewer neutrophils (Figure 4E and 4F).

CF neutrophils remain capable of engulfing bacteria similarly to WT neutrophils (Figure S4A), suggesting that their phagocytosis activity toward *Mabs* is independent of CFTR. However, they fail to control intracellular *Mabs* (Figure S4B), leading to increased cell death (Figure S4C), as reported for MΦ (Figure 3). To determine whether the impaired neutrophilic response to *Mabs* infection in *cftr*-deficient ZF is linked to a possible intrinsic alteration in neutrophil recruitment, we performed a neutrophil mobilization assay using fMLP, a synthetic neutrophil chemoattractant. Injection of fMLP

into the otic vesicle induced similar neutrophil recruitment in *cftr* morphants and control larvae (Figure S5A). Previous findings highlighted also the requirement of IL8 for neutrophil trafficking during Mabs infections and its crucial role in elaborating Mabs-granuloma (Bernut et al., 2016). That CFTR impairs the early mobilization of neutrophils into Mabs-lesion led us to enquire whether CFTR depletion influences *cxc18* expression. Quantitative RT-PCR analysis revealed a similar level of *cxc18* expression in *cftr* morphants and control animals infected with both S and R variants (Figure S5B), indicating that impaired neutrophil trafficking is not caused by an alteration in the Cxcl8 pathway-mediated neutrophil mobilization.

Overall, these results suggest CFTR is required for early and late neutrophil recruitment to localized Mabs infection and to control the intracellular growth of Mabs.

### **Intracellular killing of *M. abscessus* is mediated by NOX2-dependent ROS production**

Mabs induces an oxidative stress response with the generation of intracellular reactive oxygen species (ROS) by MΦ and neutrophils (Bernut et al., 2016; Malcolm et al., 2018). Loss of functional CFTR reduces the MΦ respiratory burst response and impairs killing of intracellular *B. cenocepacia* (Assani et al., 2017). Thus, intracellular ROS generation in CFTR-deleted larvae was investigated as a plausible mechanism through which CFTR-mediated oxidative stress controls Mabs infections. Since production of intracellular ROS by professional phagocytes relies mainly on *nox2* (Warnatsch et al., 2017), knock-down experiments abolishing initial NOX2-mediated ROS production (Figure 5A) were conducted using a specific MO targeting *nox2* (Roca and Ramakrishnan, 2013). Whereas injection of *nox2*-MO failed to affect early MΦ mobilization into the HBV (Figure 5B) or MΦ phagocytosis (Figure 5C), it, however, led to higher numbers of heavily infected cells (Figure 5D). Consistent with these findings, global mycobacterial loads increased in the absence of NOX2 (Figure 5E). Consistently, the *nox2* morphants developed more abscesses than the control embryos, leading to the premature larval death (data not shown). This implies that reduced ROS production in *nox2* morphants is deleterious for the host and that the enhanced susceptibility of these fish to the infection highlight the key role of NADPH-produced intracellular ROS for clearing Mabs.

To determine whether an impaired oxidative defence is involved in defective CFTR-associated reduced bacterial killing in early infection stages, we examined whether heat-killed and intact Mabs induce intracellular ROS production in MΦ using the CellROX dye. ROS-labelled phagocytes were rapidly detected but the number of ROS-positive MΦ harbouring either living or heat-killed R or S variants were lower in *cftr* morphant than in control ZF at 2 hpi (Figure 6A and 6B), consistent with the altered ROS production reported in the CF context (Assani et al., 2017; Phennicie et al., 2010). Of note, the higher number of ROS-positive MΦ infected with heat-killed bacteria suggests that Mabs has developed mechanisms to overcome the host oxidative killing mechanisms. Moreover, similar results were observed regarding reduced ROS production in CF-infected neutrophils (Figure

S6A and S6B). To further inquire whether the impaired intracellular ROS production in absence of CFTR is linked to a possible decrease in NADPH oxidase activity, the NOX2-mediated ROS signalling pathway was dissected in defective *cftr*-embryos. While Mabs infection triggers an upregulation of *nox2* in control fish, qRT-PCR analysis confirmed a reduced *nox2* expression in *cftr* morphants (Figure 6C), suggesting that CFTR orchestrates the early regulation of ROS induction by modulating the NOX2/NADPH oxidase activity. That the intracellular bacterial profiles were similar in the *cftr* morphants and in the double *cftr/nox2* morphants (Figure 6D) suggests that the enhanced intracellular bacterial growth in *cftr* morphants is directly linked to defective NOX2-NADPH oxidase-related ROS production. Overall, these results demonstrate that CFTR deficiency alters NOX2/NADPH oxidase-dependent ROS production which, in turn, fails to restrict intracellular growth of Mabs and that the release of NOX2-derived ROS participates in clearing Mabs in both MΦ and neutrophils (Figure 6 and Figure S6).

Activated MΦ restrict mycobacterial growth through a TNF-mediated ROS dependent pathway (Bernut et al., 2016; Dewas et al., 2003). Combined with the fact that CFTR dysfunction is associated with alterations in the innate immune regulation network and pro-inflammatory cascades (Döring and Gulbins, 2009; Cantin, 1995), we next explored whether abnormal *tnf* induction could be involved in reduced ROS-mediated bacterial killing in *cftr* morphants during the early stages of infection. To assess the effect of *cftr* loss-of-function on *tnfa* production by Mabs-infected MΦ, Mabs E2-Crimson were injected into the muscle of either WT or *cftr* morphants *tnfa:GFP-F/mpeg1:mCherry-F* double transgenic. Microscopy observations indicate that both control or CF animals exhibited equal proportions of GFP-positive infected phagocytes containing either Mabs S or R at 4 hpi (Figure S7A and S7B), suggesting that the early impaired ROS generation associated with CFTR ablation is TNF-independent. At later stages, however, CFTR ablation triggers a hyper-inflammatory response following infection with Mabs, with qRT-PCR revealing upregulation of *tnfa* expression, especially after infection with the R form (Figure S7C), similar to findings reported previously in mice (Catherinot et al., 2009).

Collectively, these data indicate that NOX2/NADPH oxidase-dependent ROS production by infected phagocytes represents a critical host defence mechanism against Mabs and suggests that the inherent deficit in NOX2-derived oxidative stress in CF leukocytes is responsible for their defective bacterial killing responses.

## DISCUSSION

Pulmonary disease is the leading cause of morbidity and mortality in CF and is characterized by a vicious circle of chronic infections and persistent inflammation. Among the deleterious bacteria found in CF-airway, the fast-growing multidrug-resistant *Mabs* has emerged as an important respiratory pathogen of major concern in CF centers worldwide ([Parkins and Floto, 2015](#)). However, our understanding of the particular vulnerability of CF patients to *Mabs* infection remains limited by the lack of suitable animal models mimicking the immune abnormalities found in the CF population. Nevertheless, important insights into the pathophysiology of *Mabs* diseases have recently been obtained in the ZF ([Bernut et al., 2014](#); [Bernut et al., 2016](#)) and the very close structural relatedness between ZF and human CFTR emphasizes further the relevance of ZF to study CFTR functions ([Zhang and Chen, 2016](#); [Liu et al., 2017](#)). To address these unmet needs, we exploited here CF-ZF as an innovative vertebrate recapitulating aspects of CF immuno-pathogenesis. Thanks to genetic and high resolution imaging approaches, we report the direct stepwise dissection of *Mabs* infection in an animal lacking CFTR to elucidate the biological implication of CFTR in innate immunity to *Mabs*.

*Mabs*-infected CFTR-depleted ZF rapidly succumb to infection, reflecting a hypersusceptibility to this mycobacterium in CF, providing a first glimpse into CFTR-mediated host defences to *Mabs* infection. Mechanisms leading to the formation of protective *Mabs* granuloma depends on efficient M $\Phi$  and neutrophil cooperation, orchestrated by fine-tuning innate immune responses ([Bernut et al., 2016](#)). The spatiotemporal events associated with CFTR ablation ([Figure 7](#)) reveal a mechanism whereby CFTR participates in neutrophil chemotaxis to the infected sites and the adjustment of oxidative host defences, conditioning efficient phagocyte-mediated bacterial killing, together generating a protective granulomatous response. Infection of ZF with *Mabs* is characterized by: *i*) rapid engulfment of the bacilli by M $\Phi$ ; *ii*) activation of M $\Phi$ , resulting in chemotaxis guiding neutrophils to pre-forming granulomas and ROS production by NOX2 for intracellular killing of *Mabs*; *iii*) homeostatic granuloma formation to sequester *Mabs*, containing the infection and favouring the development of chronic disease. Conversely, CFTR deletion promotes increased susceptibility to *Mabs* infections, correlating with *i*) deficiency in ROS production altering phagocyte-mediated *Mabs* killing, resulting in increased intracellular bacterial loads and premature cell death and *ii*) impaired neutrophil chemotaxis toward nascent granulomas. These two factors conspire to alter the maintenance of protective granuloma with uncontrolled extracellular mycobacterial spread, conducting to acute infection and larval death. Other studies suggested that dysfunction of CFTR dampens the microbicidal activity of immune cells ([Assani et al., 2017](#); [Di et al., 2006](#); [Duranton et al., 2012](#)), promoting infectious pathology in CF airways. However, existing CF models have failed to reproduce the hypersusceptibility phenotype associated with mycobacterial infections in CF ([Roux et al., 2016](#); [Le Moigne et al., 2015](#)). Using the CF-ZF model, we report here that primary

alterations in innate immunity directly contribute to increased susceptibility to the infection. Whereas ROS produced by the NADPH oxidase during the respiratory burst participate in the elimination of pathogens, Mabs has been reported to withstand the hostile oxidative environments inside phagocytes (Oberley-Deegan et al., 2010), although this awaits *in vivo* confirmation. In ZF, inhibition of the NOX2/NADPH oxidase pathway enhances intracellular growth. Given the importance of NOX2-derived ROS production in Mabs killing, a reduced oxidative response in CF-ZF is very likely to explain the increase susceptibility to Mabs. Of note, the bacterial burden in *nox2*-defective MΦ is lower than in CFTR-MΦ (Figure 6D), suggesting that other CFTR-mediated microbicidal mechanisms participate in Mabs clearance.

Overly exuberant neutrophil influx associated with harmful oxidative stress is a hallmark of the inflammatory CF lungs (Cantin, 1995; Hector et al., 2014). The increased number of neutrophils in *cfr* morphants mirrors the neutrophilia seen in CF. Our results emphasize also the neutrophil chemotaxis impairment to Mabs, as shown previously in *P. aeruginosa*-infected ZF (Phennicie et al., 2010), but raising also controversial questions regarding inflammatory and infectious CF pathologies. Supporting the view that neutrophilic inflammation-mediated Mabs infection plays a critical role in host defence against this pathogen by maintaining granuloma integrity and preventing extracellular bacterial multiplication (Bernut et al., 2016), we show that the capacity of neutrophils to migrate in a CFTR-dependant manner is involved in the formation of protective granulomas. Release of ROS by epithelial cells through NOX2-NADPH oxidase has been implicated in neutrophil chemotaxis to wounds (Braunersreuther et al., 2013). We provide here evidence for CFTR in modulating the NOX2 oxidative pathway, in which a local redox imbalance in *cfr* morphants may account for the reduced number of infiltrated neutrophils toward Mabs. This suggests that NOX2 dependent-ROS production from activated leucocytes triggers an oxidative environment sustaining neutrophilic mobilization to efficiently contain bacteria within homeostatic granulomas. The NOX/DUOX family NADPH oxidases-mediated oxidative defences play a critical role to control invading pathogens by triggering neutrophil chemotaxis to infected tissues or bacterial killing (Brothers et al., 2013; Rada and Leto, 2008). In addition to the reduced *nox2* expression in absence of CFTR, presumably contributing to imbalanced adjustments of redox signalling, other mechanisms altering ROS production *via* other NADPH oxidase complexes by epithelial or immune cells may also contribute to the impaired bactericidal function and/or chemotaxis of leucocytes during Mabs infection.

The clinical relevance of ROS production in host defence, notably the NOX2/NADPH oxidase in granuloma formation, is consistent with mutations in *nox2* leading to chronic granulomatous disease (CGD) typified by the development of large size and poorly structured granulomas that are unable to sequester mycobacteria (Deffert et al., 2014) and associated with severe inflammation (Rieber et al., 2012). However, although the altered NADPH oxidase function in CGD can lead to increased disease severity following infection with *M. tuberculosis* or vaccination with *M. bovis* BCG (Deffert

et al., 2014), reports of infection with NTM remain anecdotal (Ohga et al., 1997; Weening et al., 2000; Chusid et al., 1975) and no particular link with *M. abscessus* has been ascribed yet.

This, together with clinical cases of CF patients heavily infected with *Mabs* R (Jönsson et al., 2007; Catherinot et al., 2009) demonstrates that *Mabs* exacerbates inflammation in absence of CFTR, implicating a critical inflammatory pathology associated with tissue damage and persistent *Mabs* infections. Thus, at the later stages of infection, it is possible that an imbalance of ROS production or neutrophil chemotaxis could be caused by CF-mediated hyperinflammation acting as a negative-feedback loop which would undo the fine tuning of immune responses.

Our study indicates that *cftr* is a regulator of host immunity to MABSC, but not to other saprophytic (*M. smegmatis*) or pathogenic (*M. marinum*) NTM, suggesting that species-specific restriction mechanisms may exist for these organisms. In fact, we show that *M. marinum*-infected *cftr* morphant succumbed to infection more slowly than controls, suggesting that inactivation of *cftr* triggers a mild protective immunity against tuberculosis. Indeed, it has been proposed that the high carrier rate for CFTR mutations among Caucasians is due to an evolutionary selective advantage against infectious disease, with candidate agents including cholera (Gabriel et al., 1994), typhoid fever (Pier et al., 1998) and tuberculosis (Meindl, 1987), in which the pandemic in the early 1600s could explain the modern-day CF incidence rates in Euro-descendent populations (Poolman and Galvani, 2007). Recent studies support the hypothesis that carrying the most common F508del *cftr* allele protects against *M. tuberculosis* infection (Bosch et al., 2017). This species-specific susceptibility to different mycobacteria is particularly intriguing and deserves further attention.

In summary, we demonstrate that CFTR dysfunction leads to hypersusceptibility to *Mabs* infection *in vivo*, potentially explaining the high rates of infection seen clinically in CF patients. We anticipate that insights obtained from ZF may guide the development of future therapies targeting innate immune defects in CF.

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## AUTHORS CONTRIBUTIONS

A.B. and L.K. conceived the study, and wrote the manuscript with input from J.L.H., R.A.F., and S.A.R.; A.B., S.A.R. and L.K. designed experiments and analysed data; A.B. performed experiments and produced the figures; C.D. performed PCR experiments; N.V.O. made and provided *Tg(mpeg1:NLSclover)sh436* line; A.N. performed EM preparation and imaging experiments; All authors reviewed the manuscript.

## DECLARATION OF INTEREST

The authors declare no competing interests.

## SUPPLEMENTAL INFORMATION

Supplemental Information includes 1 table and 7 figures.



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## FIGURE LEGENDS

### Figure 1. *cftr* deficiency increases susceptibility to *M. abscessus* infection

(A-H) WT, *cftr* mutant or *cftr* morphant were *iv* infected with either Mabs R or S expressing tdTomato.

(A) Survival analysis of R- (top graph) or S-infected larvae (bottom graph). Data are plotted as percentage of surviving animal over 10 days (n=60, three experiments).

(B) Mean bacterial loads as fluorescence pixel counts (FPC, average of three independent experiments) of 3 dpi larvae infected by either Mabs R (top) or S (bottom).

(C) Representative overlay fluorescence microscopy images of Mabs R- (top) or S-infected larvae (bottom) at 3 dpi. Scale bars, 200  $\mu$ m.

(D-E) Percentage of 3 dpi larvae with abscess (D) and associated mean  $\pm$  standard error of the mean (SEM) number of abscess per infected animal (E) (n=45, three experiments).

(F) EM showing a sagittal section through a Mabs S-abscess in a *cftr* morphant. Overview image of the abscess into the spinal cord (top; scale bar, 25  $\mu$ m). Representative EM image of the abscess lesion with many extracellular bacterial in an area of acellular necrotic debris (bottom; scale bar, 0.5  $\mu$ m). Close-up showing bacterial division with yellow arrows indicating septum of division (scale bar, 0.5  $\mu$ m).

(G-H) Kinetic of Mabs R cording in whole embryos over 4 days of infection (G) and associated mean  $\pm$  SEM number of cords per infected animal (H) (n=45, three experiments).

See also [Figure S1](#) and [S2](#).

### Figure 2. Absence of CFTR impairs *M. abscessus* infection-mediated granuloma integrity and maintenance

(A-D) WT, *cftr* mutant or *cftr* morphant were *iv* infected with S variant of Mabs expressing tdTomato and monitored by confocal microscopy for granuloma formation, number and size (n=35-45, data are plotted as mean  $\pm$  SEM from three experiments).

(A-B) Kinetics of granuloma formation in whole embryos over 4 days of infection (A) and associated number of granuloma per animal (B).

(C) Confocal images showing a representative early Mabs granuloma in 3 dpi larvae. Scale bars, 10  $\mu$ m. For a similar number of infected phagocytes (asterisk) in a granuloma, the CF granuloma exhibits a higher bacterial burden than the WT granuloma.

(D) Granuloma volume analysis in whole larvae over 4 days of infection.

(E-H) Control larvae or *cftr* morphants infected with Mabs S expressing tdTomato observed by confocal microscopy and EM for granuloma at 4 dpi.

(E and G) Representative confocal imaging of granulomas within the spinal cord of an infected animal showing the development of a compact and organized phagocyte aggregates moderately

infected in control fish (E) compared to a *cftr* morphant harbouring a heavily-loaded granuloma whose disruption leads to bacterial spread characterized by abscess formation (G). Scale bars, 10  $\mu$ m.

(F) EM showing a sagittal section through a WT Mabs granuloma. Overview image of granuloma (arrow) into the brain of infected WT (top; scale bar, 50  $\mu$ m). Representative EM image of a compact and well-organised WT granuloma (bottom and right panels; scale bars, 2  $\mu$ m) showing a stable mycobacterial-containing structure with a central necrotic area and a typical dense region of cellular debris and surrounded by numerous infected phagocytes (asterisk) and giant cells (arrow).

(H) EM showing a sagittal section through a Mabs granuloma-like lesion in absence of CFTR. Overview image of granuloma in the spinal cord (arrow) of infected *cftr* morphant (top; scale bar, 25  $\mu$ m). Representative EM image of a CFTR-depleted granuloma (bottom and right panels; scale bar, 2  $\mu$ m) showing a necrotic structure with replicating bacteria and numerous dead infected phagocytes harboring a typical apoptotic nucleus (asterisk) with chromatin "superaggregation". Most bacilli stay extracellular.

### **Figure 3. *cftr* knockdown diminishes intracellular killing of *M. abscessus* and promotes macrophage death**

(A) Control or *cftr* morphants *Tg(mpeg1:NLSmClover)sh436* were infected with Mabs R or S expressing tdTomato into the hindbrain ventricle (HBV). Confocal microscopy was used to monitor the cell recruitment at 2 hpi. Mean  $\pm$  SEM number of M $\Phi$  recruited to the infected HBV (n=20, two experiments).

(B) *mpeg1:NLSmClover* control or *cftr* morphants were *iv* infected with Mabs R or S expressing tdTomato. Mean  $\pm$  SEM number of infected M $\Phi$  in the caudal hematopoietic tissue (CHT) at 4 hpi (n=20, two experiments).

(C-D) *mpeg1:NLSmClover* control or *cftr* morphants were *iv* infected with Mabs R or S expressing dtTomato imaged at 1 dpi using confocal microscopy to quantify the intracellular bacterial loads.

(C) Average proportions of infected M $\Phi$  containing <5, 5 to 10, or >10 bacteria in the CHT (n=16, two experiments).

(D) Confocal images showing infected M $\Phi$ . While WT-M $\Phi$  efficiently contain intracellular bacilli, CF M $\Phi$  fail to control Mabs growth. Arrow indicates intracellular Mabs R cording. Scale bars, 2  $\mu$ m.

(E) Control or *cftr* morphants *Tg(mpeg1:mCherry-F)ump2* were *iv* infected with Mabs R or S expressing E2-Crimson and stained with acridine orange (AO). Dead infected M $\Phi$  in the CHT were counted using confocal microscopy at 2 dpi. Data are plotted as mean  $\pm$  SEM from two experiments (n=20-22).

#### Figure 4. *cftr* deficiency reduces chemoattraction of neutrophils to infection sites

(A-G) Controls or *cftr* morphants *Tg(mpx:GFP)i114* were infected in the otic cavity (A-B), the muscle (C-D) or the caudal vein (E-G) with Mabs R or S expressing tdTomato and monitored by confocal microscopy to follow the neutrophil behavior toward infection sites.

(A-D) Representative images (A and C) and mean  $\pm$  SEM number (B and D) of neutrophils recruited to infection sites after 3 hpi (n= 30, three experiments). Scale bars, 50  $\mu$ m.

(E) Number of neutrophils recruited to WT (upper panel) or CFTR-depleted (lower panel) nascent granulomas as a function of granuloma volume.

(F-G) Confocal images showing the distribution of a neutrophil-associated granuloma (F) and abscess (G) in a control animal *versus* a *cftr* morphant. Scale bars, 25  $\mu$ m.

See also [Figures S3](#), [S4](#) and [S5](#).

#### Figure 5. NADPH-oxidase activity-mediated intracellular ROS production restricts *M. abscessus* growth

(A) *mpeg1:NLSmcllover* controls or *nox2* morphants were infected with Mabs expressing dtTomato and stained for ROS production using CellRox Deep red. Representative ROS-producing infected M $\Phi$  revealed by confocal microscopy in a *nox2* morphant *versus* a WT larvae. Scale bars, 5  $\mu$ m.

(B) *mpeg1:NLSmcllover* controls or *nox2* morphants were infected with Mabs expressing tdTomato into the HBV and monitored by confocal microscopy to determine cell recruitment. Mean  $\pm$  SEM number of M $\Phi$  recruited to the infected HBV at 2 hpi (n=20, two experiments).

(C-E) *mpeg1:NLSmcllover* controls or *nox2* morphants were *iv* infected with Mabs S (C and D) or R (D) expressing tdTomato. (C) Mean number of infected M $\Phi$  in the CHT at 4 hpi (n=20, two experiments).

(D) Average proportions of infected M $\Phi$  containing <5, 5 to 10, or >10 bacteria in the CHT at 1 dpi (n=16, two experiments).

(E) Mean FPC of 3 dpi larvae *iv* infected by either Mabs R (left) or S (right) expressing tdTomato from three experiments.

#### Figure 6. CFTR modulates NADPH oxidase-mediated ROS production

(A-B) *mpeg1:NLSmcllover* controls or *cftr* morphants were infected with either living or heat killed Mabs R or S expressing dtTomato into the muscle and stained for ROS production using CellRox Deep red and analyzed by confocal microscopy.

(A) Proportion of ROS-positive infected M $\Phi$  at 2 hpi (n=16, two experiments).

(B) Distribution of representative ROS-producing M $\Phi$  within the muscle at 2 hpi. Scale bars, 15  $\mu$ m.

(C) qRT-PCR measurement in whole embryos *iv* infected with Mabs and plotted as fold increase over mock injection for *nox2*. Mean relative  $\pm$  SEM gene expression of three independent replicates.

(D) *mpeg1:NLSclover* controls, *cftr*, *nox2* or double *cftr/nox2* morphants were *iv* infected with Mabs expressing dtTomato and intracellular bacterial loads were quantified at 1 dpi using confocal microscopy. Graph represents the average proportions of infected MΦ containing <5, 5 to 10, or >10 bacteria in the CHT (n=15, two experiments).

See also [Figures S6](#) and [S7](#).

**Figure 7. Critical role of the CFTR/NADPH oxidase axis for efficient ROS production and protective immunity against *M. abscessus* infection**

Schematic overview summarizing the CFTR/NOX2 axis-dependent ROS production in Mabs infection control.



## STAR★METHODS

### Key Resources Table

REAGENT or RESOURCE	Sources	Identifiant
<b>Experimental Model: Bacterial Strains</b>		
<i>Mycobacterium abscessus sensu stricto</i> , strain CIP104536 <sup>T</sup> , smooth	Laboratoire de Référence des Mycobactéries (IP, France)	ATCC19977 <sup>T</sup>
<i>Mycobacterium abscessus sensu stricto</i> , strain CIP104536 <sup>T</sup> , rough	Laboratoire de Référence des Mycobactéries (IP, France)	ATCC19977 <sup>T</sup>
<i>Mycobacterium abscessus subsp. bolletii</i> , CIP108541 <sup>T</sup> , smooth	Laboratoire de Référence des Mycobactéries (IP, France)	CIP108541 <sup>T</sup>
<i>Mycobacterium abscessus subsp. bolletii</i> , CIP108541 <sup>T</sup> , rough	<a href="#">Bernut et al., 2016b</a>	CIP108541 <sup>T</sup>
<i>Mycobacterium abscessus subsp. massiliense</i> , CIP108297 <sup>T</sup> , smooth	Laboratoire de Référence des Mycobactéries (IP, France)	CIP108297 <sup>T</sup>
<i>Mycobacterium abscessus subsp. massiliense</i> , CIP108297 <sup>T</sup> , rough	Laboratoire de Référence des Mycobactéries (IP, France)	CIP108297 <sup>T</sup>
<i>Mycobacterium marinum</i> strain M	<a href="#">Stinear et al., 2008</a>	ATCC BAA-535
<i>Mycobacterium chelonae</i> strain A6	Laboratoire de Référence des Mycobactéries (IP, France)	N/A
<i>Mycobacterium smegmatis</i> mc <sup>2</sup> 155	<a href="#">Snapper et al., 1990</a>	N/A
<b>Experimental Model: Zebrafish lines</b>		
golden mutant	<a href="#">Lamasson et al., 2005</a>	N/A
<i>cftr</i> <sup>pd1049</sup> mutant	<a href="#">Navis et al., 2013</a>	N/A
<i>gBAC(cftr-RFP)pd1042</i>	<a href="#">Navis et al., 2013</a>	N/A
<i>Tg(mpx:GFP)i114</i>	<a href="#">Renshaw et al., 2006</a>	N/A
<i>Tg(LysC_DSred)nz5</i>	<a href="#">Hall et al., 2007</a>	N/A
<i>Tg(mpeg1:NLSmClover)sh436</i>	This study	N/A
<i>Tg(mpeg1:mCherry-F)ump2</i>	<a href="#">Bernut et al., 2014</a>	N/A
<i>Tg(tnfr:GFP-F)ump5</i>	<a href="#">Nguyen-Chi et al., 2015</a>	N/A
<b>Chemicals, Peptides, and Recombinant Proteins</b>		
Hygromycin B	Sigma-Aldrich	Cat# H3274
Tricaine	Sigma-Aldrich	Cat# E10521
Difco™ Middlebrook 7H9 Broth	Thermo Fisher Scientific	Cat# DF0713-17-9
Middlebrook OADC Growth Supplement	Sigma-Aldrich	Cat# M0678
Tween-80	Sigma-Aldrich	Cat# P1754
f-Met-Leu-Phe (fMLP)	Sigma-Aldrich	Cat# F3506
Acridine Orange	Invitrogen	Cat# 93001
CellROX™ Deep Red	Invitrogen	Cat# C10422
LR Clonase II Plus	Invitrogen	Cat# 12538
SuperScript™ IV First-Strand Synthesis System	Invitrogen	Cat# 18091050
LightCycler® 480 SYBR Green I Master	Roche	Cat# 04887352001
<b>Recombinant DNA</b>		
pTEC15	Addgene	Cat# 30174
pTEC19	Addgene	Cat# 30178
pTEC27	Addgene	Cat# 30182
pcDNA3.1-Clover-mRuby2	Addgene	Cat# 49089
Gateway™ pDONR™221 Vector	Invitrogen	Cat# 12536017
pCSTP	<a href="#">Kawakami et al., 2014</a>	N/A

<b>Critical Commercial Assays</b>		
Nucleospin RNAII kit	Macherey-Nagel	Cat# 740955
<b>Oligonucleotides</b>		
Primers used in this study are listed in Table S1		
<b>Software and Algorithms</b>		
Prism 7.0	Graphpad	<a href="https://www.graphpad.com/">https://www.graphpad.com/</a> ; RRID:SCR_002798
R 3.5.0	R core team	<a href="http://www.r-project.org/">http://www.r-project.org/</a> ; RRID:SCR_001905
ImageJ	NIH	<a href="https://imagej.nih.gov/ij/">https://imagej.nih.gov/ij/</a> ; RRID:SCR_003070
TIA Software	Thermo Scientific™ Tecnai	<a href="https://www.fei.com">https://www.fei.com</a>
Volocity 6.3	PerkinElmer Life and Analytical Sciences, Cambridge, UK	<a href="http://www.perkinelmer.com/fr/lab-products-and-services/resources/whats-new-volocity-6-3.html">http://www.perkinelmer.com/fr/lab-products-and-services/resources/whats-new-volocity-6-3.html</a> RRID:SCR_002668
LAS-AF	Leica Microsystems	<a href="https://www.leica-microsystems.com/products/microscope-software.html">https://www.leica-microsystems.com/products/microscope-software.html</a>
Zen (Blue edition)	Zeiss	<a href="https://www.zeiss.com/microscopy/int/products/microscope-software/zen.html">https://www.zeiss.com/microscopy/int/products/microscope-software/zen.html</a>
LightCycler® 480 Software	Roche	<a href="https://lifescience.roche.com/en_gb/products/lightcycler14301-480-software-version-15.html">https://lifescience.roche.com/en_gb/products/lightcycler14301-480-software-version-15.html</a>

## CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Laurent Kremer, ([laurent.kremer@irim.cnrs.fr](mailto:laurent.kremer@irim.cnrs.fr)).

## EXPERIMENTAL MODELS AND SUBJECT DETAILS

### Bacterial Strains

Mycobacterial strains carrying pTEC15 (Addgene, plasmid 30174), pTEC27 (Addgene, plasmid 30182) or pTEC19 (Addgene, plasmid 30178) that express green fluorescent protein (Wasabi), red fluorescent protein (tdTomato) or bright far-red fluorescent protein (E2-Crimson), respectively, were grown under hygromycin B selection in Middlebrook 7H9 supplemented with oleic acid, albumin,

dextrose, catalase (OADC), and 0.05% Tween-80. To prepare heat-killed Mabs, bacteria were incubated at 80°C for 20 min.

### **Zebrafish Husbandry and Ethic statements**

Experimental procedures were performed using the *golden* mutant (Lamasson et al., 2005), the *cftr*<sup>pd1049</sup> mutant (Navis et al., 2013), the transgenic lines *Tg(mpx:GFP)i114* (Renshaw et al., 2006) and *Tg(LysC\_DSred)nz5* (Hall et al., 2007) to visualize neutrophils; *Tg(mpeg1:NLSclover)sh436* and *Tg(mpeg1:mCherry-F)ump2* (Bernut et al., 2014) to visualize MΦ ; *gBAC(cftr-RFP)pd1042* (Navis et al., 2013) to visualize *cftr* expression, and *Tg(tnfα:GFP-F)ump5* (Nguyen-Chi et al., 2015) to visualize *tnf-α* expression. ZF were raised and maintained according to standard protocols (Nusslein-Volhard and Dahm, 2002). Eggs were obtained from pairs of adult fish by natural spawning and raised at 28.5°C in tank water. The ZF husbandry and all ZF experiments described in the present study were conducted in accordance with guidelines from the UK Home Office (Bateson Centre, University of Sheffield) and in compliance with the European Union guidelines for handling of laboratory animals (CNRS, Montpellier) and were approved by the Direction Sanitaire et Vétérinaire de l'Hérault et Comité d'Ethique pour l'Expérimentation Animale de la région Languedoc Roussillon under the reference CEEA-LR-1145 (Montpellier experiments).

### **METHOD DETAILS**

#### **Creation of the *Tg(mpeg1:NLSmClover)* transgenic line**

The Tol2kit multisite Gateway-based transposon system (Kwan et al., 2007) was used to generate a construct from which a stable transgenic line was raised. mClover was amplified from pcDNA3.1-Clover-mRuby2, a gift from Kurt Beam (Addgene, plasmid 49089), using forward primer 5'-GGGGACAAGTTTGTACAAAAAAGGCTCAATGGCTCCAAAGAAGAAGCGTAAGGTA-3' and reverse primer 5'-GGGGACCACTTTGTACAAGAAAGCTGGGTCTACTTGTACAGCTCGTCCA-3' and cloned into pDONR 221 donor vector (Invitrogen) to produce pME-NLSmClover. An LR Clonase II Plus (Invitrogen) Gateway reaction was performed with p5E-mpeg1 (Ellet et al., 2011), pME-NLSmClover and p3E-polyA inserted into pDestTol2pA2 destination vector (Kwan et al., 2007) to produce *mpeg1:NLSmClover* construct. tol2-transposase mRNA was synthesised from pCSTP (Kawakami et al., 2014) then co-injected with *mpeg1:NLSmClover* construct into zebrafish one-cell-stage embryos to create the *Tg(mpeg1:NLSmClover)sh436* transgenic line harboring green fluorescent MΦ.

#### **Morpholino injection**

MO were purchased from Gene Tools. *cftr* splice-blocking morpholino targeting *cftr* (ZFIN, ZDB-GENE-050517-20) (5'-GACACATTTTGGACACTCACACCAA-3') was injected into one-cell-stage

zebrafish embryos (1 mM, 2 nl). The efficiency of gene knockdown was confirmed by RT-PCR and sequencing with the following primers for both sides of the MO target sequence (forward and reverse): CCTGTGGAGGATGCCAACTGCC and TGCATGCCCAGGTGGTGCAG. Total RNA from 2 dpf embryos (pools of 10 each) was isolated and purified with Nucleospin RNAII kit (Macherey-Nagel), following the manufacturer's instructions. The SuperScript IV reverse transcriptase (Invitrogen) was used to synthesize first-strand cDNA with oligo(dT) primer from 1 µg of total RNA at 50°C for 50 min. MO for *nox2* (*cybb*, ZFIN, ZDB-GENE-040426-1380) knockdown (5'-CATAATCCCGATAGCTTACGATAAC-3') was prepared and injected as described earlier ([Roca and Ramakrishnan, 2013](#)). A standard control-MO was included as a negative control in all experiments.

### **Zebrafish infection**

Bacterial inoculate were prepared for infection challenges in ZF embryos/larvae, according to procedures described earlier ([Bernut et al., 2015](#)). Briefly, systemic infections were carried out by the injection of single-cell suspensions of known titer (100-200 colony-forming units) into the caudal vein of 30 hrs post-fertilization (hpf) embryos. For leucocyte mobilization assays, mycobacteria were locally injected into the hindbrain ventricle at 30 hpf, the otic vesicle or the muscle compartment of 3 days post-fertilization (dpf) larvae. The number of animals used for each procedure was guided by pilot experiments or by past results ([Bernut et al., 2016a](#); [Bernut et al., 2014](#)).

### **qRT-PCR**

Total RNA was prepared and first-strand cDNA synthesized. Real-time RT-PCRs were performed with an LightCycler® 480 system instrument using LightCycler® 480 SYBR Green I Master (Roche) and gene expressions were detected with gene-specific primers listed in [Table S1](#). Reaction mixtures were incubated for 5 min at 95°C, followed by 45 cycles of 5s at 95°C, 20s at 65°C, and finally 10s at 95°C. Each experiment was run in triplicate. qRT-PCR data are analysed with the LightCycler® 480 Software, normalized to the housekeeping gene *ef1α* and calculated using the  $\Delta\Delta C_t$  method.

### **Epifluorescence, Confocal Microscopy and Imaging**

To quantify bacterial loads, granulomas (defined as cellular aggregates comprising at least 10 infected cells), cords, leukocyte mobilization, and evaluate oxidative stress, infected larvae were tricaine-anesthetized and mounted in 0.8% low melting point agarose with 0.016% tricaine for real-time microscopy observations. To evaluate intracellular mycobacterial growth, phagocytosis, cellular mortality, and granuloma organization, infected animals were tricaine-anesthetized, fixed overnight at 4°C in 4% paraformaldehyde in PBS, washed twice in PBS and then transferred gradually from PBS to 50% glycerol for microscopy observation.

Epifluorescence microscopy was performed using a Zeiss Plan Neo Fluor Z 1x/0.25 FWD objective and equipped with an Axiocam503 monochrome (Zeiss) camera. Pictures were taken and processed using ZEN 2 (blue edition). Confocal microscopy was performed using a Spinning disc confocal Perkin Elmer Ultraview VoX inverted Olympus IX81 with a UplanSAPO 20x/0.8 and a UplanSAPO 40x/1.3 oil objective, equipped with a Hamamatsu C9100-50 EM-CCD camera or with a Leica SPE upright microscope with a ACS APO 40x/1.15 oil objective. Images were captured and processed using the Velocity or LASAFS softwares.

### **Transmission Electron Microscopy**

For TEM, ZF larvae were tricaine-anesthetized and fixed overnight at 4°C in 2.5% glutaraldehyde in 0.1M phosphate buffer. After a post-fixation in 1% osmic acid 1 hr at 4°C and 0.5% tannic acid for 30 min at 4°C, animals were dehydrated in successive ethanol baths, infiltrated with mixes of epon 812/propylene oxide, embedded in epon 812 resin, and then polymerised at 60°C for 48 hrs. Ultrathin sections were cut with a Reichert Ultracut ultramicrotome (Leica) and collected on nickel grids. Finally, the grids were stained 20 min in 2% uranyl acetate and 3 min in 3% lead citrate, then examined on a Tecnai G2 F20 (200kV, FEG) electron microscope. Images were captured using TIA imaging software.

### **Neutrophils Recruitment Assay**

Neutrophil mobilization was elicited through injection of 300 nM f-Met-Leu-Phe (fMLP) chemoattractant into the otic cavity of 3 dpf larvae (Bernut et al., 2016) and counting the recruited neutrophils at the injection site using fluorescence microscopy.

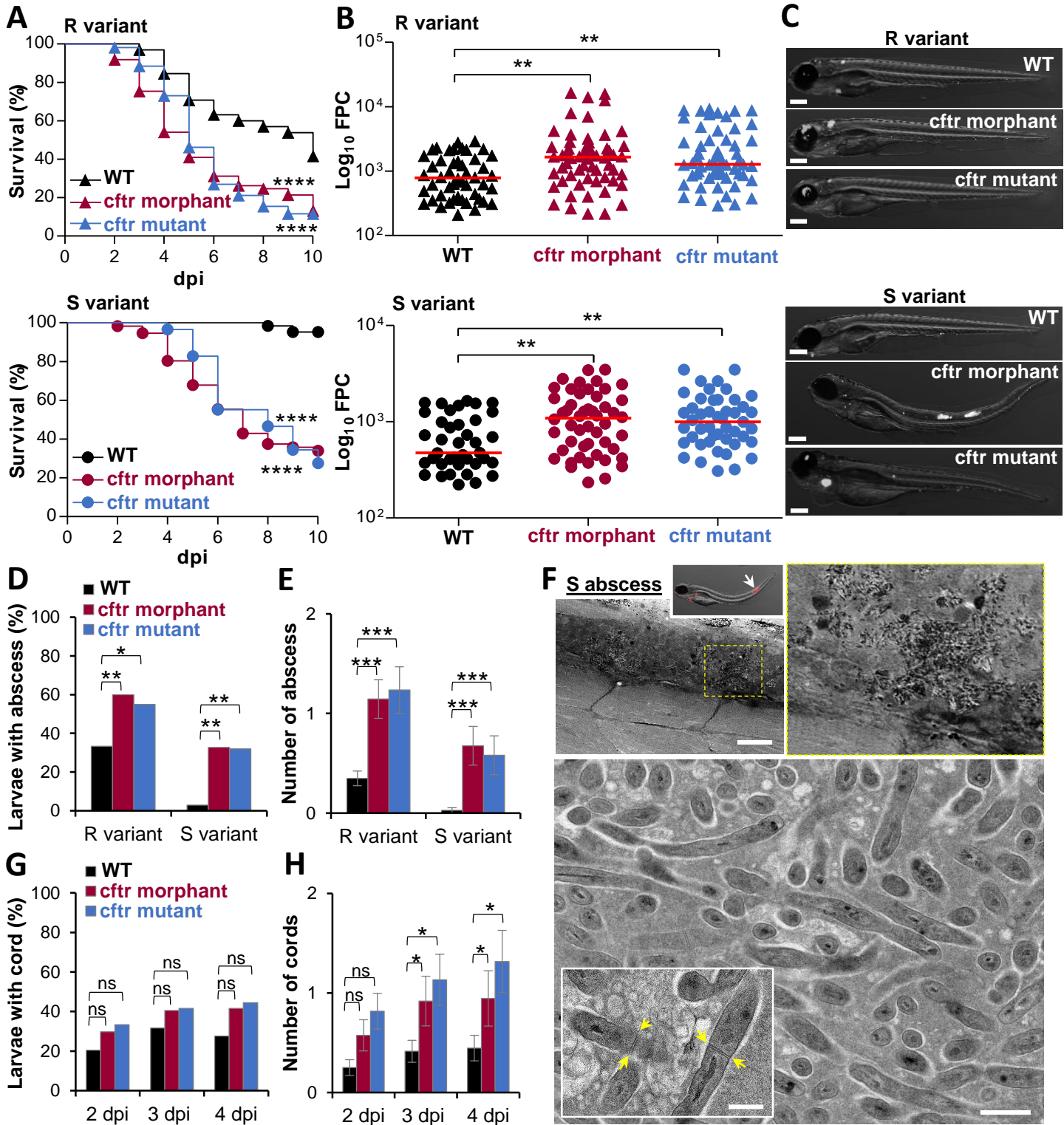
### **Cell Death and Reactive Oxygen Species Detection**

Leucocyte killing in ZF was detected using Acridine Orange (AO), as previously described (Bernut et al., 2014). Living embryos were soaked in 10 mg/ml AO in fish water for 30 min at 28.5°C, followed by two washes, then replaced at 28.5°C until observation and analyzed by confocal microscopy. For ROS detection, living larvae were soaked in 5 µM CellROX Deep Red Reagent in PBS for 30 min at 28.5°C, rinsed twice with PBS and directly prepared for confocal microscopy observations.

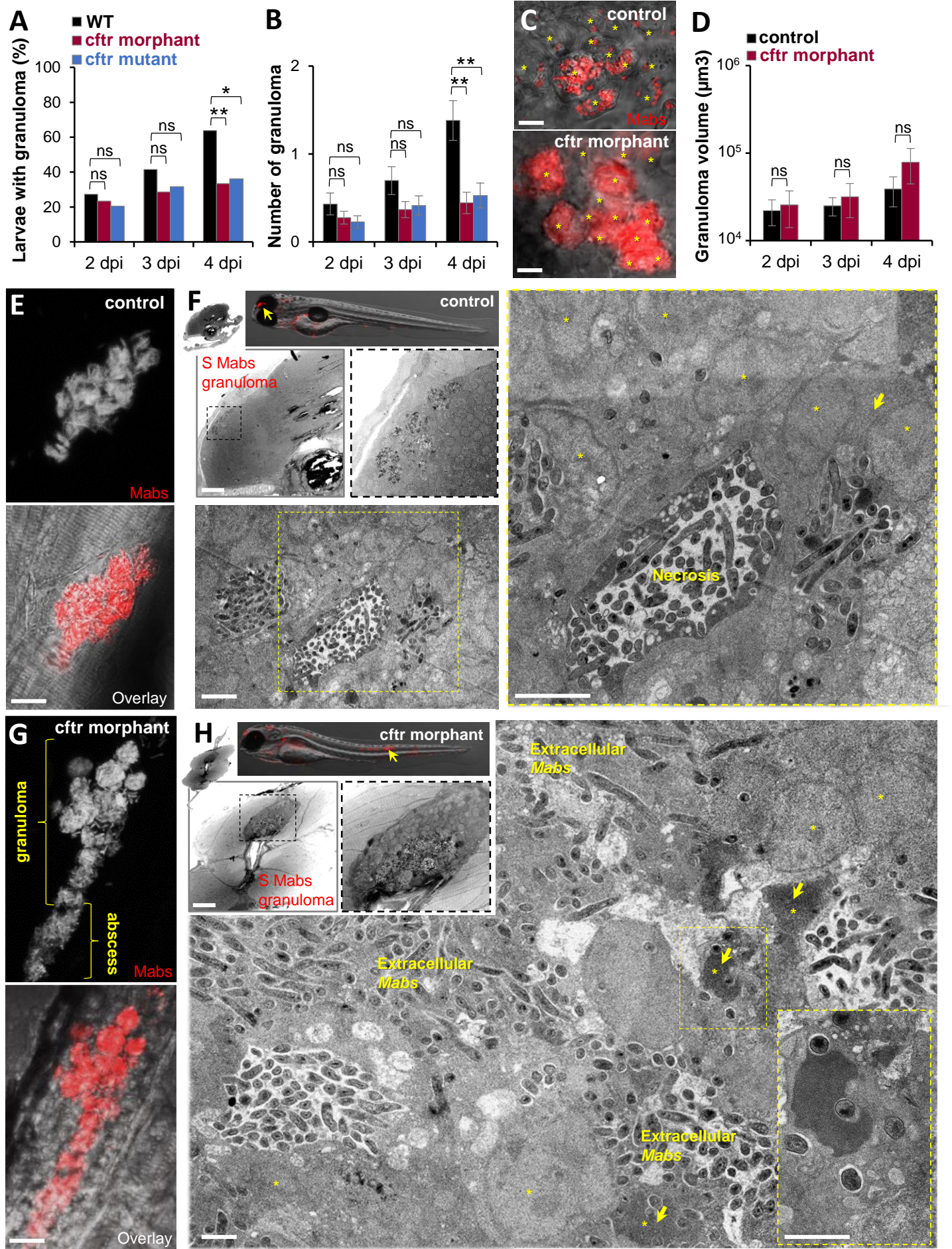
### **QUANTIFICATION AND STATISTICAL ANALYSIS**

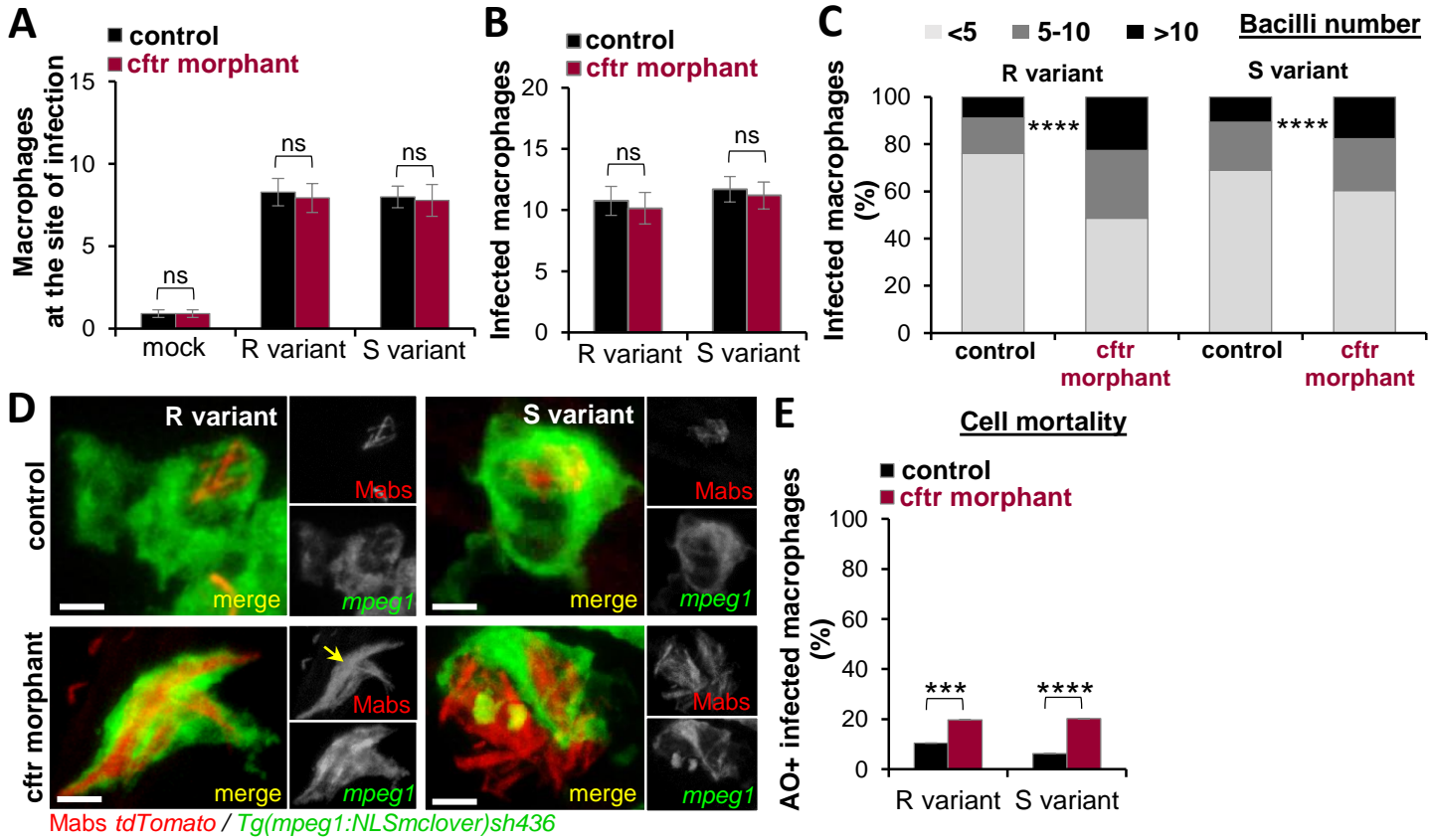
Statistical analysis (Prism 7.0; GraphPad Software or R 3.5.0) was performed using  $\chi^2$ Fisher's exact test, two-tailed unpaired Student's *t* test for comparisons between two groups and one-way or two-way ANOVA (with appropriate post-test adjustment) for other data. Mantel-Cox Log-rank test was used to compare survival curves. All data are plotted as average of two or three independent experiments. All error bars indicate standard errors of means (SEM). ns, not significant ( $p \geq 0.05$ ); \* $p < 0.05$ ; \*\* $p < 0.01$ ; \*\*\* $p < 0.001$ ; \*\*\*\* $p < 0.0001$



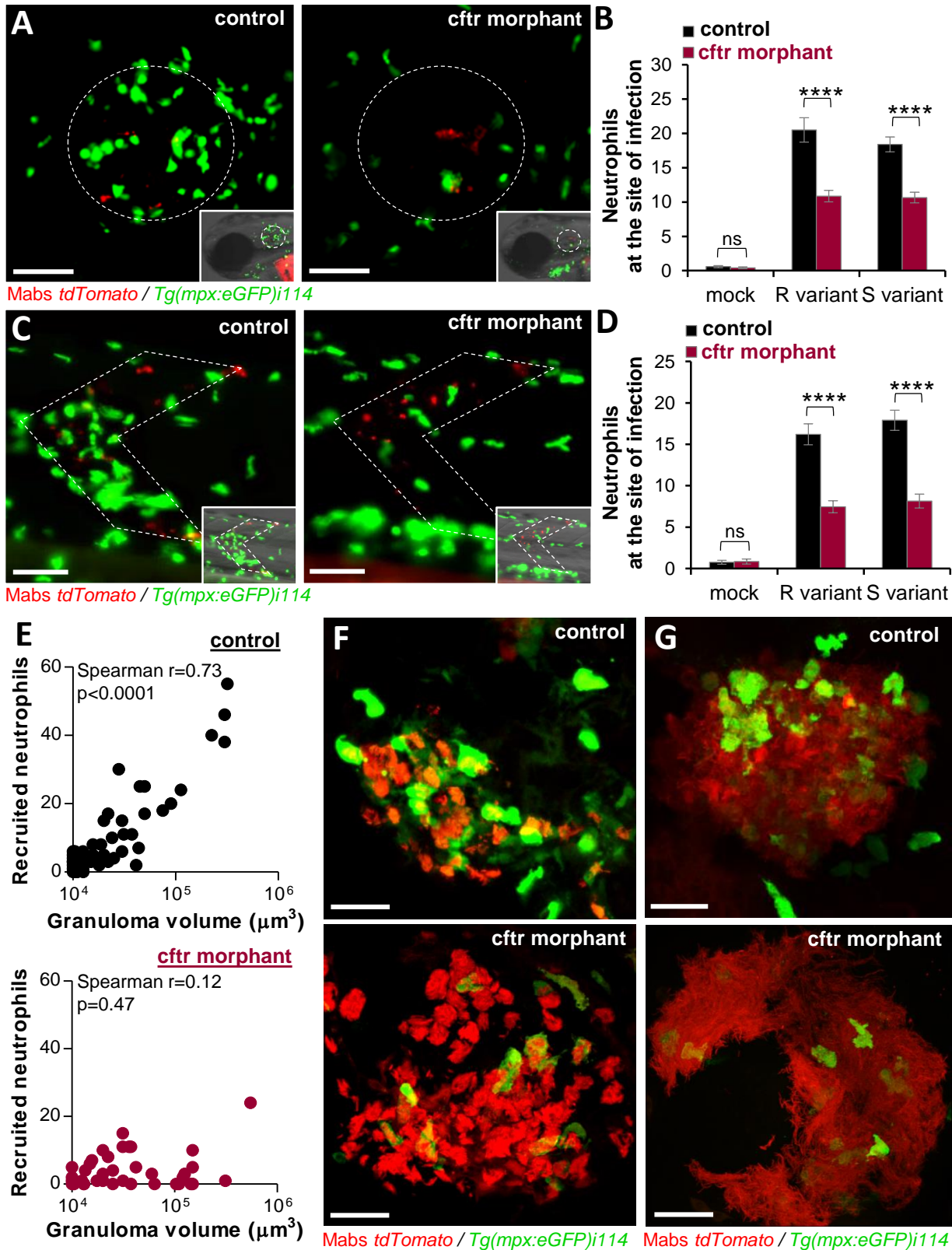


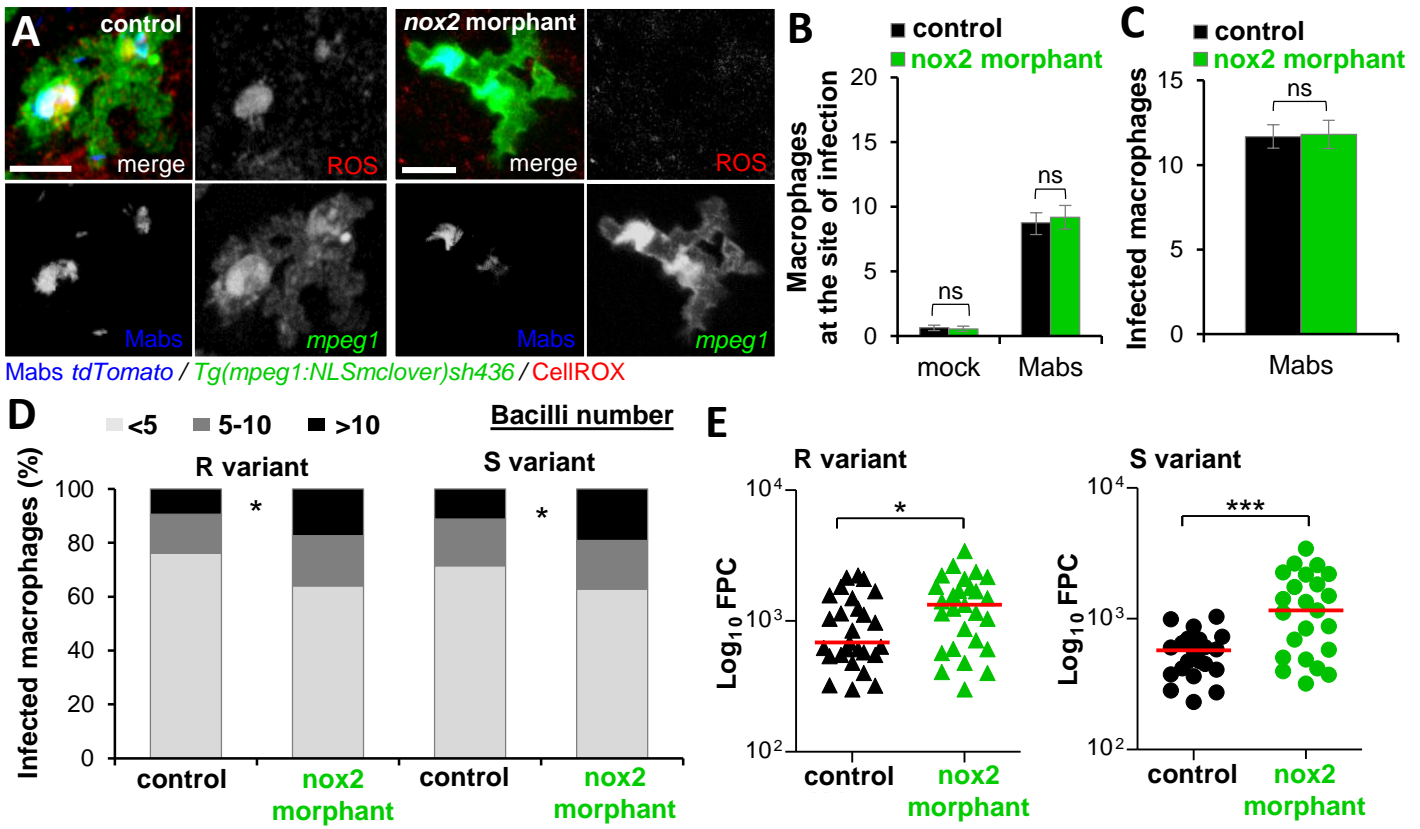


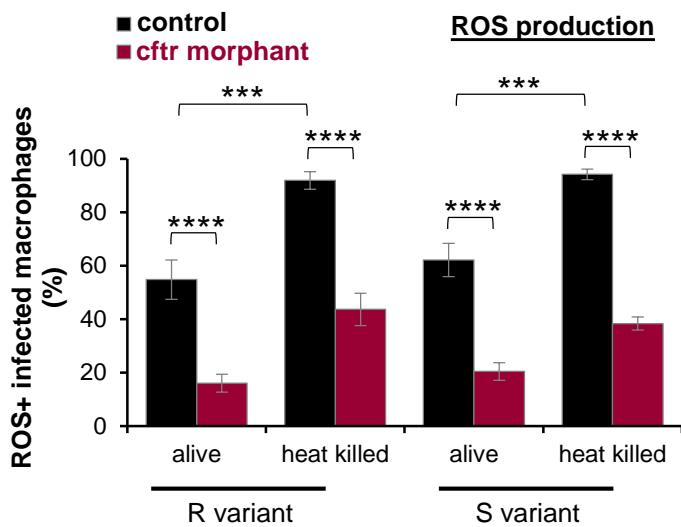
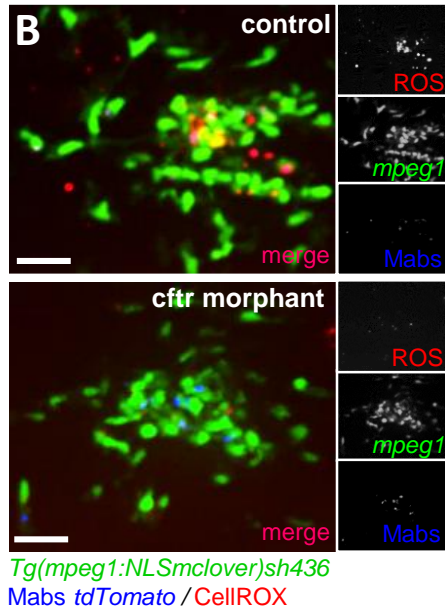
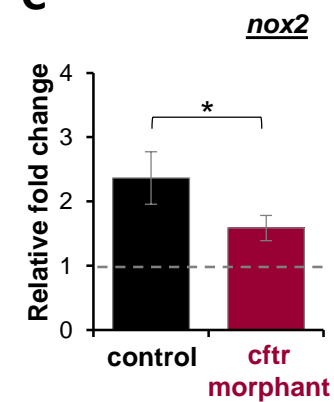
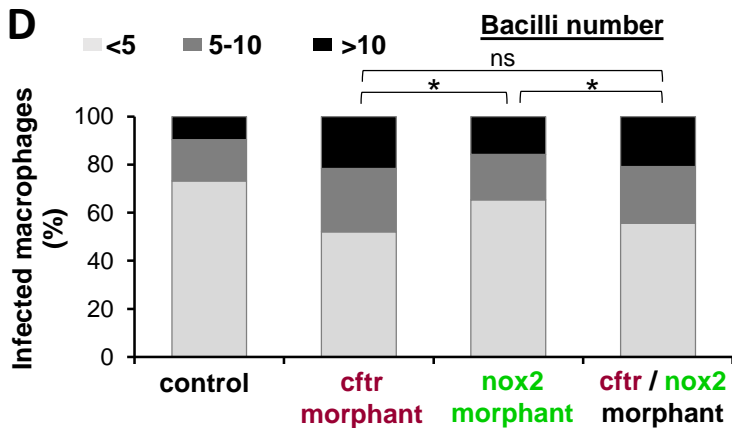


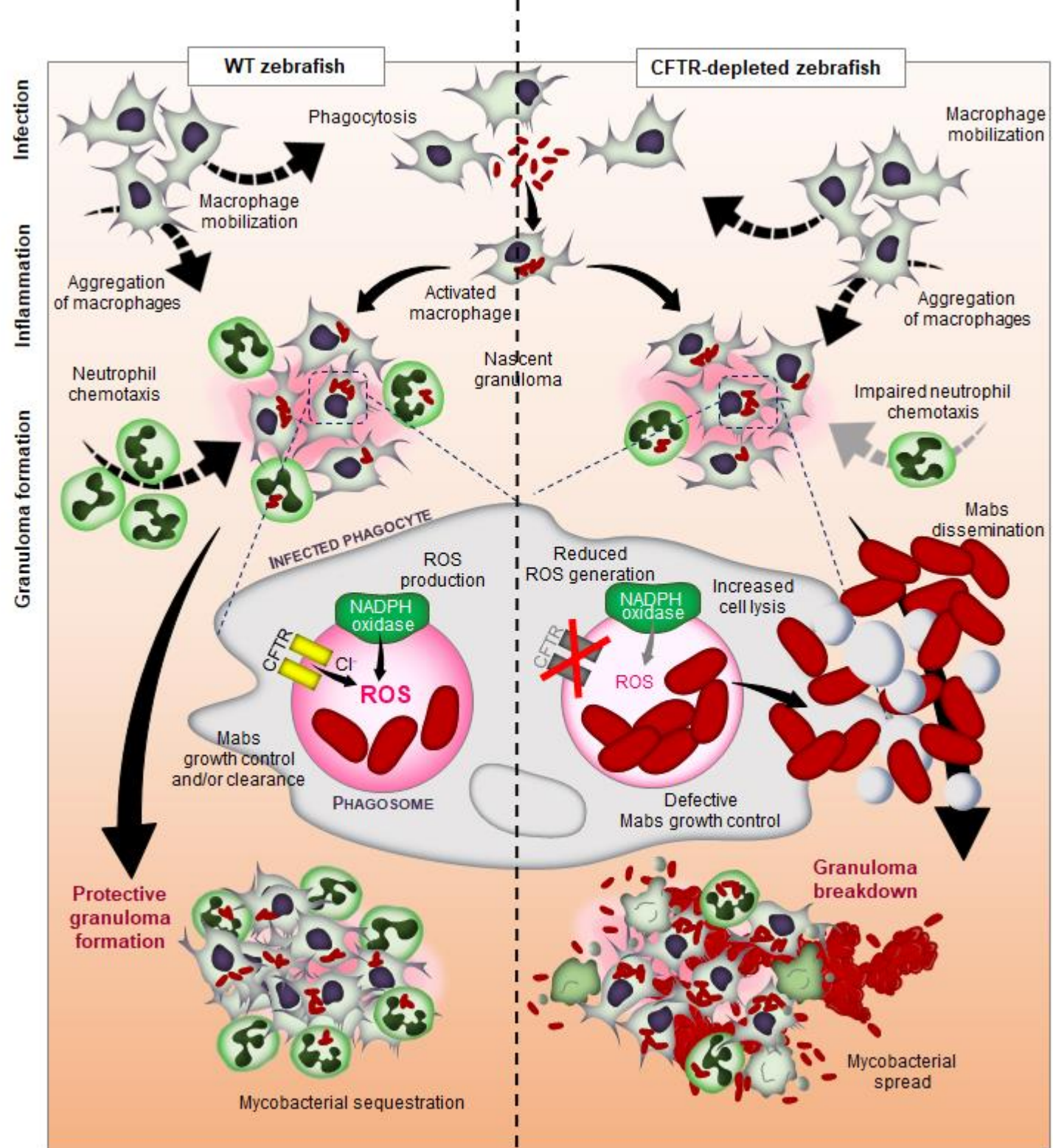








**A****B****C****D**





**WT-context**

**CF-context**

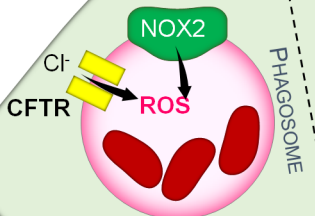
**Granuloma  
breakdown**

**Bacterial  
dissemination**

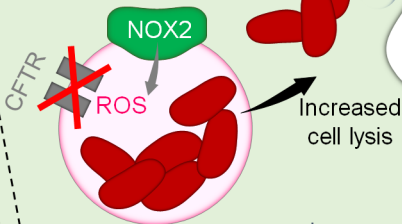
**Protective  
granuloma  
formation**

ROS  
production

Reduced ROS  
production



**Bacterial growth control  
and/or clearance**



**Defective bacterial  
growth control**

**Increased  
cell lysis**

**INFECTED  
PHAGOCYTE**