The origin of extracellular DNA in bacterial biofilm infections in vivo

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One sentence summary: eDNA is not observed within in vivo biofilms, but surrounding the biofilms. A PMN-derived layer of eDNA may provide a passive physical shield for the biofilm against antibiotics and phagocytes.

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ABSTRACT

Extracellular DNA (eDNA) plays an important role in both the aggregation of bacteria and in the interaction of the resulting biofilms with polymorphonuclear leukocytes (PMNs) during an inflammatory response. Here, transmission electron and confocal scanning laser microscopy were used to examine the interaction between biofilms of Pseudomonas aeruginosa and PMNs in a murine implant model and in lung tissue from chronically infected cystic fibrosis patients. PNA FISH, DNA staining, labeling of PMN DNA with a thymidine analogue and immunohistochemistry were applied to localize bacteria, eDNA, PMN-derived eDNA, PMN-derived histone H3 (H3), neutrophil elastase (NE) and citrullinated H3 (citH3). Host-derived eDNA was observed surrounding bacterial biofilms but not within the biofilms. H3 localized to the lining of biofilms while NE was found throughout biofilms. CitH3, a marker for neutrophil extracellular traps (NETs) was detected only sporadically indicating that most host-derived eDNA in vivo was not a result of NETosis. Together these observations show that, in these in vivo biofilm infections with P. aeruginosa, the majority of eDNA is found external to the biofilm and derives from the host.

Keywords: polymorphonuclear leukocyte; neutrophil extracellular traps; NETosis; histone; elastase

INTRODUCTION

Persistent bacterial infections present an increasing challenge to health care practitioners worldwide. For Pseudomonas aeruginosa bacterial aggregation shows consistency with biofilm formation in both the tolerance and resistance to both antibiotics and the host immune system (Alhede et al. 2011; Bjarnsholt et al. 2013). Most studies of bacteria are based on planktonic cultures consisting primarily of single cells grown in vitro where both aggregation and host factors are missing (Moser et al. 2017). To understand the underlying mechanism(s) causing persistence of chronic bacterial infections, it is important to investigate the
interplay between bacterial biofilms and the host immune system. A common denominator of both bacterial biofilms and cells of the innate host defense is extracellular DNA (eDNA). In vitro studies have shown eDNA, produced by the bacteria themselves, to be important primarily for initial biofilm formation as a major structural component of bacterial biofilms (Whitchurch et al. 2002) and as a stabilizer that contributes to increased tolerance of biofilms to antibiotics in vitro (Allesen-Holm et al. 2006; Izano et al. 2008; Chiang et al. 2013; Das, Sehar and Manefeld 2013).

The role of eDNA in vivo has still not been established, but during the last decade the use of animal model to study eDNA has been published (Jurcisek et al. 2007; Conover, Mishra and Deora 2011; Jung et al. 2017; Sovaelomandrosio et al. 2017). However, most of the studies only focus on the eDNA as a component originating from the bacteria and being a part of the biofilm, because this has been shown in vitro. Unfortunately, the host response and/or the in vivo environment is not taken into account. The common feature of chronic bacterial infections is persistent inflammation dominated by polymorphonuclear leukocytes (PMNs), which border bacterial biofilms but are unable to eradicate them (Bjarnsholt et al. 2009). This type of inflammation is classified as acute inflammation. Successful resolution of acute inflammation involves apoptosis and engulfment of demised PMNs by macrophages (Cox, Crossley and Xing 1995). However, during chronic bacterial infections, PMNs are thought to undergo necrosis, during which release of toxic compounds such as NE, oxidants and eDNA actually increase inflammation (Cox, Crossley and Xing 1995). Furthermore, components such as NE contribute to the destruction of tissue in chronic wounds and at sites of inflammation in the lungs of cystic fibrosis (CF) patients (Ohlsson and Olsson 1977; Yager and Nwomeh 1999; Voynow, Fischer and Zheng 2008; McCarty et al. 2012; Suleman 2016). PMNs actively secrete neutrophil elastase (NE) and histones attached to DNA, via a process termed NETosis (Steinberg and Grinstein 2007), both in vitro and in vivo in acute infections (Brinkmann et al. 2004; Yipp et al. 2012). This process is collectively referred to as neutrophil extracellular traps (NETs), which capture and kill bacteria. Since 2004, formation of NETs has been proposed as a function of neutrophils and other immune cells (Goldmann and Medina 2012) activated in vitro and in vivo in response to various bacterial components (Yipp et al. 2012; Kenny et al. 2017).

However, a recent review questioned whether NETs play a role in innate immunity and thereby trapping of bacteria (Sørensen and Borregaard 2016). Release of eDNA by NETosis has, to the best of our knowledge, never been demonstrated in association with inflammation caused by chronic bacterial infections, despite evidence of close contact between PMNs and bacteria.

At present, our knowledge about the interaction between in vivo biofilms and the host immune system during chronic infections is limited and we do not know the role or localization of eDNA in relation to the biofilms in vivo. Unsuccessful eradication of infectious biofilms despite accumulation of PMNs has not been explained fully. Thus, to obtain more detailed information about the interplay between bacterial biofilms and PMNs during chronic infection and to increase our understanding of the role of eDNA, we studied these phenomena for the first time, directly in a murine implant model of biofilm infection and directly in human samples of chronic-infected CF patients. By using transmission electron microscopy (TEM) we were able to study the interaction between PMNs and biofilm of Pseudomonas aeruginosa formed on a silicone implant inserted in the peritoneal cavity of mice. Furthermore, labeling the DNA of PMNs during their S-phase using the Click-iT technology in vivo and confocal scanning laser microscopy (CLSM), showed PMN-derived DNA did not localize with biofilms. To establish the localization of specific components originating from PMNs we used immunohistochemistry and found that PMN-derived histone H3 (H3) and NE co-localized with biofilms, but citrullinated H3 (citH3) and PMN-derived DNA did not. The results obtained using the mouse model could be correlated directly to chronic bacterial infections in human CF lungs.

RESULTS

TEM-based examination of the interaction between biofilms and PMNs in a murine implant model shows damaged PMNs

Previously, we used scanning electron microscopy (SEM) (van Gennip et al. 2012) to examine the interaction between immune cells and P. aeruginosa biofilms in a murine implant model and found a marked influx of PMNs toward the biofilms. We then used CSLM to identify PMNs according to their segmented lobular nucleus. Here, we used TEM to examine the in-depth additional detailed interaction between bacteria and PMNs, and to visualize the matrix of the biofilm. Implants were inspected at three different time points (6, 24, and 48 h post-insertion) to illustrate progression of infection and the immune response.

At 6 h post-insertion, we noted intact PMNs containing internalized bacteria within vacuoles, which indicates active phagocytosis (Fig. 1A). PMNs with internalized bacteria were confined to areas with a low bacterial density. Areas with a higher bacterial density decreased the presence of intact PMNs (Fig. 1B). At 24 h post-insertion, bacterial aggregation resulted in the formation of large biofilms lining the surface of the inside of the implants (Fig. 1C). PMNs frequently adjoined the bacteria. These PMNs were enlarged and had damaged cell membranes (Fig. 1C). Intact PMNs at this time point were rare. Matrix material surrounding the bacteria within the biofilm was evident (Fig. 1E–G). At 48 h post-insertion, the biofilm appeared denser and the PMNs were tightly interfaced with the bacteria (Fig. 1D). Matrix material was clearly visible (Fig. S1, Supporting Information).

In vitro interaction of P. aeruginosa and PMNs results in lysis of PMNs

When we examine the in vitro interaction between PMNs and aggregated P. aeruginosa microscopically, time series reveal that PMNs lyse resulting in release, expansion and dissolution of DNA (Movie 1 and Fig. S5, Supporting Information). The staining of the nuclear content, DNA, with propidium iodide (PI), increases over time as the membrane of the PMNs becomes damaged. At the end the DNA disappears (the red color from PI fades) (Movie 1 and Fig. S5, Supporting Information).

PMN-derived eDNA surrounds in vivo biofilms in a murine implant model

To examine eDNA in vivo, we stained silicone implants ex vivo from the murine implant model with SYTO9 (green) which binds to DNA together with the Click-iT® DNA labeling technology kit to determine the origin (i.e. murine or bacterial) of the eDNA in biofilms in vivo. This method utilizes a modified thymidine analogue, 5-ethynyl-2’-deoxyuridine (EdU), which is incorporated into DNA during active DNA synthesis in the S-phase of the eukaryotic cell cycle. Ex vivo, PMN DNA was detected by labeling with a fluorescent Alexa Fluor® dye (pink). Treatment of healthy mice showed the presence of Edu-labeled PMNs in
Figure 1. (A), Pseudomonas aeruginosa is internalized by PMNs in the murine implant model. Top row: TEM images showing intact and active PMNs containing internalized P. aeruginosa (arrows) 6 h post-insertion of a pre-coated silicone implant (Bar: 2 μm). Bottom row: internalized bacteria are magnified (Bar: 500 nm). Images represents sections obtained from two biological samples (two implants). (B–D), Interaction between PMNs and P. aeruginosa in the murine implant model. Pre-coated silicone implants were inserted into the peritoneal cavity of BALB/c mice. The interaction between PMNs and bacteria was imaged by TEM at 6 h (B), 24 h (C) and 48 h (D) post-insertion. In (B) black arrows points to bacteria and in (C) to a P. aeruginosa biofilm. Arrow heads: PMNs (Bar: 5 μm). Images represents sections obtained from two biological samples (two implants). (E–G), Matrix material surrounding P. aeruginosa on a silicone implant at 24 h post-insertion. TEM images showing matrix material surrounding P. aeruginosa bacteria at different magnifications. Matrix material (black arrows) can be seen between the bacteria. (E) Bar: 2 μm; (F) Bar: 1 μm, and (G) Bar: 200 nm. Images represents sections obtained from two biological samples (two implants).
The release of PMNs into the blood increases during an infection (Hansen, Karle and Valerius 1978), we decided to treat the mice two days pre-infection to increase the probability of EdU-labeled PMNs would reach the implant and biofilm post-infection. The percentage of EdU-labeled PMNs in the images presented in Figs 2–4 indicated 45% (165 pink/370 green), 42% (28 pink/66 green) and 89% (32 pink/36 green), respectively.

Long strings of eDNA (green) were clearly visible in the PMN accumulations with interacting bacteria (Fig. 2) and surrounding the biofilms (Fig. 3), but not as part of the biofilm itself (Figs 3 and 4).

As seen in Fig. 2, eDNA strings labeled with pink dye was clearly visible in close proximity to PMNs 24 h post-insertion of the implant. However, no pink eDNA was detected in areas occupied by bacterial biofilms (Figs 3 and 4) at 24 h or 48 h post-insertion, indicating that PMN-derived eDNA is not present internally within these structures.

eDNA surrounds in vivo biofilms in lung tissue of chronically infected CF patients

Previously, peptide nuclear acid (PNA) fluorescence in situ hybridization (FISH) and 4’,6-diamidino-2-phenylindole (DAPI) have been used to identify bacterial biofilms surrounded by PMNs in deparaffinated sections of CF lung tissue (Bjarnsholt et al. 2009). The PNA probes bind to bacterial ribosomal 16S RNA (Stender et al. 2002) and DAPI binds to the A-T regions of double-stranded DNA. Here, we used PNA FISH (red) in combination with DAPI (blue) to show that eDNA was localized external to the biofilm in CF lung tissue (Fig. 5A–C). The biofilm stained red by a Texas Red-labeled P. aeruginosa-specific PNA probe, and the nuclei of PMNs were clearly stained blue, as were the strings of DNA outside the biofilm (Fig. 5A). Furthermore, we used the DNA stain propidium iodide (PI), which stain all cells red due to the fixation of the tissue, whereas only strings of DNA were observed outside the biofilms.

Localization of histone H3 (H3), citrullinated H3 (citH3) and NE in a murine implant model

To localize components originating from murine PMNs during a bacterial biofilm infection, we stained deparaffinated sections of silicone implants isolated 24 h post-insertion in the peritoneal cavity of mice with antibodies specific for citH3 (Fig. 6), H3 (Fig. 7) or NE (Fig. 8). All of these components, especially citH3, have also been associated with NETs. Hence, we used citH3 as a marker for NETs (Neeli, Khan and Radic 2008; Wang et al. 2009; Wong et al. 2015; Barliya et al. 2017).

H3 localized to areas surrounding bacterial biofilms. Only weak staining was observed inside the biofilm; these findings are in agreement with those described above (i.e., no eDNA originating from PMNs was observed within biofilms). citH3 localized with some PMNs, but the lack of citH3 signifies that the eDNA is not generated by NETosis.

In contrast staining with specific antibodies against NE revealed distinct co-localization of biofilms and NE, indicating that NE is incorporated into biofilms (Fig. 8). Control images are presented in Fig. S4A. For specific identification of P. aeruginosa a PNA FISH image is presented in Fig. S4C (Supporting Information).
Figure 3. The *in vivo* biofilm lack PMN-derived DNA 24 h post-insertion in the murine implant model. PMNs in the murine implant model were labeled with Click-iTR®, in which a modified thymidine analogue, EdU (5-ethyl-2′-deoxyuridine), is incorporated into DNA during active DNA synthesis in murine immune cells in *in vivo*, and hence will label only DNA originating from murine PMNs. DNA is labeled *ex vivo* with Alexa Fluor® 647 (pink) and counterstained with SYTO9 (green) on an implant 24 h post-insertion in the peritoneal cavity. The number of PMNs stained with SYTO9 were estimated to 66 whereas the EdU-labeled (pink) PMNs were estimated to 28. Thereby 42% PMNs in the images were EdU labeled. In the PMN accumulations SYTO9 (green)-stained DNA strings (white arrows) were observed. EdU labeling was observed in PMNs (pink), but was absent from biofilms (double headed arrows), suggesting that PMNs are not a source of eDNA in biofilms. (A, D), merged images showing both EDU (pink) and SYTO9 (green) staining. (B, E), images showing only the SYTO9 staining. (C, F), images showing only the EDU staining. Red squares in A indicates magnified area in D–F and G–I. Images represents staining obtained from four biological samples (four implants).

Localization of H3, citH3 and NE in lung tissue of chronically infected CF patients

Co-localization of NE and H3 with *P. aeruginosa* biofilms in the murine implant model led us to examine deparaffinized lung sections from explanted CF patients with chronic bacterial infections. To verify function and correct binding of the three antibodies when using human PMNs, we stained *in vitro* phorbol myristate acetate (PMA) generated NETs from human PMNs. We were able to show co-localization of NE and citH3 with NETs, but not with H3 (Fig. S2, Supporting Information). The lack of H3 co-localization correlates well with the fact that NETosis results in a partial degradation of H3 and post-translational histone modifications and when using PMA as stimulation a total loss of full length H3 is observed (Papayannopoulos et al. 2010; Kenny et al. 2017). However, we did find co-localization of H3 with the nucleus of unstimulated PMNs, which indicates a functional antibody (Fig. S3, Supporting Information). NE also co-localized with the cytoplasm of unstimulated PMNs of deparaffinated sections (Fig. S3, Supporting Information).

In CF lungs the distribution of citH3, H3 and NE was similar to that observed in the murine implant model. CitH3 was observed in a few PMNs but did not co-localize with bacterial biofilms (Fig. 9A–C). H3 was observed at the periphery of biofilms but not co-localized with the biofilm (Fig. 9D–F). NE co-localized with bacteria within the biofilm (Fig. 9G–I). Control images are shown in Fig. S4B (Supporting Information). To estimate the co-localization of NE, H3 and citH3 with *in vivo* biofilms we used Manders’ co-localization coefficient as seen in Fig. 9J. A value of 1, means that the green pixels (SYTO9) are perfectly co-localized with the pink pixels (antibody). The co-localization coefficient confirmed what we visually observed, H3 localized outside of biofilms (*P* < 0.0001), citH3 sometimes to outside of biofilms (*P* < 0.0007) and NE localized primarily to biofilms (*P* < 0.25). There was not a significant difference between the outside and biofilm areas for NE. Some of the biofilms seemed to be located in a substance material, presumably alginate as previously shown (Bjarnsholt et al. 2009) due to mucoid *P. aeruginosa*. These biofilms did not co-localize with NE. Instead NE was co-localized with the surrounding PMNs.
in which a modified thymidine analogue, EdU (5-ethynyl-2'-deoxyuridine), is incorporated into DNA during active DNA synthesis in murine immune cells
32. Thereby 89% PMNs in the images were EdU labeled. Labeling was observed in PMNs (pink) but was absent from biofilms (double-headed arrows), suggesting that
hence will label only DNA originating from murine PMNs

**DISCUSSION**

Surprisingly, our visual findings show that, in vivo, eDNA is concentrated external to bacterial biofilms rather than inside as shown previously in vitro by us and others (Whitchurch et al. 2002; Alhede et al. 2011). In fact, the lack of SYTO9 and DAPI staining within the bacterial biofilms in chronic bacterial infections truly questions the significance of the, otherwise in vitro important, role of eDNA in biofilms in vivo as a scaffold of the biofilm. Instead it highlights the important and detrimental feature of host-derived eDNA as a protective shell surrounding the bacteria. This barrier of host eDNA surrounding a biofilm in vivo, which could both limit dissemination of bacteria and shield the biofilm from phagocytosis, may be manifested by necrotic lysis of PMNs rather than through NETosis. Thus, our results are in support of deposition of PMN content in a zone between the PMNs and the biofilm as recently demonstrated in experimental *P. aeruginosa* ocular biofilm infection where the deposition of NETosis derived material prevented dissemination (Thanabalasuriar et al. 2019). In our study, we found only scarce amount of host-derived eDNA and citH3 suggesting a modest contribution by NETosis to the material in the zone between PMNs and biofilm. However, dissemination of the bacteria may be prevented by the NE localized to the biofilm and therefore we propose other mechanisms than NETosis to contribute to the stalemate and chronicity of biofilm infection as illustrated in Fig. 10.

Using TEM we observed biofilm formation and swollen PMNs with damaged membranes, which is in agreement with our former SEM observations (van Gennip et al. 2012) showing PMNs failing to eradicate bacterial biofilms. The matrix material seen between the bacteria in the biofilm has not been fully characterized, but in vitro studies show the importance of eDNA for initial formation of young surface biofilms in vitro (Whitchurch et al. 2002); accordingly, it is believed to be an important structural component (Allesen-Holm et al. 2006; Izano et al. 2008). Furthermore, *P. aeruginosa* has been shown to be attached to eDNA in CF sputum (Walker et al. 2005) and incorporate external eDNA into the biofilm matrix in vitro (Chiang et al. 2013). PMNs were enlarged and had damaged cell membranes, presumably due to exposure to the bacterial virulence factor, rhamnolipid, which is detrimental to PMNs (Alhede et al. 2009; van Gennip et al. 2009). It has previously been shown that *P. aeruginosa* produce rhamnolipids which cause PMNs to swell and burst, thereby releasing their cytoplasmic and nuclear content into the surrounding area (Jensen et al. 2007). In an unpublished study, we observed that when PMNs were exposed to aggregates of *P. aeruginosa* in vitro approx. 60% PMNs were killed within 8 h (unpublished Alhede et al.). The cellular components including eDNA from necrotic PMNs are believed to serve as a biological matrix for biofilm formation and a protective shield within the biofilm increasing tolerance to treatment to tobramycin in vitro (Walker et al. 2005; Chiang et al. 2013). This was confirmed in a mouse model where a rhlA mutant (unable to excrete rhamnolipid) did not cause any PMN lysis and no release of cytoplasmic and nuclear content, thus consistent with a mechanism of lysis by the biosurfactant rather than a programmed NET pathway for eDNA release (van Gennip et al. 2009).

The lack of strings in in vivo biofilms indicate that unlike in vitro biofilms, eDNA is not a direct part of the biofilm. Visually eDNA from *P. aeruginosa* has been shown to form a net both in

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**Figure 4.** The in vivo biofilm lack PMN-derived DNA 48 h post-insertion in the murine implant model. PMNs in the murine implant model were labeled with Click-iTR®, in which a modified thymidine analogue, EdU (5-ethynyl-2'-deoxyuridine), is incorporated into DNA during active DNA synthesis in murine immune cells in vivo, and hence will label only DNA originating from murine PMNs. DNA is labeled ex vivo with Alexa Fluor® 647 (pink) and counterstained with SYTO9 (green) on an implant 48 h post-insertion in the peritoneal cavity. The number of PMNs stained with SYTO9 were estimated to 36 whereas the EdU labeled (pink) PMNs were estimated to 32. Thereby 89% PMNs in the images were EdU labeled. Labeling was observed in PMNs (pink) but was absent from biofilms (double-headed arrows), suggesting that PMNs are not a source of eDNA in biofilms (A, D). Merged images showing both EDU (pink) and SYTO9 (green) staining (B, E). Images showing only the SYTO9 staining (C, F), images showing only the EDU staining. Red square in A indicates magnified area in D-F. Images represents staining obtained from four biological samples (four implants).
a stationary grown biofilm and in the flow cell biofilm (Allesen-Holm et al. 2006; Alhede et al. 2011). Furthermore, eDNA from P. aeruginosa is believed to be double-stranded DNA (Allesen-Holm et al. 2006) and since DAPI bind to double-stranded DNA the lack of DAPI staining signifies the lack of eDNA in the in vivo biofilm. Also, the measurement of double-stranded DNA has been shown to decrease when P. aeruginosa change from the planktonic state to the aggregating state (Das et al. 2014). However, the eDNA of a plate grown colony of P. aeruginosa have been shown to be 5 μm long (Chiba et al. 2015) which mean that visually the eDNA fragment would be almost as small as a single P. aeruginosa bacterium. Visually this does not correlate to our in vivo and others in vitro findings. However, we might have to rethink the need for a structural matrix depending on the environment of the biofilm outside the laboratory. Maybe when the biofilm is exposed to high forces such as fluid it needs to establish a solid structure, but in the CF lung another type of structure is established within the mucus to withstand the immune cells. Accordingly, this indicates that neither bacterial nor PMN-derived eDNA is incorporated into biofilms in vivo, but eDNA from necrotic PMNs is more likely accumulating outside the biofilms.

Since we only found host-derived eDNA outside infectious biofilms in vivo, we speculated whether in vivo biofilm matrix structures comprised other components originating from PMNs (e.g. H3, citH3 or NE). PMNs undergoing apoptosis, necrotic cell death or NETosis release different histones (i.e. H3 during necrotic cell death and citH3 during NETosis), either into the surrounding area or attached to NETs (Brinkmann et al. 2004). Release of these extracellular histones promotes secretion of pro-inflammatory cytokines and recruitment of immune cells to sites of infection (Westman et al. 2015). Histones bind to lipopolysaccharide (LPS) derived from P. aeruginosa, which neutralizes the positive charge of the histone molecule (Rose-Martel and Hincke 2014); with the outcome of antimicrobial activity (Rose-Martel et al. 2017). The fact that histones are antimicrobial (Hirsch 1958; Brinkmann et al. 2004), might not be the case in this study. Since H3 is rich in arginine it may rather play a role in the arginine pathway in P. aeruginosa, which breaks down arginine to ornithine via the arginine deiminase pathway to sustain growth under anoxic conditions. Ornithine is a carbon source used to maintain growth during periods in which nitrate is either limited or lacking (Vander Wauven et al. 1984; Stalon et al. 1987; Eschbach et al. 2004). Arginine has also been shown to contribute to biofilm formation (Bernier et al. 2011).

Here, we showed that H3 is mostly present along the periphery of bacterial biofilms, but not within the biofilms. This could indicate that H3 is still bound to eDNA. By contrast, citH3 did not localize with biofilms in either the mouse model or CF lung samples. CitH3 is a post-translational modification of H3. Conversion of arginine to citrulline is accomplished by peptidylarginine deiminase 4 (PAD4) and is a prerequisite for in vitro generated NETosis since the modification of arginine residues to citrulline changes the charge of the histones, leading to massive chromatin de-condensation, NETs (Wang et al. 2009; Lewis et al.
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Figure 6. citH3 antibody staining of the murine implant model. CSLM images of deparaffinized sections of silicone implants from the murine implant model 24 h post-insertion. The sections were stained with primary antibodies specific for citrullinated histone H3 (citH3). The secondary antibody was conjugated to Alexa Fluor 647 (pink). SYTO9 (green) was used as a counterstain. SYTO9 stains DNA in bacteria and eukaryotic cells. (A, D), Merged images of both the antibody (pink) and SYTO9 (green) staining. (B, E), Are only the SYTO9 staining. (C, F), are only the antibody staining. The images represent two implants. The letter B indicates biofilm. Images represents staining of 2-4 sections obtained from two biological samples (two implants).

NETs are highly de-condensed chromatin structures that comprise mainly DNA, although histones and antimicrobial granular substances such as NE are also present (Brinkmann et al. 2004). Histone citrullination in vitro depends on stimulation of PMNs (Neeli and Radic 2013; König and Andrade 2016); indeed, an increase in histone citrullination is associated with chromatin de-condensation during NETs formation and LPS-induced early responses to inflammatory stimuli of PMNs (Neeli, Khan and Radic 2008; Wang et al. 2009). However, one widely used stimulus, PMA, has resulted in conflicting results in the literature and has been shown to induce NETs with and without co-localized citH3 and H3 (Brinkmann et al. 2004; König and Andrade 2016; Neeli and Radic 2016). In addition citrullination of H3 is strongly reduced when PMNs are stimulated during anoxic conditions (Lopes et al. 2018), which is present in the endobronchial secretion in infected CF lungs (Worlitzsch et al. 2002; Kolpen et al. 2014).

The lack of host-derived eDNA and citH3 associated with bacterial biofilms in the present study demonstrates that NETosis probably does not occur in chronic biofilm infections. Release of DNA under these conditions is not an active process as in NETosis; rather, it is more likely due to lysis of PMNs (Jensen et al. 2007) which correlates to the presence of H3 and absence of citH3. In addition, our findings that NETosis seemingly plays no role in chronic P. aeruginosa infections correlates to the depletion of molecular oxygen at the site of chronic biofilm infections (Kolpen et al. 2010), thereby preventing NADPH oxidase from generating the reactive oxygen species (ROS) needed to release NETs (Fuchs et al. 2007). Thus, PAD4-dependent NET formation might not be responsible for DNA release in CF airways and we speculate whether citH3 may serve as a biomarker for discriminating chronic from acute infections. However, NET generation independent of ROS generated by NADPH oxidase has been shown in vivo (Kolaczkowska et al. 2015) therefore further work is needed to confirm this. Nevertheless, eDNA in sputum samples from CF patients have been shown to originate from PMNs (Lethem et al. 1990) with NETs as the major component (Manzenreiter et al. 2012). However, since NETs are antimicrobial this should benefit the CF patients since this should lead to eradication of the present bacteria. On the contrary eDNA has been shown to increase the viscosity of the mucus and bind antibiotics such as tobramycin (Potter et al. 1965; Ramphal et al. 1988). Furthermore, by treating the CF patients with DNase an improved lung function in CF patients and an increased effect of antibiotics have been reported (Shak et al. 1990; Hunt et al. 1995).

We did observe NE within biofilms, although this may have been derived from a necrotic cell death of PMNs external to the biofilm, followed by passive diffusion into the biofilm structure. NE is stored in primary (azurophilic) granules and released upon degranulation, phagocytosis and cell death. NE cleaves a wide variety of proteins, including bacterial virulence factors.
Figure 7. H3 antibody staining of the murine implant model. CSLM images of deparaffinized sections of silicone implants from the murine implant model 24 h post-insertion. The sections were stained with primary antibodies specific for histone H3 (H3). The secondary antibody was conjugated to Alexa Fluor 647 (pink). SYTO9 (green) was used as a counterstain. SYTO9 stains DNA in bacteria and eukaryotic cells. (A, D), Merged images of both the antibody (pink) and SYTO9 (green) staining. (B, E), is only the SYTO9 staining. (C, F), is only the antibody staining. The images represent two implants. The letter B indicates biofilm. Images represents staining of 2–4 sections obtained from two biological samples (two implants).

(Weinrauch et al. 2002). The purpose of NE is to degrade phagocytosed proteins; however, in CF patients NE damages the airways (Janoff et al. 1979). Furthermore, NE is an important component of NETs (Brinkmann et al. 2004). However, since NE was found within the biofilms and eDNA was not supports the lack of NETs, since in these structures NE is bound to NETs. Furthermore, unlike in acute infections caused by Gram-negative bacteria P. aeruginosa is not killed by NE following aggregation and biofilm formation (Belaauou et al. 1998). The absence of NE-mediated killing is likely attributed to production of the protease inhibitor ecotin by bacteria (Eggers et al. 2004). In addition, NET formation induced by P. aeruginosa depends strongly on flagella motility (Floyd et al. 2016), which may further explain the absence of NETs in CF lungs in which the flagella of P. aeruginosa are down-regulated (Ahmed et al. 1993) and in which flagellin, which is the major component of flagella, may be cleaved by NE (Lopez-Boado et al. 2004). Since a decrease and/or absence of flagella motility is associated with formation of bacterial biofilms (Chaban, Hughes and Beeby 2015), a lack of motile flagella may lead to a lack of detectable NETs.

Limitations: Our study applies different microscopic techniques for visual inspection of different model systems and patient samples. An image-based study as this has limitations as to showing representing images of the investigated question. However, as in this case appropriate experimental setups are not available and since the imaging technique continual improves, we are now able to study small details such as DNA. We think that the visual information obtained significantly contributes to new information on the in vivo setting of chronic bacterial infections. The in vitro models available also have limitations when studying the interaction between immune cells and biofilms since most often the immune cells are isolated from their natural environment and the bacteria are grown in a media not resembling the in vivo environment.

In conclusion, this study is the first to examine the direct distribution of host and bacterial eDNA during chronic bacterial infections in vivo. Unlike after stimulation in vitro, stimulation of PMNs by bacterial biofilms during chronic infections in vivo does not seem to trigger active release of NETs; however, necrotic PMNs do release eDNA, histone H3 and antibacterial enzymes such as NE.

We did not find evidence of eDNA being part of in vivo biofilms, instead the PMN-derived layer of eDNA, which constitutes a secondary matrix, may provide a passive physical shield for the biofilm against cationic antibiotics such as tobramycin and additional phagocytes. A similar spatial arrangement has recently been described in a murine model of P. aeruginosa infection of the cornea in which the bacterial biofilm and neutrophil layers are separated by a ‘dead zone’ rich in eDNA and NE (Thanabalasuriar et al. 2019).

Thus, we believe that eDNA released by PMNs due to a necrotic cell death contribute to the protection afforded to bacteria by biofilms (Fig. 10). These novel findings shed new light on the origin and role of eDNA in in vivo biofilms and unexpectedly
question the role of extracellular host DNA during inflammation associated with chronic bacterial infections and of NET activation.

MATERIALS AND METHODS

Murine implant model

All experiments were performed using a wild-type (WT) P. aeruginosa strain obtained from Professor Barbara Iglewski (University of Rochester Medical Center, NY, USA). The strain is QS proficient, except for the reduced production of C4-HSL previously noted for this P. aeruginosa variant (Köhler et al. 2001). Bacteria obtained from freezer stocks were plated onto blue agar plates (State Serum Institute, Denmark) and incubated overnight at 37°C. Blue agar plates are used to select Gram-negative bacilli (Høiby 1974). One colony was used to inoculate overnight cultures grown in Luria-Bertani (LB) medium at 37°C with shaking at 180 rpm. Female BALB/c mice (6 weeks-of-age) were purchased from Taconic M&B A/S (Ry, Denmark) and maintained on standard mouse chow and water ad libitum for 2 weeks before challenge. The murine implant model was generated as described previously (Bjarnsholt et al. 2010; van Gennip et al. 2012), with some modifications. Briefly, the silicon tubes (Ole Dich Instrumentmakers Aps. Silicone Pumpselange 0 Shore A (ID: 4.0 mm, OD: 6.0 mm, wall: 1.0 mm) were cut to a length of 4 mm, and the bacterial pellet from a centrifuged overnight culture was resuspended in 0.9% NaCl (OD600nm = 0.1). Mice were anesthetized by subcutaneous (s.c.) injection of hypnorm/midazolam (Roche) [hypnorm (0.315 mg ml⁻¹ fentanyl citrate; 10 mg ml⁻¹ fluanisone); midazolam (5 mg ml⁻¹)] and sterile water; 1:1:2 v/v. The pre-coated silicone implants were inserted in the peritoneal cavity of the mice. The mice were treated with bupivacaine and Temgesic® as post-operative pain relief. Experiments were terminated with euthanization of the mice with intraperitoneal injection of pentobarbital (DAK; 10.0 ml kg⁻¹ body weight).

Preparation of samples for TEM

The silicone implants were fixed in 2% glutaraldehyde in 0.05 M sodium phosphate buffer pH 7.4. Growing biofilms on a silicone tube presents a challenge with respect to sample preparation prior to the different microscopy techniques. First, a traditional preparation method was used for TEM to get an indication of the interaction between PMNs and the biofilm. However, some of the steps involved in the embedding process were shortened. Samples were observed with a CM 100 TEM (Philips, the Netherlands) operated at 80 kV. Imaged were recorded with a side-mounted Olympus Veleta camera (resolution, 2048 × 2048 pixels (2 K × 2 K)) and the iTEM (Olympus, Germany) software package.
Figure 9. Immunostaining of CF lung tissue. CSLM images of deparaffinized sections of CF lung tissue. The sections were stained with primary antibodies specific for citrullinated histone H3 (citH3) (A-C), Histone H3 (D-F) or NE (G-I). The secondary antibody was conjugated to Alexa Fluor 647 (pink). SYTO9 (green) was used as a counterstain. SYTO9 stains DNA in bacteria and eukaryotic cells. (A, D, G), Merged images of both the antibody (pink) and SYTO9 (green) staining. (B, E, H), Are only the SYTO9 staining. (C, F, I), Are only the antibody staining. The images represent staining of 2–4 sections from three different CF lungs. B indicates biofilm. J: Co-localization of biofilm with antibodies in CF lungs shown as Manders’ co-localization coefficient. ‘Outside biofilm’ are areas containing PMNs and ‘biofilms’ are ROI defined biofilms. Sample size (n): NE outside biofilms (16), NE biofilms (20), H3 outside biofilms (18), H3 biofilms (17), citH3 outside biofilms (15), citH3 biofilms (18). There was significant difference between H3 outside biofilms and H3 biofilms (P < 0.0001) and citH3 outside biofilms and biofilms (P < 0.0007). No significant difference was found between NE outside biofilms and biofilms (P < 0.25). P < 0.05 was considered significant.
In vitro experiments with PMNs and P. aeruginosa

Wild-type P. aeruginosa (strain PAO1) was obtained from the Pseudomonas Genetic Stock Center (https://www.pseudomonas.med.ecu.edu; strain PA0001). Bacteria were tagged with a stable green fluorescent protein (GFP) constitutively expressed by plasmid pMRP9 (Davies et al. 1998). PAO1 cultures were grown for 24 h at 37°C in 12 well microtiter plates containing LB medium. Human blood was collected from healthy volunteers with the approval of the Danish Scientific Ethical Board (H-3-2011-117). PMNs were isolated as described (Bjarnsholt et al. 2005). Isolated PMNs were resuspended in Krebs ringer buffer containing 10 mM glucose at 37°C (final density, 2.5 x 10⁷ PMNs/ml). Propidium iodide (2.5 µg/ml) was used as an indicator of cell death. PAO1 (350 µl) and PMNs (50 µl) were mixed in a 12 well microtiter plate and observed under a confocal microscope for 35 min.

Click-iTTM

Mice were treated with EdU (Click-iTTM EdU Alexa Fluor™ 647 Imaging Kit) (C10640, ThermoFisher) for 2 days pre-infection and until termination of the experiment. Each mouse was injected i.p. twice a day with 25 mg/g EdU dissolved in 0.9% NaCl (Zeng et al. 2010). Control mice received 0.9% NaCl without EdU. Ex vivo implants were fluorescently labeled using the Click-iTT kit, according to the manufacturer’s instructions, and stained with SYTO9 (S34934, ThermoFisher) prior to observation under a microscope.

The percentage of SYTO9 stained PMN nuclei and EdU labeled cells were estimated using the Measuring PRO add on to Imaris 8.5 (Bitplane, Switzerland). Individual nuclei were identified by applying isometric surfaces and separated with the build in object separator as spheres with a 5–10 µm diameter in green (SYTO9) spectra. Similarly, antibody stained cells were separated and counted as ~10 µm diameter spheres in pink (EdU) spectra. The results were returned as simple numeric counts.

In vitro generated NETs

In vitro NETs were generated according to Vong et al. (Vong, Sherman and Glogauer 2013) and Brinkmann et al. (Brinkmann et al. 2012) with modifications. PMNs were isolated according to Hu et al. (Hu 2012) using polymorphprep with modifications. Isolated PMNs were suspended in RPMI 1640 with glutamine and were seated in an ibidi slide coated with poly-L-lysine (80 604, ibidi) for 30 min before stimulation with PMA for 3 h. The PMNs were incubated at 37 degrees and 5% CO₂. PMNs was washed with PBS and fixed in 4% PFA pH 7.2 for 15 min, washed in 0.025% Triton X-100 twice before proceeding with antibody staining as described below for antibody staining of deparaffinized tissue sections. Before imaging the channels were filled with Ibidi mounting medium (50001, ibidi).

Ex vivo paraffin samples

Samples from the mouse model, the lungs of CF patients and unactivated human PMNs were fixed in 4% formalin and paraffin-embedded according to standard protocols. Sections (4 µm thick) were cut using a standard microtome, fixed on glass slides and stored at 4°C until required.

Lung tissue from chronically infected CF patients

Explanted lungs tissue from three contemporary intensively treated P. aeruginosa infected CF patients were examined (Bjarnsholt et al. 2009).

PNA FISH

PNA FISH staining was performed as previously described (Kirketerp-Møller et al. 2008; Bjarnsholt et al. 2009). Texas Red-labeled P. aeruginosa-specific PNA probe both in hybridization solution (AdvanDx, Inc., Woburn, MA), was added to each section and hybridized in a PNA FISH workstation covered by a lid at 55°C for 90 min. The slides were washed for 30 min at 55°C in wash solution (AdvanDx) and counterstained with DAPI (D1306, ThermoFisher).

Antibody staining of deparaffinated tissue sections

Sections were deparaffinized in 99.9% xylene (twice for 5 min each time) and subsequently hydrolyzed in 99.9% ethanol (twice for 5 min each time), followed by 96.5% ethanol (twice for 3 min each time). Finally, sections were rinsed in water (twice for 5 min each time). Heat-induced antigen retrieval was performed in citrate buffer, pH 6.0 (Sigma C9999) for 15 min in a domestic microwave oven. For unstimulated human PMNs a permabilization step was included using 0.1% Triton X-100 for 20 min. Sections were washed twice (5 min each time) in 0.025% Triton X-100, drained and blocked for 2 h at room temp in 10% goat serum in 1% BSA/PBS. Samples were drained and incubated for 16–18 h at 4°C with a primary antibody (Abcam Cat# ab1791, RRID: AB_302613, Abcam Cat# ab5103, RRID: AB_304752 or Abcam Cat# ab21595, RRID: AB_446409), diluted 1:200 in 2% goat serum/1% BSA in PBS). Negative controls were isotype IgG (Abcam Cat# ab27478, RRID: AB_2616600) or absence of primary antibody. Samples were washed and incubated in 0.025% Triton X-100 (twice for 5 min each) with gentle agitation. Samples were then incubated (1 h at room temp in the dark) with a secondary antibody (Abcam Cat# ab150083, RRID: AB_2714032) diluted 1:1000 in 1% BSA/PBS. Finally, samples were washed three times in PBS (5 min each time before imaging).
time) and counterstained with SYTO9 (S34934, ThermoFisher). ProLong™ gold antifade mountant (Thermofisher, P36934) was applied before adding the coverslip.

Co-localization coefficient

Co-localization refers to the geometric co-distribution of two fluorescent labels or color channels. Co-localization was acquired in Zen (version 2.1 SP3 FP2 (black)) as described by Zeiss ’Acquiring and analyzing data for co-localization experiments in AIM or Zen software’. The Manders’ co-localization coefficient (Manders, Verbeek and Aten 1993) for SYTO9 (green channel) was used to show the co-localization of the antibody (pink channel) using 2D images. Perfect co-localization will give the value 1. The crosshair was set to 1003 for the green channel and 2500 for the pink channel based on single stained deparaffinated sections according to the manual by Zeiss. Sections from three different patients were used. Biofilm and PMN areas were selected by region of interest (ROIs) in each deparaffinated section. All biofilm areas in one image were included. GraphPad Prism was used for statistical analysis using unpaired t-test. P < 0.05 was considered significant.

Confocal scanning laser microscopy

All observations and image acquisition were performed using a confocal scanning laser microscope (LSM510, LSM 710 or LSM 880; Carl Zeiss GmbH, Germany). Images were obtained using 40 × 63 × 100 × oil objective lenses. Images were scanned at 405 nm (blue), 488 nm (green) and 633 nm (far red). Images of the biofilms were generated using Imaris software including co-localizations according to the manual by Zeiss. Sections from three different patients were used. Biofilm and PMN areas were selected by region of interest (ROIs) in each deparaffinated section. All biofilm areas in one image were included. GraphPad Prism was used for statistical analysis using unpaired t-test. P < 0.05 was considered significant.

SUPPLEMENTARY DATA

Supplementary data are available at FEMSPD online.

ETHICS STATEMENT

All animal studies were carried out in accordance with the European convention and Directive for the Protection of Vertebrate Animals used for Experimental and Other Scientific Purposes, and with Danish law on animal experimentation. All experiments were authorized and approved by the National Animal Ethics Committee, Denmark (The Animal Experiments Inspectorate, dyreforosgisticslyset.dk; permit numbers: 2012–15–2934–00677 and 2014–15–0021–00259).

Human blood was collected from healthy volunteers with the approval of the Danish Scientific Ethical Board (H-3-2011-117). All donors provided written informed consent.

Infected tissue was collected following approval by the Danish Scientific Ethical Board (KF465 01278432). The samples used were collected with written informed consent for a previous study [9], and only from adults. The samples were anonymized for this study.

AUTHOR CONTRIBUTIONS

M.A. conducted the mouse experiments and antibody staining experiments and wrote and edited the manuscript. M.A. undertook microscopic analysis of Click-It and edited the manuscript. K.Q. assisted with TEM preparation and image recording and edited the manuscript. PØJ. performed the in vitro PMN experiments, contributed to data interpretation and edited the manuscript. K.N.K estimated EdU labeled cells with Imaris, performed PNA FISH, helped with the co-localization, illustrated Fig. 10 and edited the manuscript. P.S. contributed to data interpretation and edited the manuscript. T.B. performed the in vitro PMN experiments and PNA FISH, contributed to data interpretation and edited the manuscript.

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Conflicts of interest. None declared.

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