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Research Article

Microbiology

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Bacterial neuraminidase facilitates mucosal infection by participating in biofilm production

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Many respiratory pathogens, including *Hemophilus influenzae*, *Streptococcus pneumoniae*, and *Pseudomonas aeruginosa*, express neuraminidases that can cleave α 2,3-linked sialic acids from glycoconjugates. As mucosal surfaces are heavily sialylated, neuraminidases have been thought to modify epithelial cells by exposing potential bacterial receptors. However, in contrast to neuraminidase produced by the influenza virus, a role for bacterial neuraminidase in pathogenesis has not yet been clearly established. We constructed a mutant of *P. aeruginosa* PAO1 by deleting the PA2794 neuraminidase locus (Δ 2794) and tested its virulence and immunostimulatory capabilities in a mouse model of infection. Although fully virulent when introduced i.p., the Δ 2794 mutant was unable to establish respiratory infection by i.n. inoculation. The inability to colonize the respiratory tract correlated with diminished production of biofilm, as assessed by scanning electron microscopy and in vitro assays. The importance of neuraminidase in biofilm production was further demonstrated by showing that viral neuraminidase inhibitors in clinical use blocked *P. aeruginosa* biofilm production in vitro as well. The *P. aeruginosa* neuraminidase has a key role in the initial stages of pulmonary infection by targeting bacterial glycoconjugates and contributing to the formation of biofilm. Inhibiting bacterial neuraminidases could provide a novel mechanism to prevent bacterial pneumonia.

Introduction

Neuraminidases (sialidases) are produced by a wide variety of mucosal pathogens, ranging from *Streptococcus pneumoniae* in the airway to *Vibrio cholerae* in the gut (1). While the central role of viral neuraminidase in pathogenesis of influenza is well established (2) and provides a target for both vaccines and chemotherapy, the contribution of bacterial neuraminidase to the pathogenesis of infection is not as clearly defined. Neuraminidase-producing species such as *Hemophilus* (3), *S. pneumoniae* (4, 5), and *Pseudomonas aeruginosa* (6) share a common ecological niche, colonizing the heavily sialylated secretions and surfaces of the upper respiratory tract. Although each can bind to asialylated glycolipids exposed by neuraminidase activity (7), they differ substantially in their ability to either metabolize (8) or incorporate sialic acid into surface structures (9). Thus it is likely that bacterial neuraminidases interact with both microbial and eukaryotic glycoconjugates (1).

P. aeruginosa is a major opportunistic pathogen, an important cause of nosocomial pneumonia as well as the chief cause of lung infection in cystic fibrosis (CF). Over 2 decades ago, neuraminidase production in isolates of *P. aeruginosa* from CF patients was described and suggested to contribute to pulmonary infection (10). In vitro studies documented that many pulmonary pathogens including *P. aeruginosa* bind to the GalNAc β 1,4Gal moiety exposed on asialylated glycolipids (7), suggesting that the abil-

ity to desialylate mucosal surfaces could contribute to bacterial colonization of the airways. The *P. aeruginosa* neuraminidase was cloned and characterized. It was shown to be osmoregulated, thought to be consistent with expression in the milieu of the CF lung (6) and capable of exposing the receptor asialoganglioside gangliotetraosylceramide (Gal β 1,2GalNAc β 1,4Gal β 1,4Glc β 1,1Cer) (asialoGM1) on the surface of CF airway cells in vitro, implying a role in pathogenesis (11). However, data confirming *P. aeruginosa* adherence to the airway surface in CF patients has been lacking (12). The current consensus suggests that organisms are predominantly entrapped in dehydrated secretions of the lung and – by shedding proinflammatory products – activate airway inflammation (13), a model that does not require direct attachment of organisms to the epithelial surface. Nonetheless, analyses of *P. aeruginosa* gene expression in CF patients document that the PA2794 neuraminidase locus is one of the most highly expressed genes in this patient population in vivo (14). Unlike other respiratory pathogens, *P. aeruginosa* cannot use sialic acid as a carbon source, nor does it contain sialic acid as a component of its LPS (15). Thus it seemed likely that there was some additional function for the enzyme relevant to the pathogenesis of respiratory tract infection. To better define the importance of bacterial neuraminidases, we constructed a *P. aeruginosa* mutant lacking the PA2794 *nanA* locus and analyzed it in a mouse model of infection. Our studies showed that the *P. aeruginosa* neuraminidase is involved in biofilm formation contributing to initial colonization of the airway. Furthermore, we demonstrated that this activity can be blocked by viral neuraminidase inhibitors in clinical use indicating a novel therapeutic target for preventing bacterial pneumonia.

Nonstandard abbreviations used: asialoGM1, Gal β 1,2GalNAc β 1,4Gal β 1,4Glc β 1,1Cer, asialoganglioside gangliotetraosylceramide; CF, cystic fibrosis; LB, Luria broth; OMP, outer membrane protein; PMN, polymorphonuclear leukocyte.

Conflict of interest: The authors have declared that no conflict of interest exists.

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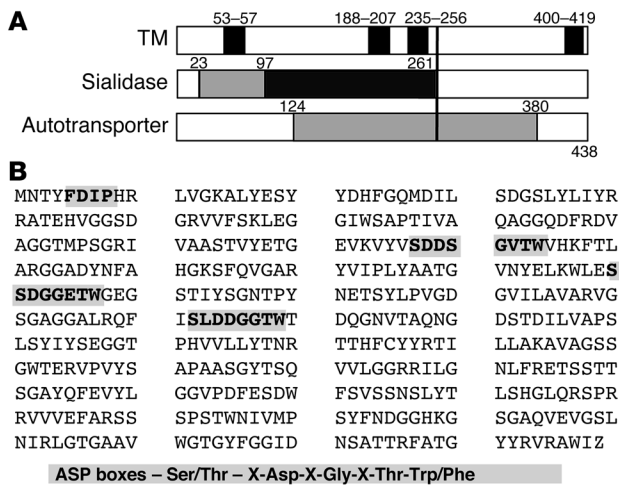


Figure 1 Properties of the PA2794 locus. (A) Sequence predictions for PA2794. Shown are the results of structural prediction and sequence similarity searching techniques. The gene PA2794 is depicted 3 times, and shaded regions demonstrate distinct functional predictions. Regions are indicated using amino acid positions. Predicted transmembrane (TM) regions are shown first. These predictions were made using the programs TMPred (http://www.ch.embnet.org/software/TMPRED_form.html) and TMAP (<http://bioweb.pasteur.fr/seqanal/interfaces/tmap.html>). The second gene depiction shows the region most similar to other sialidase genes. Lighter gray shading indicates the region predicted by BLAST analysis. For an inclusive definition of the boundaries of this region, we used the most extreme amino- and carboxy-terminal positions of BLAST alignments for the first 50 hits that were also labeled as sialidase or neuraminidase genes. The darker shading corresponds to the sialidase region predicted by the SMART program (<http://smart.embl-heidelberg.de/>). The third gene depiction shows the region similar to autotransporter genes. This region was determined by the BLAST alignment with a putative autotransporter (Magn03008734, GenBank accession no. ZP_00054112) from the *Magnetospirillum magnetotacticum* genome. This region had an E value score of 6×10^{-4} in our BLAST search. (B) Predicted PAO1 neuraminidase amino acid sequence. ASP boxes are shaded gray.

Results

Construction of a nanA null mutant. Sequence predictions for the PA2794 locus were analyzed using ORFcurator (16), which identified both the sialidase region and a domain expected to have autotransporter function (Figure 1A) (17) consistent with previous reports (18). The predicted sequence included the ASP boxes expected to interact with sialic acid (Figure 1B) (19). An in-frame nonpolar deletion allele of the predicted neuraminidase open reading frame (PA2794, *nanA*) was constructed and used to replace the wild-type gene in PAO1 (Figure 2A). Loss of neuraminidase activity was documented by an assay monitoring the ability of culture supernatants to expose asialoGM1 from human airway epithelial cells (Figure 2B) (6). The deletion was shown not to impose any metabolic consequences on the fitness of the mutant, as growth curves of the wild-type, mutant, and complemented strains were comparable (Figure 2C).

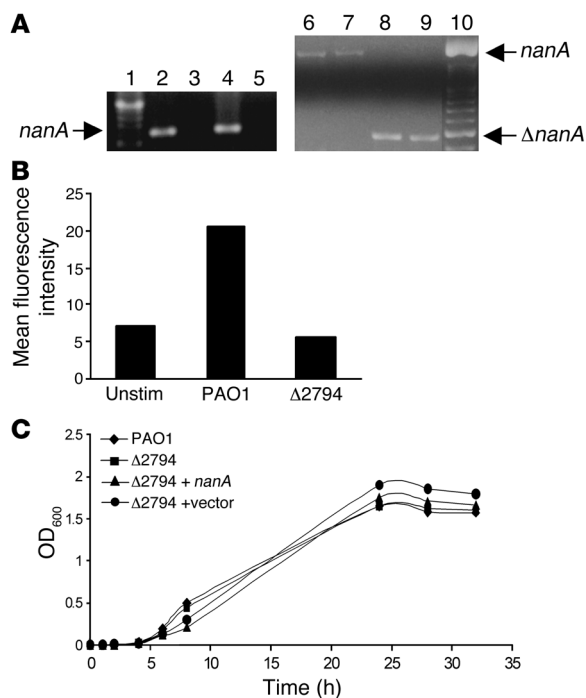
Virulence properties of Δ2794 neuraminidase mutant differ depending upon the route of inoculation. To evaluate the role of the neuraminidase in respiratory tract infection, 2×10^8 CFU inocula of the wild-type PAO1, Δ2794, and Δ2794 + *nanA* strains were used to

infect 7- to 10-day-old BALB/c mice by the i.n. route, and morbidity and mortality were assessed 18 hours after infection (Figure 3, A–D) (20). In contrast to PAO1, the Δ2794 mutant was readily cleared from the respiratory tract: 33% of the mice infected with Δ2794 developed pneumonia versus 78% of the mice infected with PAO1 ($P < 0.05$). Significantly fewer mice infected with the Δ2794 mutant became bacteremic: 10% versus 44% ($P < 0.05$). Mortality rates were not significantly different (Figure 3A). Quantification of lung mRNA indicated less chemokine KC expression, which correlated with fewer polymorphonuclear leukocytes (PMNs) recruited to the lungs, in mice inoculated with Δ2794 compared with PAO1 ($P < 0.05$; Figure 3, B and C). Histopathology similarly demonstrated substantially decreased inflammation in Δ2794-infected lungs (Figure 3D).

We also compared the virulence of wild-type and Δ2794 strains when introduced into mice by the i.p. route. This route of administration could potentially uncover differences in the immunogenicity of the mutant due to changes in glycosylation that might affect LPS or other surface structures. However, there were no differences in virulence when equal inocula of the wild-type and mutant strains were injected by the i.p. route (Figure 3E).

Immunostimulatory properties of the PA2794 mutant. To account for the attenuated phenotype of the Δ2794 mutant when introduced into the airways, we characterized its ability to attach to and stimulate chemokine and cytokine expression in both airway epithelial cells and macrophages (Figures 4 and 5). Although we had documented that PAO1 culture supernatant exposed more asialoGM1 on the surface of human airway cells than did Δ2794 (Figure 2B), there were no significant differences in either bacterial adherence or the induction of IL-8 expression by the wild-type and Δ2794 mutant strains (Figure 4, A and B), in contrast to published reports that used concentrated purified enzyme (6). As bacterial modification of surface glycoconjugates could affect interactions with phagocytic cells, we compared the uptake and killing by RAW cells. Both wild-type and mutant bacteria were efficiently phagocytosed by RAW cells and both induced equivalent amounts of TNF-α production (Figure 5, A and B). In addition, we compared outer membrane proteins (OMPs) and secreted exoproducts from the wild-type and mutant strains and detected no differences (data not shown).

Effects of the PA2794 neuraminidase locus on LPS. Having found no biologically important alteration in eukaryotic surface glycosylation that could be attributed to effects of the neuraminidase, we next examined possible effects on surface structures including LPS. An extensive biochemical analysis of *P. aeruginosa* LPS had previously failed to identify sialic acid (15), in contrast to other neuraminidase-producing organisms such as *Hemophilus* (21). Although the enzyme could target a different amino sugar involved in LPS biosynthesis, a GC-mass spectroscopic analysis kindly performed by R. Ernst (University of Washington, Seattle, Washington, USA) did not reveal any differences in LPS lipid A structures among the wild-type, mutant, or complemented strains (data not shown) (22). As LPS structure is responsible for the resistance of *P. aeruginosa* to the lytic effects of normal human complement, we compared PAO1 and Δ2794 sensitivity to 10% human serum and also found no differences (data not shown). These negative results, along with the observation that the wild-type and mutant strains were equally virulent when injected i.p., indicate that the Δ2794 mutation does not have a major effect on LPS or its immunogenicity.

**Figure 2**

Characterization of the $\Delta 2794$ mutant. (A) Colony PCR for expression of *nanA*. Lanes 2–5 show PCR using primers from within the *nanA* gene; lanes 6–9 show PCR using primers flanking *nanA*. Lane 2, PAO1; lane 3, $\Delta 2794$; lane 4, $\Delta 2794 + nanA$; lane 5, $\Delta 2794$ with vector alone; lane 6, PAO1; lane 7, PAK; lane 8, $\Delta 2794$; lane 9, $\Delta 2794$ PAK. Lanes 1 and 10 show molecular weight markers. (B) Amount of superficial asialoGM1 on 16HBE cells following exposure to bacterial supernatant from the indicated strains was quantified by flow cytometry. A representative experiment is shown. Unstim, unstimulated (i.e., treated with media alone). (C) Growth of strains in M9 media as determined by OD₆₀₀. A representative experiment is shown.

The PA2794 locus affects biofilm formation. Since the $\Delta 2794$ mutant appeared to be deficient solely in its ability to initiate infection by the mucosal route, we postulated that the neuraminidase might target other bacterial exopolysaccharides, such as those involved in biofilm formation. Bacteria-bacteria interactions were examined using crystal violet staining to quantify biofilm production in 96-well plates (Figure 6A) (23). The wild-type strain PAO1 containing an empty vector was used as a control, and all strains were grown under identical conditions in gentamicin selection. The $\Delta 2794$ mutant with the empty vector produced significantly less biofilm than PAO1 also containing the control vector ($P < 0.001$). Biofilm production by the $\Delta 2794$ mutant expressing the cloned *nanA* locus was significantly greater than that of the parental strain ($P < 0.0001$), consistent with the constitutive expression of the cloned gene. Similarly, overexpression of *nanA* by expressing the cloned gene in PAO1 also resulted in significantly greater biofilm production ($P < 0.0001$). In addition, we tested an independently derived mutation in strain PAK and a $\Delta 2794$ PAK mutant. Although PAK did not produce significant amounts of biofilm under control conditions, and deletion of the 2794 locus had no apparent effect, overexpression of the cloned 2794 locus in $\Delta 2794$ PAK resulted in a significant increase in the production of biofilm compared with the PAK parental strain ($P < 0.0001$). Biofilm assays were then performed in a flow cell (Figure 6C) (24) and on a rotating disc reactor (Figure 6D) (25) in order to formally evaluate the ability of the bacteria to form structured communities. In each assay system, the biofilm produced by $\Delta 2794$ exhibited dramatically changed architecture, and complementation of the phenotype by the cloned gene was observed. To establish that the mutant $\Delta 2794$ exhibits similarly impaired biofilm formation on epithelial cells, we incubated GFP-expressing bacteria with monolayers of airway epithelial cells and observed by fluorescence imaging that the $\Delta 2794$ organisms did not cluster or autoagglutinate, in contrast to PAO1 (Figure 6E).

Neuraminidase inhibitors block biofilm production. Drugs that target the neuraminidase produced by influenza viruses are an important component of antiviral chemotherapy, used for both prophylaxis and treatment. As bacterial and viral neuraminidases can share common ASP boxes that interact with sialic acid (19), we postulated that the neuraminidase inhibitors designed to block the influenza enzyme (26) might have sufficient avidity for the active site of the bacterial neuraminidase to inhibit biofilm formation. We tested the effects of the influenza virus neuraminidase inhibitors oseltamivir (27) and peramivir (28) on PAO1 biofilm formation using the crystal violet assay (Figure 7, A and B). A dose-dependent effect was observed with each of the 2 drugs, suggesting that *P. aeruginosa* biofilm formation may be a target to prevent infections in patients at risk by using neuraminidase inhibitors.

To document that these viral neuraminidase inhibitors are specifically interacting with the bacterial neuraminidase, we also tested their effect in blocking neuraminidase activity using the fluorescent substrate 2'-(4-methylumbelliferyl)- α -D-N-acetylneuraminic acid (Figure 7C). The *V. cholerae* enzyme as well as the *P. aeruginosa* PAO1 enzyme were inhibited by peramivir, suggesting that the active sites of both types of neuraminidases have some affinity for the available viral neuraminidase inhibitors.

Discussion

Bacterial neuraminidases have long been implicated in the pathogenesis of mucosal infection. Their prevalence and conservation among a very broad range of important human pathogens such as *S. pneumoniae* and *V. cholerae* as well as commensal flora, *Mycoplasma* (29), and even the gut flora of fish (30) implies a critical role in microbial ecology, which may vary according to bacterial species and sites of infection (1). The contribution of neuraminidase expression to the pathogenesis of respiratory infection is complex. The expected analyses of isogenic neuraminidase mutants has been accomplished in few bacterial species, due to gene redundancies (31, 32), involvement in metabolic pathways, and perhaps technical issues, as the construction of mutants has been unexpectedly difficult.

Initial studies of the *P. aeruginosa* neuraminidase performed with purified enzyme and in vitro analyses were entirely consistent with a role for the enzyme in modifying airway epithelial cell surfaces to facilitate bacterial attachment (6). Moreover, as CF airways were more readily modified than were normal airway cells (11), the *Pseudomonas* enzyme seemed likely to be important in that disease. However, in the studies presented herein, performed under more physiological conditions in vivo using isogenic mutants, we now find an entirely different function for the *P. aeruginosa* neuraminidase. The neuraminidase appears to

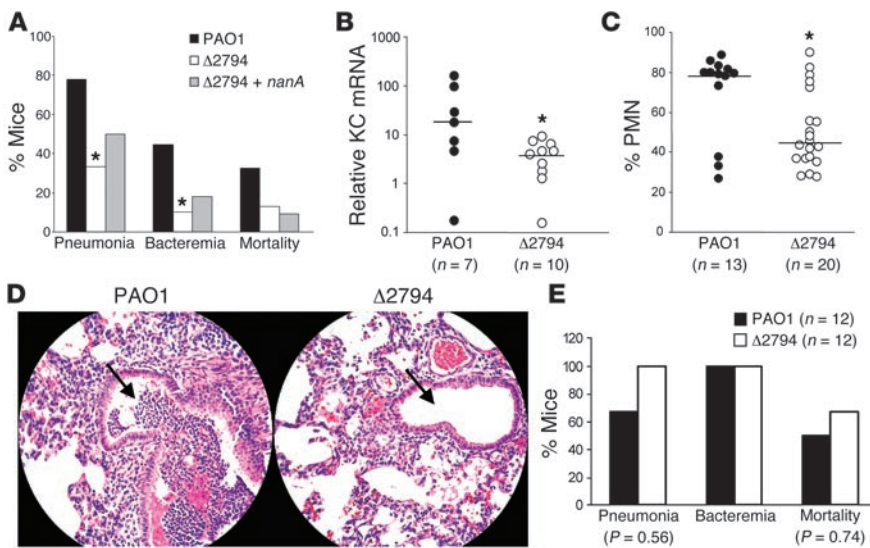


Figure 3
 Comparison of PAO1 and $\Delta 2794$ virulence. (A–D) Mouse model of pneumonia. (A) The percentage of the total number of mice inoculated i.n. with each strain that developed pneumonia or bacteremia or died is indicated. (B) KC mRNA expression was determined by real-time PCR and standardized to actin. Individual mouse values are shown; horizontal lines indicate the median value of each group. (C) Lung cell suspensions from mice inoculated with *P. aeruginosa* PAO1 or $\Delta 2794$ were stained for CD45 and Ly6G (i.e., PMNs). Leukocytes were gated based on CD45 expression, and the percentage of Ly6G-positive cells was determined. Individual mouse values are shown; horizontal lines indicate the median value of each group. * $P < 0.05$, Fisher’s exact test (A); nonparametric Mann-Whitney test (B and C). (D) H&E-stained sections of murine lungs 16 hours following inoculation with *P. aeruginosa* PAO1 or $\Delta 2794$. Arrows demonstrate that airways are full of PMNs in PAO1-infected lung but clear in $\Delta 2794$ -infected lung. (E) Comparison of PAO1 and $\Delta 2794$ virulence when introduced i.p. BALB/c mice (7–10 days old) were inoculated i.p. with PAO1 or $\Delta 2794$ ($n = 12$ per group). Mice were euthanized 16 hours later, and lungs and spleens were harvested for plating of single-cell suspensions. The percentage of mice inoculated with each strain that developed pneumonia or bacteremia or died is indicated. Pneumonia was defined as the recovery of more than 1,000 CFU per lung; bacteremia was defined as the presence of bacteria in the spleen.

be important for biofilm production, the cell-cell interactions which are critical even in the initial colonization process. Exactly how the PA2794 neuraminidase is involved in biofilm synthesis is unclear. Although the components of the PAO1 biofilm have not been fully defined, those of another *P. aeruginosa*, PA14, do not include sialic acid (33), indicating that the neuraminidase activity may be directed against a different sugar linkage on the bacterial surface. Pseudaminic acid – or a structure containing pseudaminic acid, a 9-carbon acidic sugar with structural similarity to the neuraminic acids – is a potential substrate and modifies several surface structures including LPS (34), pili (35), and flagella (36) in *P. aeruginosa*. Recent studies indicate that there are significant homologies among the genes involved in sialic acid O-acetylation in many bacterial species including *P. aeruginosa* strain 012, which produces pseudaminic acid but not sialic acid (37). Pseudaminic acid is involved in the glycosylation of *Campylobacter pylori* flagella (38), and mutants that lack pseudaminic acid fail to autoagglutinate (as we found in the $\Delta 2794$ mutant) and are attenuated in a ferret model of diarrheal disease. Mutations in the biosynthetic pathway involved in the addition of pseudaminic acid to flagella appear to contribute to this phenotype. Just as autolysins are necessary for cell wall biosynthesis, enzymes capable of cleaving carbohydrate linkages are necessary for the growth and modifica-

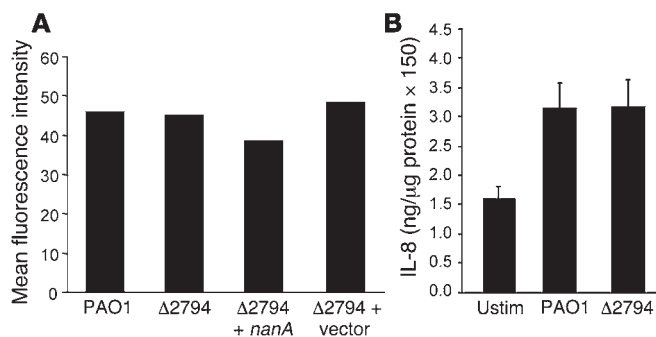
tion of extracellular polysaccharides during biofilm biosynthesis (39).

A recent survey of virulence factors expressed by *P. aeruginosa* isolated from CF patients indicated that the PA2794 neuraminidase locus is one of the most highly conserved (14). This is consistent with its role in biofilm production, as the primary mode of *P. aeruginosa* growth in the CF lung is in biofilm (40, 41). However, in contrast to current paradigms regarding *P. aeruginosa* gene expression in the CF lung, our data implies that cell-cell communication, as manifested by biofilm production, is critical even in the initial colonization process. In vivo, both mucin (42) and the presence of neutrophils (43) contribute to biofilm formation. Although the presence of “mature” biofilms with complex secondary structures may be more typical of long-standing infection, our data suggest that neuraminidase production is involved in cell-cell interactions necessary for colonization and persistence in the airway. The involvement of the neuraminidase locus appears to contribute to the initial stages of biofilm development. The $\Delta 2794$ mutants trapped in a planktonic form of growth were fully virulent in a model of i.p. sepsis in which replication and induction of a host immune response are sufficient to cause mortality, but could not efficiently colonize the lung. Additional in vitro assays assessing immunostimulatory interactions with both macrophages and epithelial cells revealed no significant differences among the wild-type and mutant strains. The bio-

logically significant consequence of the loss of neuraminidase appears to be limited to its defect in cell-cell aggregation.

The importance of biofilm production in the pathogenesis of respiratory tract infection caused by other common respiratory pathogens such as *H. influenzae* (21, 44) and *S. pneumoniae* has been well described (45). Less clear is whether these biofilms require the participation of neuraminidases for biosynthesis or modification. *H. influenzae* are known to produce sialylated biofilms that are important in the pathogenesis of mucosal infections, such as otitis media (46). While neuraminidase production by *Hemophilus* species has been reported (47), its specific involvement in colonization or pathogenesis has not yet been established. Several functions have been ascribed to the pneumococcal neuraminidase(s), which can desialylate host proteins (5) as well as surface components from adjacent flora (48) but does not appear to be critical for invasive infection (49), as was the case for the *Pseudomonas* enzyme. The surface location of the pneumococcal enzyme is consistent with a role in biofilm biology (4) but more definitive, functional experiments are necessary.

The mucosal surface of the gut is also colonized by organisms in biofilms (50), and neuraminidase expression has been well characterized in gastrointestinal pathogens (50) including *Salmonellae* (51), *Vibriosis* (52), and *Bacteroides* (8). Neuraminidases

**Figure 4**

Interactions of PAO1 and $\Delta 2794$ with human airway epithelial cells. **(A)** Bacterial adherence following a 1-hour exposure to 16HBE airway epithelial cells was determined by flow cytometry. Cells were stained with polyclonal anti-OMP followed by Alexa Fluor 488-conjugated anti-rabbit IgG. The number of bacteria bound to the surface was quantitated as mean fluorescence intensity. **(B)** Induction of IL-8 production by confluent monolayers of 1HAEo airway epithelial cells following exposure to bacteria — compared with induction by bacterial growth media alone as a control — was quantified by ELISA as previously described (57).

contribute to bacterial metabolism in some of these organisms, and bacterial surface sialylation may provide an immune masking function as well (3). However, even though neuraminidase expression by *V. cholerae* has been characterized for 40 years (52), its specific contribution to the pathogenesis of cholera remains obscure (53). Although cholera toxin targets sialic acid, it has never been clearly established how the neuraminidase is linked to the toxin (53) or if the enzyme is also involved in cell-cell interactions and the colonization process. There may be technical issues that have limited the analysis of cloned genes that contribute to biofilm formation, as it has only recently been recognized that the TEM β -lactamase often used for selection interferes with biofilm formation (54).

There has been great interest in identifying bacterial gene products that are essential for pathogenesis, as these could be targeted in the patient populations known to be at high risk for infection. This is an especially appealing strategy to prevent *P. aeruginosa* infection in CF and intensive care patients. Viral neuraminidase inhibitors have been very useful in the prevention and treatment of influenza, targeting similar high-risk patient populations. The PA2794 neuraminidase shares many conserved elements and folds in the manner predicted for other microbial neuraminidases (19, 55). A preliminary analysis of PA2794 crystal structure indicates that the enzyme shares the same sialic acid binding fold as does the influenza enzyme, but otherwise has little homology (L. Tong

Figure 5

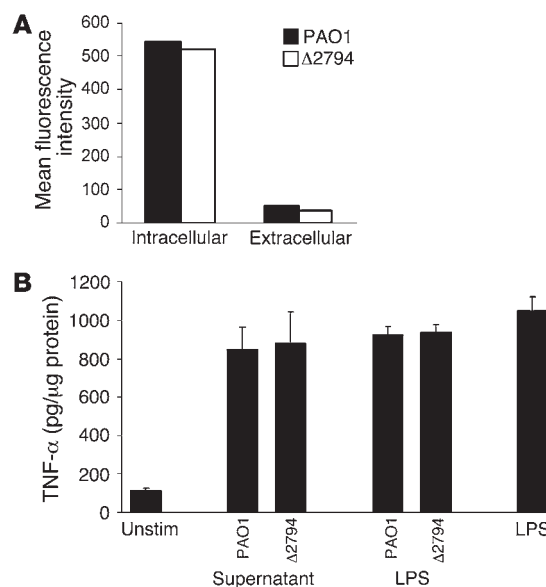
Interactions of PAO1 and $\Delta 2794$ with phagocytic cells. **(A)** Binding and phagocytosis of PAO1 and $\Delta 2794$ by RAW cells was determined by flow cytometry. Adherent organisms were quantified by staining with anti-OMP, and adherent plus internalized bacteria were quantified in RAW cells that were permeabilized prior to staining to determine total RAW cell-associated bacteria. **(B)** TNF- α production by RAW cells treated with media alone, bacterial culture supernatant, or LPS harvested from cultures of PAO1 and $\Delta 2794$ was quantified by ELISA. Purified LPS (Sigma-Aldrich) was used as a positive control.

and Y.-S. Hsiao, unpublished observations). The activity of the viral neuraminidase inhibitors in blocking *P. aeruginosa* biofilm production in vitro suggests that bacterial neuraminidases may also be a useful target to prevent infection in patients at high risk. These or similar compounds optimized for activity against bacterial enzymes could be especially useful in preventing *P. aeruginosa* colonization in CF patients. As neuraminidase expression is so highly conserved in mucosal pathogens, it should be possible to determine whether this approach blocks bacterial biofilm production and infection by other common pathogens.

Methods

Construction of a *P. aeruginosa* PAO1 neuraminidase null mutant. A *nanA* null mutant ($\Delta 2794$) was constructed by allelic replacement. An in-frame nonpolar deletion allele was constructed by removing the *nanA* coding sequence corresponding to amino acids 5–435 of the predicted 438-residue polypeptide and used to replace the full-length gene (1,317 base pairs) by the method previously described (56). Primers were designed using the published DNA sequence for the neuraminidase gene (designated PA2794) from *P. aeruginosa* strain PAO1 (GenBank accession no. AF236853). A *nanA* complementation plasmid was constructed by cloning a PCR product corresponding to the full-length neuraminidase open reading frame into plasmid pMMBGW with either a gentamicin or a penicillin resistance marker (56). The complementation clone or an empty vector control was introduced into the $\Delta 2794$ mutant by conjugation and selection on gentamicin (40 μ g/ml) or piperacillin (100 μ g/ml). The same procedure was carried out to generate a $\Delta 2794$ mutation in the *P. aeruginosa* strain PAK. The following primers were used for genotyping: internal to PA2794, 5'-CGCACTATACACAGGAACACG-3' and 5'-GCCTAGCGGAAGGATC-GTCGC-3'; external to PA2794, 5'-GATTATAAGTCTGCCGTCGG-3' and 5'-CTCGGGAAACGTGCACATCC-3'.

Bacterial strains and culture conditions. The standard laboratory strain of *P. aeruginosa* PAO1 was used as a prototype, grown in Luria broth (LB) or M9 media with Mg-glu as indicated. For complementation studies, the PA2794 locus was overexpressed in *E. coli* using pMMB67EH.gm and pMMB67EH.amp. Growth curves were obtained by growing bacteria in M9 media — with 40 μ g/ml gentamicin or 100 μ g/ml of piperacillin selection for strains containing plasmid — overnight to stationary phase, then



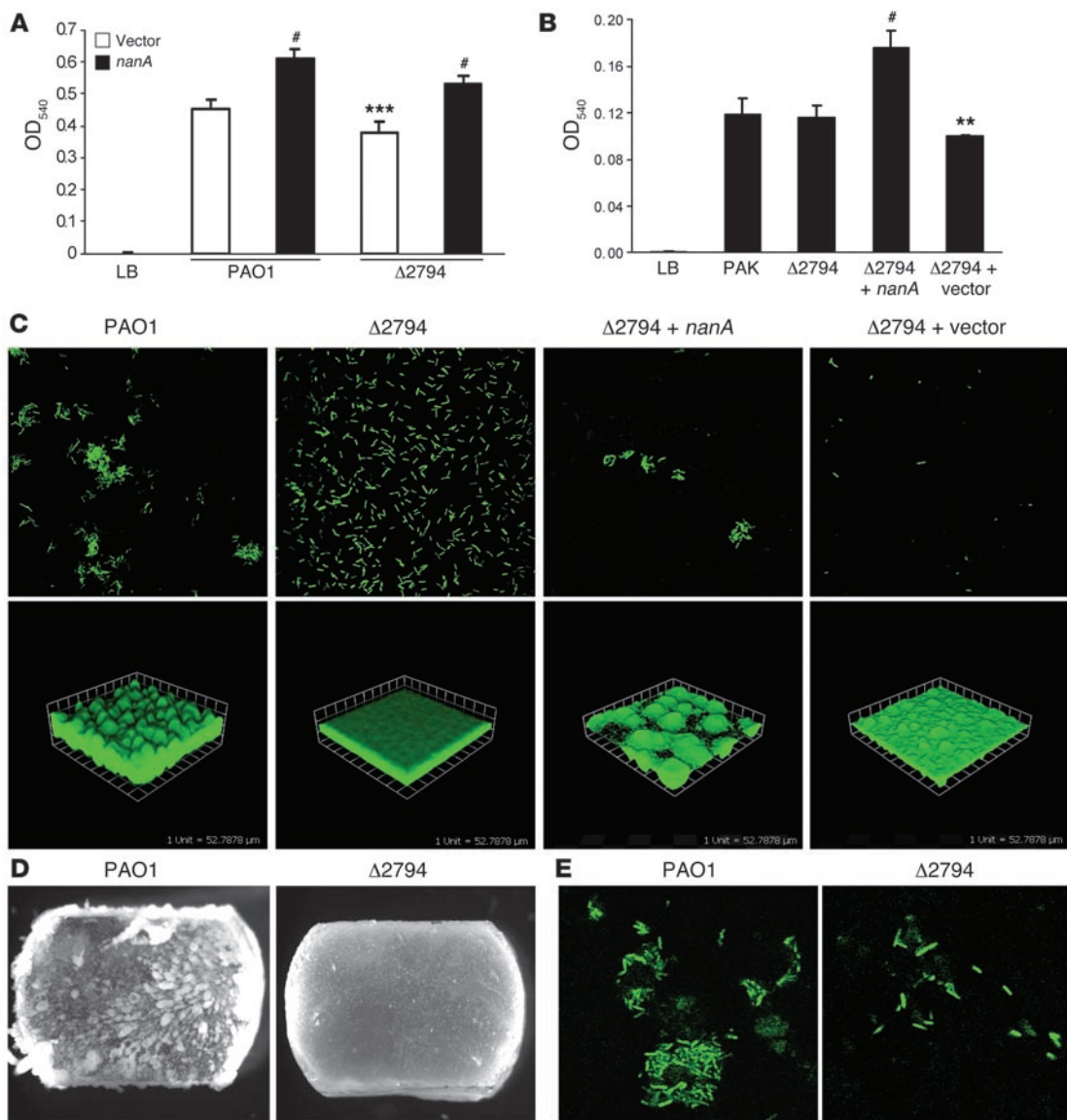


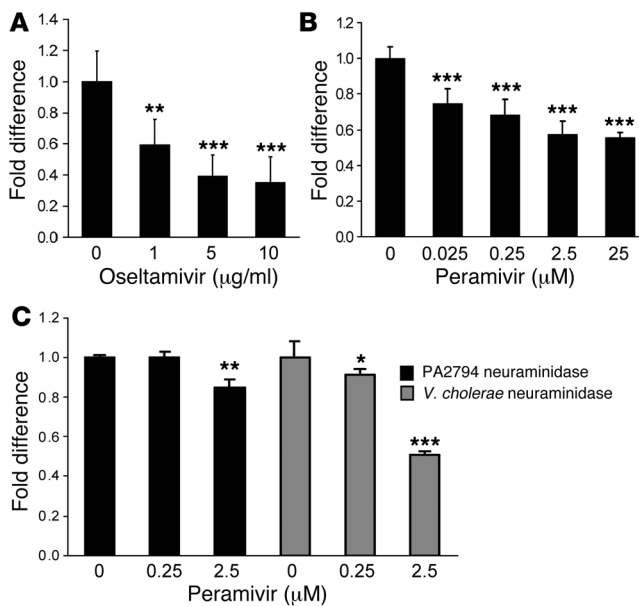
Figure 6

Comparison of biofilm production by PAO1 and $\Delta 2794$. **(A and B)** Biofilm production by bacterial strains detected by crystal violet staining. **(A)** PAO1 and $\Delta 2794$ each with either empty control vector (white bars) or cloned *nanA* (black bars) and **(B)** PAK, $\Delta 2794$ PAK, $\Delta 2794$ PAK + neuraminidase locus ($\Delta 2794 + nanA$), and $\Delta 2794$ PAK plus empty control vector are shown. $**P < 0.01$, $***P < 0.001$, $\#P < 0.0001$ versus PAO1 + vector **(A)** or PAK **(B)**. **(C)** Fluorescence microscope images (top row) and profiles (bottom row) of biofilm production in a flow cell. Compared with PAO1, $\Delta 2794$ formed less differentiated and flattened layers. The $\Delta 2794 + nanA$ complemented mutant showed restored, differentiated clumps of cells, while the $\Delta 2794$ mutant plus control vector did not. **(D)** Light microscope images of biofilms produced under more dynamic conditions using a rotating disc reactor that uses removable discs attached to a rotor in a chemostat. The disc rotation produces high-shear forces. High-shear conditions can accentuate deficiencies in attachment and matrix integrity. White, fluffy material is biofilm. The lack of biofilm formation by the mutant could be caused by defective matrix synthesis. **(E)** Bacteria expressing GFP (10^6) were incubated with confluent monolayers of 16HBE airway epithelial cells for 5 hours and visualized by confocal microscopy.

diluting 1:1,000 in fresh media and incubating at 37°C with shaking and OD₆₀₀ readings taken over time.

Epithelial cell culture. Originally obtained from D. Gruenert (California Pacific Medical Center Research Institute, San Francisco, California, USA), 1HAEo and 16HBE cells were grown in minimum essential medium with Earle’s salts supplemented with 10% fetal calf serum (Molecular Probes) as previously described (57). RAW cells were grown in RPMI medium 1640 with 10% fetal calf serum.

Biofilm assays. An overnight culture of bacteria grown in LB with shaking was diluted 1:100, and 100- μ l aliquots added to 96-well microtiter plates were incubated for 24–48 hours at 37°C (58). Crystal violet (0.1%) was added to each well for 15 minutes, rinsed 3 times with water, and then released with the addition of 200 μ l of 95% ethanol. Absorbance was determined at 540 nm. For experiments with inhibitors, the following modifications in the assay were used: for oseltamivir, an overnight culture of bacteria was diluted 1:100 in different doses of inhibitor in LB, and 100 μ l

**Figure 7**

Dose-dependent inhibition of biofilm production in response to viral neuraminidase inhibitors. PAO1 was grown in the presence of indicated concentrations of (A) osetamivir and (B) peramivir before plating for assay, and biofilm production was detected by crystal violet staining method. Fold difference compared with untreated is plotted. A representative experiment is shown. (C) Inhibition of bacterial neuraminidases by peramivir. Purified PAO1 neuraminidase and *V. cholerae* neuraminidase were incubated with and without peramivir, and neuraminidase activity was measured using the fluorescent substrate 2'-(4-methylumbelliferyl)- α -D-N-acetylneuraminic acid. Fold difference compared with untreated is plotted. A representative experiment is shown. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ versus untreated.

aliquots were plated and assayed as described above; for peramivir, an overnight culture of bacteria was diluted 1:100 in different doses of inhibitor in LB, incubated at room temperature for 48 hours, and then diluted 1:2 in LB, and 100 μ l aliquots were plated and assayed as described above. Each sample was tested in sextuplicate, the assay was repeated on 3 separate occasions, and representative data are shown. For the complementation studies, all of the PAO1 strains were transformed with either the control vector or the vector expressing the 2794 locus. Flow cell experiments and confocal microscopy were performed as previously described (24). For visualization by confocal microscopy, pMRP9-1 (which expresses GFP) was transformed into appropriate strains (59). The rotating disk reactor (25) was used for generating biofilms for microscopy and quantitative counts. Tryptic soy broth medium (1:100 strength) was used for these experiments.

Neuraminidase assays. The PAO1 enzyme (10 μ g/ml) was overexpressed and purified from *E. coli* using the pET28a vector (Novagen, EMD Biosciences), and a control *V. cholerae* neuraminidase (0.1 U/ml; Calbiochem, EMD Biosciences) was incubated with the fluorescent substrate 2'-(4-methylumbelliferyl)- α -D-N-acetylneuraminic acid (Sigma-Aldrich) (25 μ M in 0.9% NaCl) with or without peramivir (0.25 and 2.5 μ M) for 24–48 hours, and fluorescence was read at excitation 360 nm and emission 465 nm. Each data point was performed in sextuplicate. A representative experiment is shown.

IL-8 assays. Confluent monolayers of 1HAEo cells, weaned from serum overnight, were washed and stimulated with bacteria (1×10^8 CFU/ml) for 30 minutes. Fresh media plus gentamicin (100 μ g/ml) was added and then removed for chemokine analysis after 3 hours. ELISA for IL-8 (R&D Systems) was performed as previously described (56). Each data point was performed in quintuplicate and standardized by protein. Each experiment was performed at least 3 times, and a representative study is shown.

Quantification of epithelial sialylation by flow cytometry. 16HBE cells were grown in 24-well plates to confluence and exposed to bacterial supernatant concentrated 30-fold for 3–5 hours followed by 3 PBS washes. Cells were stained with rabbit polyclonal Anti asialo GM1 antibody (Wako) followed by Alexa Fluor 488 donkey anti-rabbit IgG (Molecular Probes). Cells detached from the plastic using 0.02% EGTA in HBSS were then fixed with 1% paraformaldehyde and analyzed on a FACSCalibur using CellQuest software (version 3.3; BD).

RAW cell binding and phagocytosis as determined by flow cytometry. RAW cells, a murine monocyte-macrophage cell line, were grown in 10-cm dishes and exposed to 1×10^8 bacteria for 30 minutes at 37°C. After 4 washes with PBS, 2 ml HBSS plus 0.02% EGTA was added, and cells were harvested. Cells were counted in a hemacytometer, and 1×10^6 cells were aliquoted per microfuge tube. PBS (1 ml) was added to each tube, and cells were pelleted at 400 g for 5 minutes. For extracellular binding determination, cells were incubated in 5% normal serum in PBS. For determination of total external and internalized bacteria, cells were incubated with Perm/Wash buffer (BD Biosciences – Pharmingen). In both cases, cells were then stained with rabbit anti-OMP antibody followed by Alexa Fluor 488 donkey anti-rabbit secondary and fixed with 1% paraformaldehyde and analyzed on a FACS-Calibur using CellQuest software (version 3.3; BD).

Mouse models of infection. Mouse protocol number AAAA1718 was approved by the Institutional Animal Care and Use Committee at Columbia University. Seven day old BALB/c mice were inoculated i.n. with 2×10^8 CFU of PAO1 or Δ 2794 in 10 μ l of PBS or i.p. with 5×10^5 CFU of PAO1 or Δ 2794 and euthanized 16 hours later with pentobarbital. Pneumonia was defined as the recovery of more than 1,000 CFU per lung, and bacteremia was defined as the recovery of bacteria from the spleen. The inflammatory response in vivo was assayed by flow cytometry as previously described (60). Single-cell suspensions of the lung were screened for the percentage of PMNs in the total leukocyte population by double staining with PE-labeled anti-CD45 and FITC-labeled anti-Ly6G antibodies (BD Biosciences – Pharmingen). Irrelevant, isotype-matched antibodies were used as a control. Cells were gated on the basis of their forward and side scatter profiles and analyzed for the expression of both CD45 and Ly6G.

Immunohistochemistry. Paraffin lung sections from mice infected with PAO1 and Δ 2794 were stained with H&E.

Real-time PCR. Lungs from PAO1- and Δ 2794-inoculated mice were obtained 16–18 hours after inoculation and stored in RNeasy (QIAGEN). RNA was isolated using the QIAGEN RNeasy Mini Kit. cDNA was made from 1 μ g of RNA using the iScript cDNA Synthesis Kit (Bio-Rad). For quantitative real-time PCR, amplification was performed in a LightCycler using the DNA Master SYBR Green I kit (Roche Diagnostics). Primers used for KC amplification were 5'-CCGCGCCTATCGCCAATGAGCTGCGC-3' and 5'-CTTGGGGACACCTTTTAGCATCTTTTGG-3', and 35 cycles were



run with denaturation at 95°C for 8 seconds, amplification at 56°C for 10 seconds, and extension at 72°C for 12 seconds. Actin was amplified on each individual sample and used as control for standardization. Primers used for actin amplification were 5'-GTGGGGCGCCACAGCACCACCA-3' and 5'-CGGTTGGCCTTGGGGTTCAGGGGGG-3', and 35 cycles were run with denaturation at 95°C for 8 seconds, amplification at 63°C for 10 seconds, and extension at 72°C for 12 seconds.

Adherence assay. 16HBE airway epithelial cells were stimulated with bacteria for 1 hour. After washing with PBS to remove unbound organisms, cells were stained with polyclonal anti-OMP followed by Alexa Fluor 488-conjugated anti-rabbit IgG (Molecular Probes). Fixed cells were analyzed by flow cytometry to quantitate the number of bacteria bound to the surface.

Statistics. For biofilm, neuraminidase, and IL-8 assays, means and standard deviations were calculated, and statistical significance was determined using 2-tailed unpaired Student's *t* test (biofilm assays) and 1-way analysis of variance with Bonferroni's post test (neuraminidase and IL-8 assays).

Tests of statistical significance were performed with GraphPad InStat (version 3.0; GraphPad Software).

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