



Secreted Phosphatase and Deoxyribonuclease Are Required by *Pseudomonas aeruginosa* To Defend against Neutrophil Extracellular Traps

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ABSTRACT Neutrophil extracellular traps (NETs) are produced by neutrophils as an innate immune defense mechanism to trap and kill microbial pathogens. NETs are comprised of ejected chromatin that forms a lattice structure enmeshed with numerous antimicrobial proteins. In addition to forming the structural backbone of NETs, extracellular DNA (eDNA) has membrane-disrupting antimicrobial activity that contributes to NET killing. Many pathogens produce secreted extracellular DNases to evade the antimicrobial activity of NETs. *Pseudomonas aeruginosa* encodes an operon of two secreted enzymes, a predicted alkaline phosphatase and a DNase. The DNase (*eddB*) degrades eDNA to use as a nutrient source. Here we report that both eDNA and NETs are potent inducers of this DNase-phosphatase operon. Furthermore, the secreted DNase contributes to degrading NET DNA and defends *P. aeruginosa* against NET-mediated killing. We demonstrate that EddA has both alkaline phosphatase and phosphodiesterase (PDase) activities and also protects against the antimicrobial activity of NETs. Although the phosphatase does not cause DNA degradation similar to that of the DNase, its protective function is likely a result of removing the cation-chelating phosphates from the eDNA phosphodiester backbone. Therefore, both the DNase and PDase contribute to defense against NET killing of *P. aeruginosa*, highlighting the role of DNA-manipulating enzymes in targeting the eDNA in neutrophil extracellular traps.

KEYWORDS *Pseudomonas aeruginosa*, deoxyribonuclease, neutrophil extracellular trap, phosphatase

As an opportunistic pathogen, *Pseudomonas aeruginosa* causes chronic infections in immunocompromised individuals, including those with the heritable genetic disease cystic fibrosis (CF). However, *P. aeruginosa* is also capable of infecting burn and diabetic foot wounds and causing nosocomial infections as well as eye and urinary tract infections. During chronic infection in the CF lung, the immune response is dominated by an influx and activation of neutrophils, in an effort to contain and clear *P. aeruginosa* (1, 2). The antimicrobial strategies employed by neutrophils include the phagocytosis of bacterial invaders, the release of toxic effectors, and the production of neutrophil extracellular traps (NETs) (3, 4). As recently described innate immune effectors, NETs are capable of ensnaring viruses, fungi, and both Gram-positive and Gram-negative bacteria (3, 4).

The deployment of NETs (a process termed NETosis) can result from a lytic (3) or a nonlytic (vital) (5, 6) pathway in neutrophils and consists of an ejected DNA lattice that is enmeshed with numerous neutrophil granule proteins. Lytic NETosis can be chemically stimulated with phorbol myristate acetate (PMA) treatment of isolated neutro-

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phils, and vital NETosis has been reported in neutrophils with intact membranes, which are still capable of phagocytosis and chemotaxis (5). Extracellular traps with a DNA backbone have also been observed for other innate immune cells, including mast cells, eosinophils, and macrophages (7). NETs have been observed in *P. aeruginosa* eye (8) and skin (9) infection models as well as in sputum recovered from CF lungs (10–12). Although they perform the innate immune function of trapping and killing bacteria, excessive NET production contributes to airway obstruction, inflammation, and pathophysiology and has been observed under numerous noninfectious conditions (13, 14).

NET antimicrobial activity is generally attributed to the neutrophil granular and chromatin proteins that are localized to the DNA lattice (4). Antibodies against the neutrophil-bound proteins, such as histones or cathepsin G, have been used to reduce bacterial killing, demonstrating their direct role in killing (3, 15). We previously showed that DNA has a potent antimicrobial activity by chelating surface-bound cations, disrupting outer and inner membrane integrity, which results in rapid, lytic cell death (16). DNA also contributes to bacterial killing by NETs and can be neutralized with treatment of NETs with excess cations or by alkaline phosphatase (AP) treatment (9). The cation-chelating phosphodiester backbone of DNA is responsible for the antibacterial nature of DNA, and we hypothesize a model where phosphatase treatment blocks NET killing by removing phosphates from DNA but does not alter the NET structure or the function or localization of some NET effector proteins (9).

Several microbial pathogens have been shown to defend against NETs through diverse strategies. *Streptococcus pneumoniae* produces a capsule and D-alanylated lipoteichoic acid on the bacterial surface that reduces the trapping and killing of the bacteria in NETs (17). The secretion of extracellular nucleases that degrade the chromatin backbone of NETs is a conserved evasion strategy. A growing list of bacteria that employ nucleases includes the Gram-positive bacteria *Staphylococcus aureus*, *S. pneumoniae*, group A *Streptococcus*, and *Streptococcus suis* and, recently, the Gram-negative bacteria *Vibrio cholerae*, *Neisseria gonorrhoeae*, and *Prevotella intermedia* (18–22). DNase production allows pathogens to evade killing by degrading the NET backbone and diluting the concentration of toxic NET-bound effector proteins while also directly neutralizing the antimicrobial activity of DNA (9, 16). *S. aureus* produces a nuclease that degrades DNA and RNA to 5'-phosphomononucleotides and dinucleotides, which are subsequently converted to 2'-deoxyadenosine (dAdo) through the AdsA enzyme (23). The dAdo product is cytotoxic to macrophages and therefore protects *S. aureus* by restricting these innate immune cells from entering staphylococcal abscesses (23).

An early observation of CF patient pathology was that the CF sputum is highly enriched in DNA (24). Given the strong recruitment of neutrophils in the lungs of individuals suffering from cystic fibrosis (1, 2) and the direct observation of NETs in CF sputum (10–12), this suggests that much of the sputum DNA is derived from neutrophils. *P. aeruginosa* is a dominant pathogen in CF infections and demonstrates greater resistance to NET killing than *S. aureus* and *Escherichia coli* (9). *P. aeruginosa* promotes the tolerance of NETs through the modification of the outer surface lipopolysaccharide (LPS) charge by the addition of aminoarabinose to lipid A or by the production of surface spermidine, both of which mask the negatively charged surface groups (9, 25). These surface modifications also contribute to resistance to antimicrobial peptides and aminoglycosides, both cationic antibiotics that bind to bacterial surfaces with a negative charge, as a first step in disrupting the membrane or entering the cell (16, 25–27).

We previously identified a secreted DNase that is required for DNA degradation and the utilization of DNA as a phosphate source (28). The DNase EddB (extracellular DNA degradation protein B) is downstream of a predicted alkaline phosphatase (16, 29). Since the operon is induced by phosphate limitation (28), it is likely that these two enzymes contribute to phosphate acquisition. Here we test the hypothesis that these secreted enzymes are involved in defense against the DNA component of neutrophil extracellular traps.

RESULTS

Phosphodiesterase-DNase operon induced by eDNA and NETs. We previously demonstrated that the *P. aeruginosa* xcp type II secretion system is required for the secretion of a DNase capable of degrading extracellular DNA (eDNA) (28). The DNase EddB, encoded by PA3909, promotes *P. aeruginosa* growth when DNA is supplied as the sole phosphate source, and DNA can be used as a source of carbon and nitrogen (28). Interestingly, the upstream open reading frame (ORF) PA3910 is predicted to encode an alkaline phosphatase, EddA, and shares a promoter with *eddB*, forming a bicistronic operon (16, 29). EddA contains a predicted signal peptide for the twin-arginine translation (TAT) pathway, which transports folded proteins across the inner membrane, and uses the Xcp pathway to cross the outer membrane (30). Alkaline phosphatase proteins fall into 3 groups: PhoX, PhoA, and PhoD-type phosphatases (31). EddA shares ~49% homology with *Bacillus subtilis* PhoD, the archetypal secreted alkaline phosphatase (32). Interestingly, PhoD phosphatases possess both monoesterase as well as phosphodiesterase (PDase) activities (30, 31). Moreover, EddA homologs, such as the *Zymomonas mobilis* CP4 PhoD alkaline phosphatase, have been shown to hydrolyze nucleotides more effectively than sugar phosphates (33).

Given that *eddB* are both likely involved in utilizing DNA as a nutrient, we assessed whether their expression is regulated by the presence of extracellular DNA (Fig. 1). Quantitative reverse transcription-PCR (qRT-PCR) analysis of *eddB* expression revealed that this gene is highly induced in the presence of low levels of eDNA after incubation for 60 and 120 min (Fig. 1A) (28). During infection, *P. aeruginosa* encounters and must defend against neutrophil extracellular traps, particularly in the CF lung. We sought to extend our analysis to determine whether NETs were also capable of inducing the expression of these genes. We used PMA treatment to induce the production of NETs in human neutrophils, which also induced the expression of *eddB* to comparable levels when exposed to pure DNA (Fig. 1A). There was no observable growth (optical density at 600 nm [OD₆₀₀]) during these quantitative RT-PCR (qRT-PCR) experiments, as *P. aeruginosa* was suspended in buffers that lack all nutrients. Gene expression was normalized to the expression of the 16S rRNA genes, which can normalize for any subtle differences in growth in the presence of DNA or NETs.

We confirmed the dynamic nature of the NET-induced gene expression pattern by utilizing the *P. aeruginosa eddB::lux* reporter strain in incubation experiments with neutrophils that were PMA stimulated to produce NETs (Fig. 1B). *P. aeruginosa* appears to sense the DNA in a NET and increase the gene expression level of an operon that produces a secreted phosphatase and DNase.

Secreted DNase and phosphodiesterase activities. We wanted to confirm the predicted enzymatic function of EddA and EddB, both of which are secreted extracellularly by *P. aeruginosa*. EddB was previously shown to degrade bacterial genomic DNA in the presence of excess Ca²⁺ and Mg²⁺ (28). To quantitate DNA degradation, we used a fluorescence-based DNA degradation assay using bacterial supernatants as the source of bacterial enzymes. Bacterial supernatants from *P. aeruginosa* lacking either the Xcp type II secretion system or PA3909 (*eddB*) were strongly impaired for the degradation of Sytox green-labeled extracellular salmon sperm DNA, relative to the control (Fig. 2A). Complementation with pEddB restored the capacity of *P. aeruginosa eddB::lux* to degrade eDNA, with a slight delay in the degradation kinetics (Fig. 2A).

To assess the phosphatase activity of EddA, we utilized a colorimetric assay in which bacterial supernatants were added to the AP (monoesterase) substrate nitroblue tetrazolium-5-bromo-4-chloro-3-indolylphosphate (NBT/BCIP). Coincubation of wild-type *P. aeruginosa* bacterial supernatants with NBT/BCIP led to a rapid increase in the amount of the cleaved substrate (Fig. 2B). Monoesterase activity was strongly dependent on the presence of a functional Xcp type II secretion system and on PA3910/EddA. Complementation with pEddA restored the capacity of *P. aeruginosa eddA::lacZ* to cleave the AP substrate (Fig. 2B).

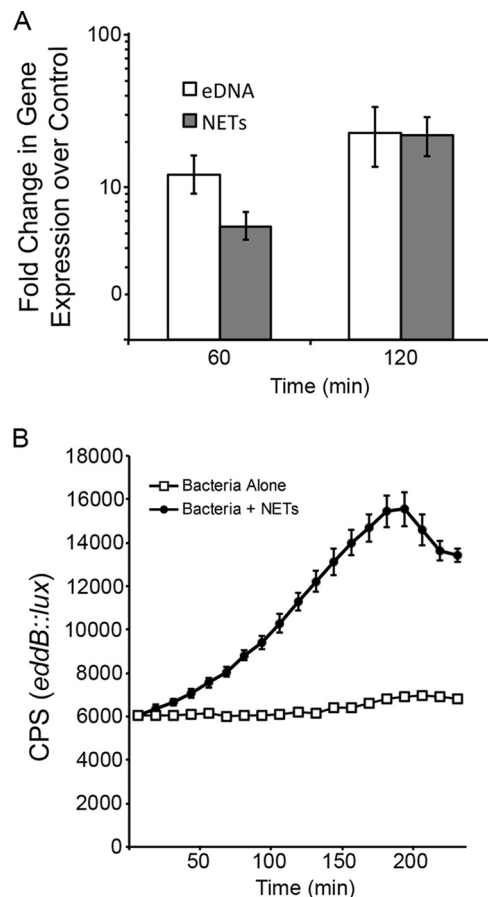


FIG 1 Expression of the *eddAB* operon is induced by eDNA and exposure to neutrophil extracellular traps. (A) qRT-PCR was used to measure *eddB* expression during exposure of *P. aeruginosa* to 0.2% eDNA or NETs. Bacteria were resuspended in 10 mM Tris buffer (pH 7) with 0.2% DNA or in HBSS with human neutrophils (MOI of 4:1) that were PMA stimulated to induce NET formation. After incubation, total bacterial RNA was extracted, converted to cDNA, and used as the template in qRT-PCR analyses, where values shown are the means and standard deviations from 8 replicates. Expression levels under the test conditions were compared to those under control conditions without eDNA or NETs and normalized to the expression level of the 16S rRNA housekeeping gene. (B) The *P. aeruginosa* *eddB::lux* reporter strain was grown in the absence or presence of PMA-activated human neutrophils (MOI of 10:1), and gene expression (counts per second) was measured every 20 min. The values shown are the means and standard deviations from 6 replicates.

We wanted to determine if EddA also had phosphodiesterase activity, which would support the cleavage of phosphates from DNA. To detect secreted phosphodiesterase activity, cell-free culture supernatants were incubated with the phosphodiesterase substrate bis(*p*-nitrophenyl)phosphate (BNPP) (31). The type II secretion mutant had low levels of phosphodiesterase activity (Fig. 2C), which were similar to those of wild-type PAO1 grown under phosphate-rich (1,600 μ M) conditions (data not shown). The *eddAB* double mutant showed a modest defect for phosphodiesterase activity, which was restored to higher-than-parental levels when overexpressing pEddA from a plasmid (Fig. 2C). In both the DNase and phosphatase assays, there appeared to be low levels of activity remaining in the supernatants of *eddA* and *eddB* mutants, suggesting the presence of other enzymes with these activities (Fig. 2A to C).

EddA was cloned as a C-terminally His-tagged protein (EddA-His₆), which permitted the purification of protein using HisPur cobalt resin. Purified EddA-His₆ was incubated with the phosphodiesterase substrate BNPP and demonstrated enzyme activity (Fig. 2D). Consistent with other PhoD-type phosphatases, EddA has both monoesterase and phosphodiesterase activities, which supports the hypothesis that EddA is capable of the cleavage of phosphates from the DNA backbone.

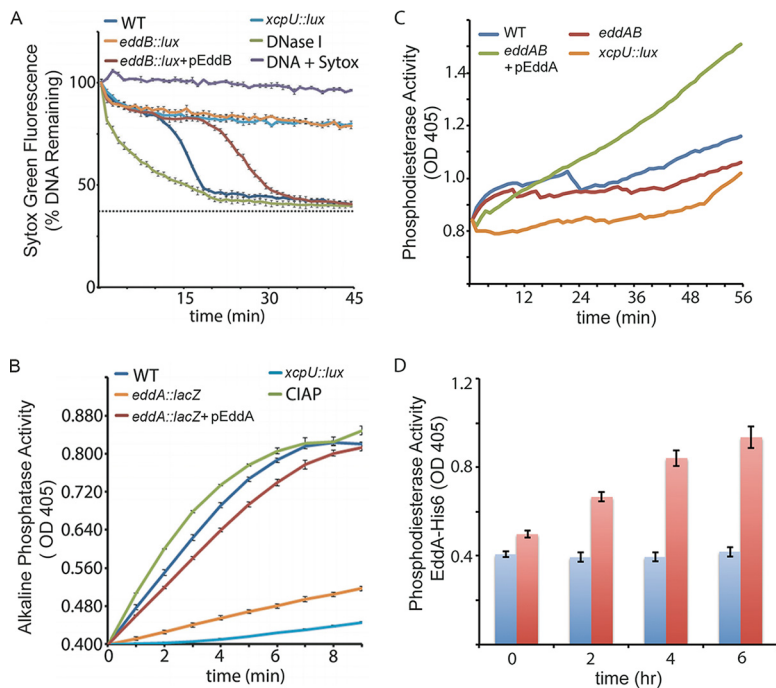


FIG 2 DNase and phosphatase activities of EddB and EddA. (A) DNase activity measured in culture supernatants that were incubated with 1 μ g DNA and 2.5 μ M Sytox green. Supernatants were isolated from cultures grown in limiting-phosphate BM2 medium. Green fluorescence was measured every minute from black-well, clear-bottom, 96-well microplates. Values shown are the means and standard deviations from quadruplicate samples. WT, wild type. (B) Alkaline phosphatase activity measured in culture supernatants incubated with the NBT/BCIP monoesterase substrate in alkaline phosphatase buffer. The colored AP reaction product was measured by monitoring the OD₄₀₅. (C and D) Phosphodiesterase activity measured in culture supernatants (C) or from purified EddA-His₆ (red bars) incubated with 25 mM BNPP compared to the negative-control sample with no EddA-His₆ (blue bars) (D). Values shown are means and standard deviations from triplicate samples.

DNase but not phosphatase dissolves neutrophil extracellular traps. NET killing of *P. aeruginosa* can be neutralized with treatments that specifically target the DNA component of the NETs (9). Exogenous DNase is frequently used to dissolve NETs and block bacterial killing (3, 9). Therefore, in an attempt to neutralize the cation-chelating activity of DNA but to retain the overall NET structure, NETs were treated with calf intestinal alkaline phosphatase (CIAP) (9). Phosphatase treatment blocked NET killing of *P. aeruginosa* and *E. coli* and blocked bacterial killing with pure DNA but did not alter NET structure or function or the localization of NET effectors (9). These experiments confirmed that the phosphates of DNA were responsible for the antimicrobial activity of DNA in a neutrophil extracellular trap. Therefore, we hypothesized that the EddA phosphodiesterase might also protect *P. aeruginosa* from NETs.

To assess whether these enzymes differed in their capacities to degrade the DNA backbone of the NETs, we performed a NET degradation assay using the same Sytox green-based approach used with pure DNA (Fig. 2A) (20). We stimulated NETosis by the use of PMA and monitored NET production using fluorescence measurements of Sytox green incorporation into NETs. After 200 min of PMA induction, NETs were formed, and bacteria were introduced for coinubation with NETs to assess their effects on the total DNA fluorescence (Fig. 3). As predicted, wild-type *P. aeruginosa* was capable of the full degradation of NETs within 7 h after addition (Fig. 3). Disruption of either the *xcp* type II secretion system or the *eddAB* enzymes resulted in the stability of NETs (Fig. 3). Complementation of the Δ *eddAB* double mutant with plasmid-encoded pEddB restored the NET degradation capacity of the Δ *eddAB* mutant to a level comparable to that of the wild-type parental strain (Fig. 3). Exogenous DNase was the most effective treatment for degrading NETs. However,

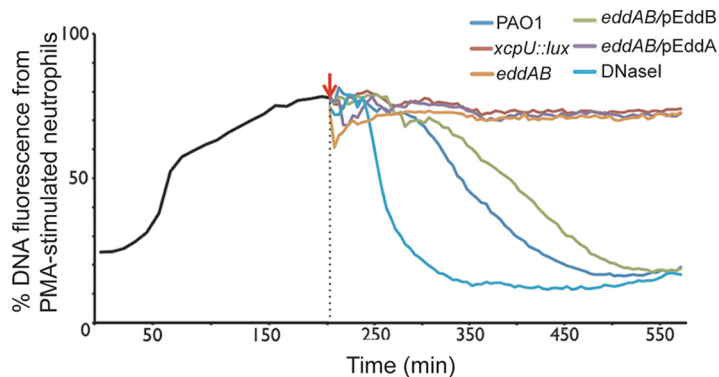


FIG 3 Degradation of DNA in neutrophil extracellular traps by secreted DNase from *P. aeruginosa*. DNA release from 1×10^6 neutrophils was stimulated with PMA, and the resulting NET DNA was stained with the cell-impermeant fluorescent DNA dye Sytox green. An increase in NET formation was seen up to 200 min, at which time *P. aeruginosa* cultures were added to the NETs to determine their effect on DNA integrity (red arrow). Staining of NET DNA with Sytox green was measured in 10-min intervals, and the values shown are the means and standard deviations from 8 replicates. The percentages of DNA fluorescence are relative to the value for the Triton X-100-treated neutrophil positive control (100% lysis) ($n = 3$).

complementation of the $\Delta eddAB$ double mutant with the plasmid-encoded pEddA phosphodiesterase had no effect on degrading the DNA from PMA-stimulated neutrophils (Fig. 3). We conclude that only the DNase can cause significant damage to and degradation of the NET structure.

Phosphatase and DNase are required to defend against NETs. Since it is common among bacteria to produce secreted DNases that degrade and disarm NETs (18–22), we tested the role of the EddA phosphatase and EddB DNase in protecting *P. aeruginosa* from neutrophil extracellular traps. Neutrophils were stimulated with PMA and exposed to *P. aeruginosa* wild-type and mutant cultures, and viable counts were determined to monitor bacterial viability. Interestingly, the *xcp* type II secretion system mutant demonstrated reduced tolerance to PMA-stimulated NETs (Fig. 4A). This result suggests that enzymes secreted by the type II secretion system are required to protect *P. aeruginosa* from NETs. The *eddAB* double mutant showed increased susceptibility to NET killing, and complementation with either pEddA or pEddB expressed in *trans* restored *P. aeruginosa* survival (Fig. 4B). While neither enzyme alone restored wild-type levels of NET tolerance, the enzymes likely work together in wild-type strain PAO1. Interestingly, the bacterial phosphodiesterase EddA significantly increased the survival of *P. aeruginosa* when exposed to NETs; however, it seems incapable of robust NET degradation relative to EddB (Fig. 3 and 4B). This protective function is congruent with the previous observation that NETs are no longer antimicrobial after treatment with CIAP (9).

DNase activity in culture supernatants of cystic fibrosis isolates. The CF lung is highly enriched in eDNA of neutrophil origin, which is the reason why inhaled recombinant DNase is used to thin CF sputum and alleviate infection symptoms (34). Therefore, we sought to characterize whether *P. aeruginosa* epidemic strains isolated from cystic fibrosis patients shared the capacity to degrade DNA. We obtained a panel of transmissible *P. aeruginosa* isolates from the Adult Cystic Fibrosis Clinic culture collection at the University of Calgary (35) and assessed their capacity to degrade DNA using the Sytox green-labeled DNA assay (Fig. 5). CF isolates produced a spectrum of activity ranging from high to low levels of DNA degradation (Fig. 5). However, 12 out of the 36 CF isolates tested did not have the capacity to degrade extracellular DNA. This phenotypic diversity is common among CF isolates, despite relatively similar genetic backgrounds, and is due to the complex selection pressure in the CF lung and the accumulation of loss-of-function mutations.

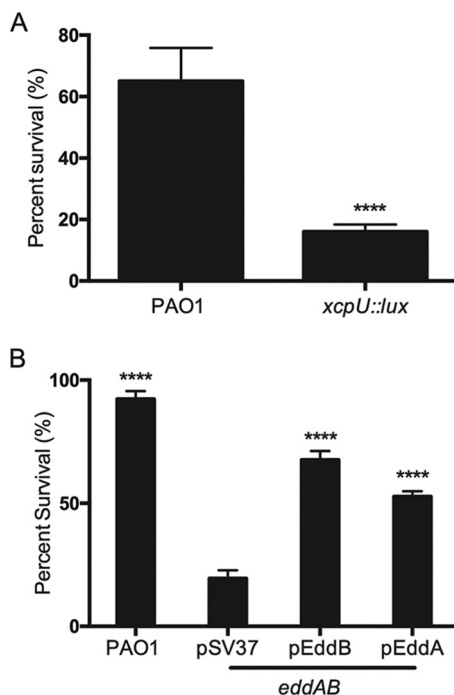


FIG 4 *P. aeruginosa* survival after exposure to neutrophil extracellular traps. A total of 1×10^7 CFU of *P. aeruginosa* bacteria were incubated with PMA-treated neutrophils (MOI of 10:1), and bacterial viability was determined by direct plate counts (CFU per milliliter) before and after 4 h of incubation with NETs. Bacterial counts were normalized to counts in the absence of neutrophils. DNase I was added exogenously 30 min prior to the end of the experiment to degrade NETs and ensure accurate counts of recoverable colonies. Data are presented as the average percentages of bacterial survival, and error bars represent the standard deviations from 6 replicates ($n = 3$). (A) Wild-type PAO1 survival compared to that of the *xcpU::lux* type II secretion mutant. Significant differences were determined by using the unpaired *t* test (****, $P < 0.0001$). (B) Bacterial survival of wild-type PAO1 and the *eddAB* double mutant, which was complemented with pEddA or pEddB. Significant differences compared to the double mutant were determined by one-way ANOVA with Bonferroni posttests (****, $P < 0.0001$).

DISCUSSION

In order to combat bacterial infection, NETs are deployed by neutrophils in numerous infection sites, including the CF lung (10–12). Although many proteins enmeshed in NETs (histones, cathepsin G, and calprotectin) have demonstrated antimicrobial properties (3, 4), here we focus on bacterial defense against the antimicrobial activity of DNA. The DNA backbone itself also contributes to the broad-spectrum antibacterial mechanism of NETs, through a mechanism of disrupting membrane integrity by chelating bacterial membrane-bound divalent cations (9, 16). The concentration of DNA in CF sputum can reach as high as 20 mg/ml, a level sufficient to promote membrane destabilization and rapid lysis of *P. aeruginosa* (16, 36, 37). This toxic effect exerted by eDNA may explain why multiple species, including CF *P. aeruginosa* isolates, share a virulence strategy of secreting DNA-degrading enzymes (Fig. 5).

Here we demonstrate that the Gram-negative, opportunistic pathogen *P. aeruginosa* employs two DNA-modifying type II secreted enzymes, the phosphodiesterase EddA and the DNase EddB, to promote bacterial evasion of NETs. Although we demonstrate that secreted EddA possesses alkaline phosphatase and phosphodiesterase activities (Fig. 2) and can promote bacterial viability (Fig. 4), the enzyme does not degrade NETs in our fluorescence-based assay (Fig. 3). The bacterial phosphatase EddA behaves similarly to calf intestinal alkaline phosphatase, which also blocks NET killing without disrupting the NET structure (9). Therefore, we propose that the bacterial phosphodiesterase EddB cleaves phosphates from DNA in a NET and neutralizes its cation-chelating, antimicrobial activity.

The DNase-PDase operon seems to have a dual role in phosphate acquisition (28)

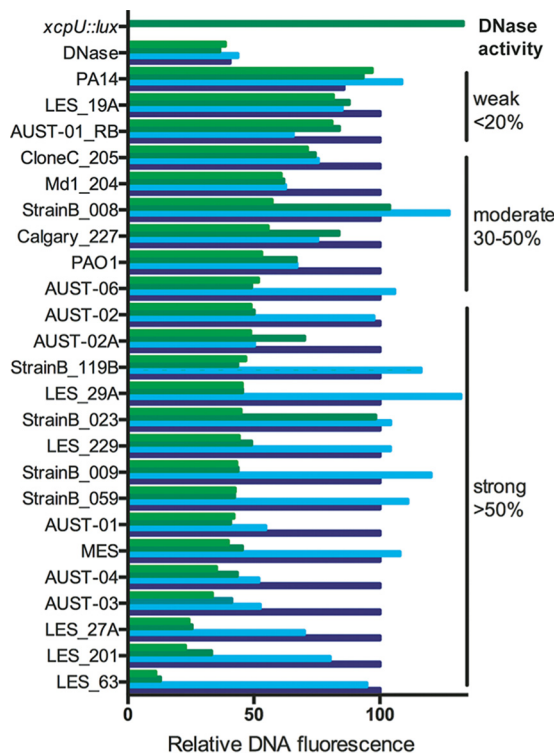


FIG 5 Clinical CF *Pseudomonas aeruginosa* isolates produce extracellular DNase activity under phosphate-limiting conditions. Cell-free supernatants were collected from cultures grown in BM2 medium with a range of phosphate concentrations, including 200 μM (light green), 400 μM (dark green), 800 μM (light blue), and 1,600 μM (dark blue). Supernatants were coincubated with Sytox green-stained DNA, and green fluorescence was measured in 10-min intervals. Relative DNA fluorescence values were derived by calculating the decrease in DNA fluorescence compared that under phosphate-rich conditions of 1,600 μM phosphate ($t = 3$ h). Pure DNase I (3 U) was used as a positive control, and the *xcpU::lux* mutant was used as a negative-control strain.

and in protection against neutrophil extracellular traps. It is also possible that secreted phosphatases cleave phosphates from the bacterial cell surface, as some enzymes can be surface localized (38). The removal of cell surface phosphates might limit DNA binding and therefore protect the outer membrane from DNA-mediated killing, which normally sequesters surface-bound cations and disturbs outer and inner membrane integrity (9).

Phosphate limitation is a nutrient cue shared by multiple Gram-negative bacteria to produce phosphate acquisition pathways, including phosphatases and phosphate uptake systems (39). The *pho* regulon is not only required for phosphate homeostasis but is also an important stress response pathway that contributes to bacterial virulence and cytotoxicity (39, 40). The DNase was originally shown to be regulated under phosphate limitation (28); however, limiting phosphate levels does not solely regulate the expression of this operon. We also detected expression or secreted enzyme activity in rich brain heart infusion (BHI) medium (Fig. 4B) and in the presence of pure eDNA or NETs, respectively (Fig. 1). Others have suggested a virulence role for the *P. aeruginosa* phosphodiesterase-DNase operon, due to its induced expression upon exposure to human airway epithelial cells (41). These observations highlight that multiple host cues can induce the expression of these virulence factors, and they are likely expressed in the CF lung.

Although the lower airways during chronic CF infections are colonized by complex multispecies communities referred to as the CF lung microbiome, *P. aeruginosa* remains a dominant pathogen (42). Chronic infection leads to a sustained inflammatory immune response and the heavy recruitment of NET-producing neutrophils (10–12). The capacity of *P. aeruginosa* to tolerate this NET-rich environment suggests that the bacterium

TABLE 1 Bacterial strains, plasmids, and primers

Strain, plasmid, or primer	Description or sequence	Source or reference
Strains		
PAO1	Wild type	R. E. Hancock
<i>eddB::lux</i>	Mini-Tn5- <i>lux</i> insertion mutant in PA3909 (EddB); transcriptional <i>lux</i> fusion	29
<i>eddA::lacZ</i>	Mini-Tn5- <i>lacZ</i> insertion mutant in PA3910 (EddA)	46
Δ <i>eddAB</i>	Unmarked deletion mutant of <i>eddAB</i>	This study
<i>xcpU::lux</i>	Mini-Tn5- <i>lux</i> insertion mutant in <i>xcpU</i> ; nontranscriptional <i>lux</i> fusion	45
Plasmids		
pEddA	EddA as an EcoRI-HindIII fragment in pPSV37 with a C-terminal His tag (EddA-His ₆)	This study
pEddB	EddB cloned as a BamHI-XhoI fragment into pUCP22	This study
pPSV37	IPTG-inducible expression vector; Gen ^r	44
pUCP22	Multicopy shuttle cloning vector; Amp ^r Gen ^r	47
Primers		
16SrtF	GAAATCCCCGGGCTCAACCTG	45
16SrtR	CCCCACGCTTTCGCACCTCA	45
EddB-F	GAGGATCCATGCACCCCTTGCCTAACGC	This study
EddB-R	GACTCGAGCTACCTGCGGTGCTTCTTCATCG	This study
EddA-soe-F	AAGCCTGAAACTCACTGAAG	This study
EddA-soe-R	ATACGCATGCCGCGATATCCGGATTGATCCCGAAAC	This study
EddB-soe-F	CGGCGGCATGCGTATAGGAAAGGGCTATTCTACG	This study
EddB-soe-R	TGGAGTGACCTCCGTTCC	This study
EddA-RBS-F	ATGCGAATTCAGGAGGAAACGATGAGTGGGATGGACCTCAAGCGC	This study
EddA-6×His-R	ATGCAAGCTTTCAGTGGTGATGGTATGATGGGCGCCGTCGGGCTGCAGTTCCTG	This study

can detoxify and tolerate the eDNA and NETs present in CF sputum. Recombinant human DNase (rhDNase) is used as an inhaled treatment in CF patients, and its beneficial effects are attributed to reducing sputum viscosity, facilitating airway clearance, improving lung function, and reducing the risk of exacerbation (24). Bacterial DNase and PDase may favor the bacterium by protecting *P. aeruginosa* from NET killing but may also promote its own clearance by thinning the CF sputum. Overall, the effects of bacterial DNA-modifying enzymes are likely insufficient to promote clearance, hence the need for exogenous rhDNase treatment. We demonstrate a virulence role for two extracellular DNA-modifying enzymes in defending against NET-mediated killing, which reinforces the concept that eDNA is an important innate immune effector that bacteria directly target to promote survival during infections.

MATERIALS AND METHODS

Bacterial strains, plasmids, and primers. The bacterial strains utilized in this study are summarized in Table 1 and in Fig. 5. *Pseudomonas aeruginosa* PAO1 was used as the wild-type strain. The mini-Tn5-*lux* mutants *eddB::lux* and *xcpU::lux* function as both transposon insertion mutants and transcriptional *luxCDABE* fusions that were constructed previously (29). *P. aeruginosa* was cultured at 37°C in defined BM2 minimal medium (0.1 M HEPES [pH 7], 7 mM ammonium chloride, 20 mM sodium succinate [pH 6.7], 2 mM magnesium sulfate, 10 μM iron sulfate, 1,600 μM phosphate buffer [pH 7.2], 1.62 μM manganese sulfate, 2.45 μM calcium chloride, 13.91 μM zinc chloride, 4.69 μM boric acid, 0.67 μM cobalt chloride) supplemented with 10 mM MgCl₂. Limiting-phosphate BM2 medium for culture or DNase and phosphatase supernatant assays contained 400 μM phosphate buffer (pH 7.2), rather than 1,600 μM. To assess protective effects of DNase or phosphodiesterase activity, *P. aeruginosa* was cultured in BHI medium (Difco) overnight at 37°C, subcultured 1:1,000 in fresh medium, and grown to mid-log phase. When added, salmon sperm DNA was used at the concentrations indicated (catalog number UB1440552; USB).

Mutant construction and complementation. The generation of the *P. aeruginosa eddAB* double mutant was conducted by allelic exchange as described previously (43). Briefly, wild-type PAO1 genomic DNA flanking the *eddAB* operon was amplified by using sequence-specific primers that added complementary regions to each amplified product. Stitching (splicing by overhang extension [SOE]) PCR was used to combine each flanking region into one PCR product that was complementary to the flanking regions of the *eddAB* operon but lacked the coding sequence. This SOE PCR product was gateway cloned (BP reaction) into a gateway-compatible vector backbone overnight, and the reaction was halted with the addition of proteinase K. Gateway constructs were transformed into electrocompetent *E. coli* DH10B cells. The isolated plasmid was then transformed into S17.1λpir and then mated with wild-type *P. aeruginosa* via puddle mating. Single-crossover events were selected for by growing *P. aeruginosa* on *Pseudomonas* isolation agar plus gentamicin (100 μg/ml). Counterselection for double recombination events was achieved by plating colonies on medium containing 15% sucrose.

The construction of the pEddA expression plasmid was completed by PCR amplifying the coding sequence from *P. aeruginosa* PAO1 genomic DNA. The amplified PCR product was cloned as an EcoRI-HindIII fragment with a C-terminal His₆ tag into the isopropyl- β -D-thiogalactopyranoside (IPTG)-inducible pPSV37 vector (44). The pEddB expression plasmid contained *eddB* cloned as a BamHI-XhoI product into the pUCP22 vector. Digested PCR products were gel purified, ligated, and directly transformed into *P. aeruginosa* PAO1 Δ eddAB, selecting for gentamicin resistance (30 μ g/ml).

Neutrophil isolation. Neutrophils were isolated from healthy donors as described previously (9). Briefly, whole blood was collected and mixed 5:1 in acid citrate dextrose, followed by the removal of red blood cells using dextran sedimentation and hypotonic lysis with KCl. After all red blood cells were lysed, the cell pellet was subjected to Ficoll-Histopaque density centrifugation. The subsequent pellet was resuspended in 2 ml of HBSS (Hanks' balanced salt solution, no cations) (catalog number 14175-095; Invitrogen) or BHI medium for NET killing assays. The viable neutrophil concentration was determined by using a hemocytometer and trypan blue staining.

Bactericidal NET assays. Quantification of bacterial viability was performed by direct plate counting methods, where a reduction in cell number indicates bacterial killing (9). NET killing experiments were performed in HBSS (Fig. 4A) or BHI medium (Fig. 4B). Isolated human neutrophils were mixed with mid-log-phase bacteria in black, clear-bottom, 96-well plates with 2.0×10^7 CFU bacteria and 2.0×10^6 neutrophils (multiplicity of infection [MOI] of 10:1). After 1 to 4 h of incubation, 50 μ l of a DNase I solution (430 kU/ml; VWR) was added to every well, mixed, and incubated for 30 min at 37°C, in order to release bacteria trapped in NETs for accurate plate counts. Fifteen microliters of the suspension was serially diluted (1/10) in a 0.9% NaCl solution in a sterile 96-well plate, and 5 μ l from each well was stamped onto LB agar plates to obtain bacterial plate count data at time zero (T_0) and after 4 h (T_4). CFU per milliliter values from the T_4 and T_0 time points were used to calculate the percent survival by subtracting the T_4 plate counts from the T_0 plate counts and dividing the value under "bacterium and neutrophil" conditions by the value under "bacterium-alone" conditions and multiplying this value by 100.

Gene expression assays. Gene expression assays were carried out with a method similar to the one described previously (9, 16, 28). In brief, fresh subcultures of PAO1 *eddB::lux* were normalized and diluted to a final concentration of 4×10^7 CFU/well into BM2–400 μ M phosphate buffer (a threshold concentration of phosphate that neither induces nor suppresses gene expression) or BM2–400 μ M phosphate supplemented with 0.2% DNA (2 mg/ml) or 3-h PMA-stimulated NETs generated by 1×10^6 neutrophils. Cultures were overlaid with mineral oil to prevent evaporation. The OD₆₀₀ and counts per second were monitored every 10 min over time in the Wallac Victor³ luminescence plate reader (Perkin-Elmer) at 37°C.

Quantitative reverse transcription-PCR was performed as described previously (45). The gene expression levels of *eddB* (*PA3909*) in triplicate 200- μ l samples of wild-type PAO1, under the conditions outlined above, were measured after 60 and 120 min of coinubation. RNA was extracted according to the manufacturer's instructions (Bacteria Protect and RNeasy; Qiagen). *Pseudomonas* gene expression levels were measured by using the iQ SYBR green supermix (Bio-Rad) and bacterium-specific primers for *eddB* and the 16S rRNA housekeeping gene. For qRT-PCR, quantification and melting curve analyses were performed with an iQ5 instrument (Bio-Rad) according to the manufacturer's instructions. Each reaction is done in triplicate, and standard deviations were used to calculate a range of fold activation values by using the $2^{-\Delta\Delta C_T}$ method (45).

DNase and phosphatase assays. To assess the production of EddA and EddB, PAO1 bacteria were cultured overnight in BM2 medium under limited-phosphate conditions (400 μ M phosphate buffer [pH 7.2], 10 mM MgCl₂) or in BHI medium (Difco) overnight at 37°C. Epidemic CF strains were cultured in BM2 medium under a range of phosphate conditions (200, 400, 800, and 1,600 μ M phosphate buffer, pH 7.2). Cultures of *P. aeruginosa* strains grown overnight were normalized to an OD₆₀₀ of 1, and supernatants were collected by centrifugation at 8,000 rpm for 10 min. To assess DNase activity, 100 μ l of the supernatant was incubated with 100 μ l of 0.2 μ g/ μ l salmon sperm DNA or 10⁶ PMA-stimulated neutrophils with 2.5 μ M Sytox green (as outlined above) in a black-walled, clear-bottom, 96-well plate and placed into a Wallac³ plate reader, and the plate was read every minute for 120 min.

For alkaline phosphatase activity, the substrate NBT/BCIP (Sigma-Aldrich) was used. For phosphodiesterase activity, 0.25 mM the substrate BNPP was used (Sigma-Aldrich). For enzyme assays, 100 μ l of the clarified bacterial supernatants was added to 100 μ l of 2 \times alkaline phosphatase buffer (1.0 M diethanolamine buffer with 0.50 mM magnesium chloride [pH 9.8]) supplemented with NBT/BCIP or BNPP in a 96-well, black-walled, clear-bottom plate, and the OD₄₀₅ was measured every minute for 30 min.

Purification of C-terminally His-tagged EddA. Mid-log-phase cultures of *P. aeruginosa*/pPSV37 expressing EddA-His₆ under the control of an IPTG-inducible promoter were grown in LB supplemented with 30 μ g/ml gentamicin until an OD₆₀₀ of ~0.5 was reached, when IPTG was added to the culture (0.2 mM) for a subsequent 3-h induction at 37°C. Cells were pelleted, the supernatant was removed, and the pellet was frozen prior to cell lysis and protein purification. The frozen pellet was gently resuspended in 5 ml 4°C wash buffer (0.5 \times phosphate-buffered saline [PBS] [pH 7.1], 300 mM NaCl, 5 mM imidazole) and sonicated for 15 min (5 s on and 10 s off) to lyse the resuspended pellet on ice. The lysate was centrifuged in a 4°C Beckman Coulter Avanti JA-20 rotor at 15,000 rpm for 30 min. The soluble fraction supernatant was removed and stored on ice. One milliliter of HisPur (ThermoFisher Scientific) cobalt Sepharose was equilibrated with 10 ml of 4°C wash buffer. The soluble lysate was then passed over the protein column twice, and the slurry was then washed by the addition of 10 ml wash buffer to the protein column. The recombinant soluble protein was eluted by the addition of 1 ml elution buffer (0.5 \times PBS [pH 7.1], 300 mM NaCl, 50 mM NaPO₄, 250 mM imidazole) and dialyzed (molecular weight cutoff [MWCO] of 4,000 Da) into the appropriate buffer for 18 h at 4°C.

Western blot detection of His-tagged EddA. Ten microliters of the eluted protein was boiled for 10 min in SDS-PAGE loading buffer. The protein was loaded onto a 10% SDS-PAGE gel, electrophoresed for 45 min at 180 V, and then transferred to nitrocellulose by using the Trans-Blot turbo transfer system (Bio-Rad). After the transfer, the nitrocellulose membrane was incubated in 1× Tris-buffered saline plus Tween (TBST) with 5% (wt/vol) skim milk powder for 18 h at 4°C. After incubation, anti-His (Sigma-Aldrich) was added at a 1:10,000 dilution for 1 h at room temperature, with rocking. The blot was then washed 3 times using 20 ml fresh TBST. The secondary anti-mouse antibody (Cell Signaling Technology) was added at a 1:10,000 dilution in 5% skim milk plus TBST, and the mixture was rocked at room temperature for an additional hour. The blot was washed with TBST as described above and then developed by using chemiluminescent Western blotting detection (Pierce ECL).

Statistical analysis. Statistical analysis was performed by using GraphPad Prism v4.0 software. One-way analyses of variance (ANOVA) with Bonferroni posttests and unpaired *t* tests were used to calculate significant differences for bacterial survival and gene expression. Significant differences refer to *P* values of ≤ 0.05 or as otherwise indicated.

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