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The Colonization Strategies of Nontypeable *Haemophilus influenzae*

Bacterial Colonization Factors and Vaccine Development



DOCTORAL DISSERTATION by due permission of the Faculty of Medicine, Lund University, Sweden.

To be defended at the main lecture hall of the Pathology building, Jan Waldenströms gata 59, Malmö, on Friday 30th of January 2015 at 13:00.

> *Faculty opponent* Professor Joachim Reidl, University of Graz

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Abstract

Nontypeable *Haemophilus influenzae* (NTHi) is a remarkably proficient colonizer of the human respiratory tract. The aim of this thesis has been to characterize currently known, and identify novel, bacterial factors involved in the key processes of colonization and pathogenesis, namely adherence to host tissue and evasion of the host innate immunity.

We have shown that the NTHi virulence factor Protein E mediates innate immune evasion by interacting with the C-terminus of host complement down-regulatory protein vitronectin. This interaction delayed the formation of the membrane attack complex on the bacterial surface and increased the bacterial serum-resistance. Furthermore, we identified a novel surface-exposed virulence factor designated *Haemophilus* Protein F that also sequesters vitronectin on the surface of the bacterial adherence to primary human bronchial cells and to the extracellular matrix protein laminin. Isogenic mutants devoid of Protein F were observed to adhere significantly less to vitronectin, epithelial cells and laminin. These interactions were characterized at the molecular level, and the N-terminus of Protein F was shown to contain the host-interacting region. Finally, we showed that immunization with Protein F conferred increased pulmonary clearance of NTHi *in vivo*.

In summary, we have identified, and at the molecular level characterized, novel virulence factors that may be suitable for future vaccine development.

Key words: ABC-transporter, bacterial colonization, *Haemophilus influenzae, Haemophilus* Protein F, laminin, NTHi, pathogenesis, vaccine, virulence, vitronectin.

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Bacterial Colonization Factors and Vaccine Development

Farshid Jalalvand



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To all my teachers in and outside of the home.

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Abbreviations

ABC-transporter	Adenosine triphosphate (ATP)-binding cassette transporter
AOM	Acute otitis media
COPD	Chronic obstructive pulmonary disease
ECM	Extracellular matrix
Fur	Ferric uptake regulator protein
Hib	<i>Haemophilus influenzae</i> serotype b
Ln	Laminin
LOS	Lipooligosaccharide
LPS	Lipopolysaccharide
MAC	Membrane attack complex
MMP-9	Metalloproteinase-9
NAD	Nicotinamide adenine dinucleotide
NLR	Nucleotide-binding oligomerization domain (NOD)-like receptor
NTHi	Nontypeable Haemophilus influenzae
ОМ	Otitis media
OMV	Outer membrane vesicle
ORF	Open reading frame
PAMP	Pathogen-associated molecular pattern
PF	Haemophilus Protein F
PRR	Pathogen pattern recognition receptor
TIMP-1	Tissue inhibitor of metalloproteinase-1
TLR	Toll-like receptor

Populärvetenskaplig sammanfattning

Bakterier är små encelliga organismer som lever i nästan varje tänkbar miljö på jorden, inklusive i och på människor. Vi är kontinuerligt koloniserade av bakterier i bland annat våra luftvägar, munhålor och mag-tarmkanaler. Denna samexistens är oftast ömsesidigt fördelaktig, eller åtminstone ofarlig för oss människor, och det är känt att bakterier hjälper oss bryta ner dietära ämnen samt förser oss med vitaminer. Men mikrober som tillhör normalfloran kan under vissa omständigheter orsaka infektioner av varierande allvarlighetsgrad.

Att kolonisera människans övre luftvägar, d.v.s. näsa, svalg och de övre delarna av broncherna, är ingen lätt uppgift för bakterier. Vi har flera skyddsmekanismer på plats för att försvara oss mot invaderande mikrober. Luftvägarna är klädda med ett slemproducerande cellhinna som kallas det *mukosala epitelet*. Epitelet i luftvägarna har dessutom ett ytterligare vapen till sitt förfogande; mikroborstar eller så kallade *cilier*. När partiklar eller mikrober andas in i luftvägarna fastnar de oftast i epitelets slemhinna och transporteras bort av cilierna till svalget för att sväljas ner till magsäcken för destruktion eller spottas ut. Utöver detta mekaniska skydd, finns människans medfödda immunförsvar på plats för att bekämpa bakteriella inkräktare. Fördelen med det medfödda immunförsvaret är att den till skillnad från det adaptiva antikroppsgenererande immunförsvaret inte behöver en uppstartstid utan är ständigt igång. Avhandlingens mål har varit att kartlägga hur den övre luftvägskoloniserande bakterien *Haemophilus influenzae* kringgår dessa antimikrobiella mekanismer för att orsaka infektioner såsom öroninflammation och perioder av försämring i KOLpatienter.

Vi har här kunnat visa att två molekyler som finns på bakteriens yta, kallade *Haemophilus* Protein F och Protein E, kamouflerar *H. influenzae* genom att binda det mänskliga proteinet vitronektin. Vitronektin är vanligtvis involverat i skyddet av kroppens egna celler från det medfödda immunförsvaret, d.v.s. den skyddar oss mot autoimmunitet. Genom att klä sig i detta protein, lurar bakterier vårt immunförsvar att de är kroppsegna celler och undgår dödande av det medfödda immunförsvaret.

Utöver det har vi visat hur Protein F fungerar som ett slags kardborre i luftvägarna. Den vidhäftar till både epitelceller samt epitelets underliggande vävnad, det så kallade *basalmembranet*. Genom att klistra sig på dessa vävnader undgår bakterien att transporteras bort av cilierna. Dessa två mekanismer, vidhäftning till epitelet och resistens mot det medfödda immunförsvaret, understödjer därmed *H. influenzaes* kolonisering av människans luftvägar.

Slutligen undersökte vi ifall Protein F kan användas som ett vaccin mot *H. influenzae*. Vi kunde visa att möss som immuniserades med Protein F rensade ut bakterierna mycket mer effektivt än möss som immuniserades med ett negativ kontrollprotein. Vår slutsats är att Protein F är en möjlig vaccinkandidat mot *H. influenzae* och bör utredas närmare.

Sammanfattningsvis har vi identifierat nya bakteriella koloniseringsfaktorer som hjälper *H. influenzae* överleva i människans luftvägar, samt visat att ett av dessa proteiner kan potentiellt utnyttjas i ett vaccin mot patogenen.

Introduction

Bacterial and Human Coexistence

Bacteria are remarkable life forms. Their presence is almost ubiquitous on Earth as they inhabit enormously diverse ecosystems ranging from oceanic and terrestrial subsurfaces devoid of sunlight, the mesosphere at altitudes exceeding 70 km above sea level, geothermally hot sulphur springs, to subglacial lakes and barren deserts (1-4). The majority, of course, live in the soil, shallow aquatic habitats and on other life forms such as animals and plants (1). Prokaryotes constitute a significant amount of the total biomass on the planet (5), "occupying" almost as much carbon as plants and an order of magnitude higher amounts of nitrogen and phosphorous according to some estimates, making them the largest living reservoir for essential nutrients (1). Simply put, their abundance and presence in every *milieu* on Earth is unquestionable. And since we humans are coinhabitants of the same planet, we too are ecological niches for bacterial life as they live in and on us (6).

Considering the ubiquity, abundance and remarkable adaptation of microbes to various environments, it does not come as a surprise that we, like all other multicellular life forms, become covered from head to toe (and inside out for that matter) in bacteria from the moment that we are born until we die, after which microbes help decompose our bodies in the proverbial "circle of life" (6, 7). Prokaryotic life traces back at least 3.4 billion years (8), and is considered to be the first cellular life form to exist on the planet (9, 10). Our own species, *Homo sapiens*, has existed for about 200 000 years (11). If we scale the duration of the bacterial inhabitation of Earth to 24 hours, humans have existed for approximately 5 seconds. With this in mind, one could argue that we are merely newly arrived visitors to an ancient microbial world.

Still, while here, the bacterial and human interplay is of huge importance for us. There are evidences of an intricate co-evolution of humans and our corresponding microbiota during our coexistence (12, 13). We humans each have a whopping 100 trillion bacterial cells living in our guts, oral cavity, upper airways, skin and, in the case of women, the vagina (6, 14). Many of these species are human-specific and cannot survive for exceedingly long periods outside of our bodies (15-21). Most commonly, our coexistence is mutually beneficial (12). We need them for, among

other things, the harvest of inaccessible nutrients from our diet and for vitamin synthesis (22). In return, we provide a predictable living-environment and act as a reliable nutrient source for them. The human genome encodes approximately 20 000 genes, but the bacterial species that live in symbiosis with us collectively provide an additional 530 000 unique genes to our biological system, many of which are believed to aid us in metabolism as mentioned (23). Indeed, some philosophically-minded scientists argue that we are multispecies "supra-organisms": part human, part microbial (22). New evidence indicating that the gut microbiota can alter human behaviour lends further merit to this notion (24, 25).

The life cycle of host-living bacteria consists of (i) reaching a host, (ii) establishing colonization and (iii) spreading to a new host. Colonization, that is adherence to host tissue and on site proliferation, all while withstanding the external stresses of that particular niche, is thus an absolutely central theme in host-microbe interactions. This interplay can be mutualistic, commensalistic, or parasitic, and depending on the bacterial species involved, may fall under multiple categories at various time points during its sojourn with us.

To colonize a healthy human host is not an easy feat for microbes. Truly, most bacteria cannot. There are nutrient limitations, potential pH-fluctuations, intramicrobial competition and the host's structural barriers. Prokaryotes that manage to colonize the host despite these factors, without inflicting substantial injury or eliciting an inflammatory response, are components of the normal flora. However, it is a well-known fact that coexistence with our microbial selves is not always frictionfree. Bacteria that disrupt the human homeostasis and cause disease during colonization, proliferation or transmission, *e.g.* are pathogenic, are the subjects of study for medical microbiologists. If colonization of the host is hard to begin with, causing disease is an order of magnitude harder as a major additional factor kicks in: the host immunity. Any bacterium that colonizes humans to subsequently cause disease has to be able to cope with all these factors. And they do.

The topic of this thesis is how one outstanding bacterial species, *Haemophilus influenzae*, overcomes the structural barriers of the host by adhering to tissue it "shouldn't" be able to adhere to, and then circumvents the host innate immunity to survive, colonize, proliferate and finally transmit to a new host before the old host has eradicated it. By identifying the molecular factors involved in this scenic drama, we could show that it is possible to "turn the table" on the pathogen, using its own colonization factors against it and stopping it in its tracks before it can establish disease.

The Human Upper Airway

The human respiratory tract is directly subjected to the unsterile and particulate external environment. We breathe 6-12 litres of air/minute, and with inhalation follow chemicals, allergens, microparticles and, naturally, microbes (26). The upper airways fulfil the function of cleaning, moistening and warming up the air before it enters the (mostly) sterile lower airways (pulmonary system) distally of the bronchi, where gas exchange and respiration occurs. Figure 1 shows a schematic representation of the upper airways that is anatomically divided in the nasal cavity, pharynx, larynx, trachea and initial bronchia (26, 27).

The upper airways are lined with a ciliated mucosal epithelium (27). This respiratory epithelium is normally colonized by a specialized microbiota constituted primarily of bacteria from the phyla *Firmicutes, Actinobacteria, Bacteroidetes* and *Proteobacteria* (28, 29). The nasopharynx, which is the primary site of colonization of *H. influenzae*, is a subcompartment of the upper airways situated dorsally of the nasal cavity and superiorly to the oropharynx. Adjacent to the nasopharynx is the entrance to the auditory tube (also known as the Eustachian tube). The route of pathogen ascension to the middle ear starts in the nasopharynx and goes via the Eustachian tube (30).



Figure 1. Schematic representation of the upper airways. The gross anatomical division of the upper airways can be observed above; air enters through the nasal cavity, continues through the naso- and oro- and laryngopharynx further down to the trachea and finally down to the lower airways. *H. influenzae* colonizes the nasopharynx, from which it can enter the autitory tube to cause otitis media, or descend down to the lower respiratory tract to produce other mucosal infections. Image used with permission (31).

The Mucociliary Elevator

The respiratory epithelium is constituted by three major cell types that jointly contribute to the physiological function of the tissue; ciliated, goblet (or mucus), and basal cells (Figure 2). The basal cells are progenitor stem cells that can differentiate into the other cell types present in the epithelium (26). They also regulate the inflammatory response and mediate attachment and communication with the underlying basement membrane. Goblet cells produce electron-lucent granules containing the acidic glycoproteins mucins that are secreted into the lumen to generate the protective mucus layer. This layer has multiple functions, including physically separating the epithelium from toxic compounds or invading microbes as well as trapping microparticles (such as infectious agents) in its mucoid consistency (26). Finally, the function of the ciliated cells is the constant removal of "dirty" mucus to the pharynx by coordinated beating of the cilia. Once the particulatecontaining mucus arrives in the pharynx, it is discarded by deglutition to the gastric tract or expulsion from the mouth (26). Collectively, this system is called the mucociliary elevator and is our first-line of defence against invading microbes in the respiratory tract.



Figure 2. Schematic representation of the mucosal epithelium. The beating cilia move particulate mucus towards the pharynx for removal to the gastric tract or oral discardation. Image cropped and used with permission (31).

The Basement Membrane and Extracellular Matrix Proteins

The respiratory epithelium rests on a rigid structure designated as the basement membrane (Figure 2). This acellular tissue is comprised of extracellular protein matrices. The two major extracellular matrix (ECM) protein components of the basement membrane are laminins (Ln) and collagens that form separate polymer sheets interconnected via molecules such as nidogen, perlecan and agrin (32-34).

The basement membrane does not merely provide a structural scaffold for epithelial and endothelial cells, but is involved in many dynamic homeostatic processes including embryonic development, cell migration, differentiation, wound healing and cell signalling (33). It is directly connected to epithelial cells via hemodesmosomes as well as an array of dynamic protein-protein connections involving, for instance, basal cell membrane integrins (26). The ECM is therefore *de facto* connected to the cytoskeletons of epithelial cells and can alter their phenotype via complex signalling pathways (32). Patients suffering from deleterious mutations in genes encoding ECM-proteins can develop disorders such as muscular dystrophies, epidermolysis bullosa and glomerular disease (35-37).



Figure 3. Schematic image of the Ln-heterotrimer (left) and histological section detecting Ln in the nasopharyngeal epithelium (right). Left: the isoforms Ln-111 and Ln-332 are shown. The different α , β and γ chains are of various lenghts but come together in a long coiled-coil region in all isoforms. At one end of the molecule, the α -chain C-terminus forms the globular domains known as LG1-5 that are important for interactions with cells. On the right: a histological section showing the distinct association of Ln α 3 (secologically dyed brown) with the healthy adult respiratory epithelium. Image cropped and used with the permission of The Human Protein Atlas (38).

As Ln forms the connective scaffold for all epithelial cells (including the skin), one can picture the human body as perpetually coated in an underlying sheet of Ln from top to bottom. Laminins are large, heterotrimeric glycoproteins that once secreted autopolymerize extracellularly to form large matrices on the basal side of the cell-

linings (32). Comprising three separate chains designated as the α -, β - and γ -chain, the Ln monomer forms a large cruciform molecule with specific biological activities assigned to specific domains (Figure 3) (32). Moreover, there are five α -, four β - and three γ -chain variants encoded by separate genes in humans. These are combined in diverse constellations that form 16 different heterotrimeric isoforms. The nomenclature for Ln is based on the chain composition; the isoform consisting of $\alpha 3$, β 3 and γ 2 thus becomes Ln-332 (39). The different isoforms have distinct tropisms and plausibly functions; Ln-332 is the primary isoform in the basement membranes whereas Ln-211 is the primary isoform in muscle tissue, for instance. Recently, the Protein Atlas (http://www.proteinatlas.org/) has serologically and Human transcriptionally showed the specific presence of the majority of Ln chains in a large variety of organ systems. Judging by the detection of singular chains in the upper airways, the prevalent isoforms in this niche appear to be Ln-111, Ln-332 and Ln-411. These findings allow the future use of anatomically relevant isoforms in the field of Ln research.

If the epithelial integrity is compromised, the basement membrane becomes exposed to the lumen. In this occurrence, ECM-proteins are attractive targets for adherence by pathogens as a mean to circumvent the mucociliary elevator. Moreover, any microbes that disseminates past the epithelium need to cross this structure. As a testament to the importance of the basement membrane in host-pathogen interactions, microbes from every biological kingdom have been shown to bind to, and in some instances degrade, ECM-proteins (34, 40). This topic will be further discussed in the following chapters.

The Innate Immunity in The Airways

The second line of human defence against invading microbes is the innate immunity. The airway epithelial cells have a range of pathogen pattern recognition receptors (PRRs) that detect conserved microbial motifs known as pathogen-associated molecular patterns (PAMPs) including lipopolysaccharides (LPS), peptidoglycan, viral/bacterial nucleic acids and lipoproteins (41). When PRRs, such as the membrane bound Toll-like receptors (TLRs) or the intracellular nucleotide-binding oligomerization domain (NOD)-like receptors (NLRs), are activated by pathogen components, innate immune responses are induced. The result is the secretion of antimicrobial agents such as lysozymes, antimicrobial peptides (AMPs), interferons, lactoferrin and nitric oxide. Another consequence is the release of cytokines, which mediate inflammation and recruitment of phagocytes to the site of pathogen detection (41, 42). Moreover, inflammation causes augmented plasma exudation to the mucosa due to increased airway epithelial permeability, resulting in the

heightened presence of complement factors (43). Together, these innate immune responses exert a potent antimicrobial activity (44).

As one of the focuses of this thesis is the *H. influenzae* interaction with the complement system, this feature of the innate immunity will be discussed in greater detail.

The Function and Regulation of the Complement System

The complement system is an organized network comprising more than 30 (mainly serum) proteins whose chief function is to prevent microbial invasion (45, 46). This arm of the innate immunity exerts its effect in three steps: recognition of microbial non-self, opsonisation of the microbe, and in the case of Gram-negatives, osmolytic destruction of the bacterial cell via insertion of the pore-forming membrane attack complex (MAC) in the bacterial outer membrane (45, 46).

In many aspects, the central component of the complement is the C3 protein. Once cleaved, it yields the chemoattractant C3a and highly potent opsonin C3b (47). The function of C3b is two-fold; opsonize bacteria for neutrophil and macrophage phagocytosis and launch the terminal pathway of the complement system.

The mechanisms of C3 cleavage are well understood. Three separate and distinct pathways can lead to activation of the terminal pathway (47). The classical pathway is dependant on bactericidal IgG and IgM antibody-recognition of microbes. C1q recognition of two proximally close Fc-regions (indicating antibody opsonisation of a particle) activates a cascade that results in the formation of the C3-convertase C4b2a. The second route, called the lectin pathway, is activated by the recognition of mannose or N-acetylglucosamine on bacterial surfaces by mannose-binding lectin. Mannose-binding lectin-associated serine proteases are consequently activated and cleave C4 and C2 to yield the C3-convertase C4b2a. Finally, the alternative pathway is induced by the spontaneous hydrolysis of C3 and the subsequent attachment of C3b to the surface of cells (45).

Once C3 undergoes hydrolyzation, and the subcomponent C3b attaches to the nearby surfaces, the terminal pathway is induced (47, 48). In sequential interactions of the components C3b, C5, C6, C7 and C8, the pore-forming polymerization of C9 will finally result in MAC-formation that lyses the bacterial cells. Bacterial species that colonize the mucosal airway epithelium, where constant plasma exudation occurs, have to address the complement system proficiently in order to resist rapid clearance.

From the host's point of view, it is absolutely imperative to prevent complementmediated autoimmunity (48). Therefore, the system is stringently regulated on almost every level. Complement down-regulatory proteins include C1-inhibitor, C4bbinding protein, clusterin, Factor H, Factor H-like protein 1, Factor I, properdin and vitronectin (47). It has long been reported that bacterial pathogens sequester these soluble complement inhibitors on their surface as a mean of evading the innate immunity (49).

Vitronectin.

Vitronectin is a multifunctional glycoprotein present as a monomer in serum and as polymers in the ECM. Comprising 478 amino acids, it has domains with homology to the proteins somatomedin-B and haemopexin as well as reported binding sites for heparin, urokinase plasminogen activator-urokinase plasminogen activator receptor complex and integrins (50). Equipped with this vast set of tools, vitronectin is involved in a series of homeostatic processes that include cell migration, woundhealing, angiogenesis and complement regulation (51, 52). So far, vitronectin deficiency has not been reported in humans, but genetically engineered vitronectinnull (Vn(-/-)) mice have been shown to suffer from fibrinolytic imbalances and decreased microvascular angiogenesis (53, 54).

As an ECM-component, vitronectin safeguards the surrounding tissue against autoimmune complement activation. The glycoprotein inhibits the terminal pathway at two distinct steps; C5b-C7 formation and C9 polymerization, thus effectively preventing MAC-formation (52). Importantly, vitronectin is present in the airway lumen of healthy individuals, and at elevated levels in patients suffering from respiratory illnesses, as detected by bronchoalveolar lavage (55, 56). These two facts make soluble vitronectin an attractive ligand for bacteria. Indeed, respiratory tract pathogens such as *H. influenzae, Moraxella catarrhalis, Pseudomonas aeruginosa, Streptococcus pneumoniae, S. pyogenes* and *Staphylococcus aureus* have been reported to interact with vitronectin (50). Many of these species also use vitronectin as a mean of adherence to the host. Vitronectin is thus an important molecule in the host-microbe interactions in the upper airways.

Haemophilus influenzae – Expert Colonizer of the Human Upper Airways

A Commensal and An Opportunist

Haemophilus influenzae is a Gram-negative, facultative anaerobic small coccobacillus belonging to the *Pasteurellaceae* family of the Gammaproteobacteria class (57). The organism is well-known to microbiologists as it has been involved in several important scientific breakthroughs: it was the source of discovery of restriction enzymes (58), it was the first free-living organism to have its genome fully sequenced (59), and the first licensed glycoconjugate vaccine for use in humans was developed against the polysaccharide capsule of the serotype b (Hib) strains of the species (60).

The organism was first discovered in 1892 by Pfeiffer and mistakenly identified as the causative agent of influenza-disease, hence the species name (originally designated as *Bacillus influenzae*, or "Pfeiffer's bacillus") (61). The pathogen did not hold up to Koch's postulate as the aetiology of influenza and was eventually disproved as such (61). It is now known that *H. influenzae* is a cause of secondary bacterial infections after prior influenza virus-mediated disease (62), explaining the initial misidentification considering that viruses were not well-known at the time. In the 1930's, the bacteriologist Margaret Pittman conducted many pioneering studies on the organism, including the identification of encapsulated (divided into serotypes a-f depending on the biochemical features of the polyribosylribitol phosphate capsule that affects agglutination with antiserum) and non-capsulated strains, the latter designated nontypeable *H. influenzae* (NTHi) (63, 64).

H. influenzae is a fastidious and highly adapted human-specific commensal (20). So much so, that it has lost the ability to biosynthesize the essential co-factor molecules heme and nicotinamide adenine dinucleotide (NAD) (65-67), relying strictly on scavenging them from the environment in our bodies. This heme- and NAD-deficient phenotype is so rare that it has historically been used for clinical diagnostics of the species (65, 67). Clinical samples would be plated out together with a known "feeder" species that could provide these co-factors, and growth of "satellite colonies" in the immediate vicinity of the streak of "feeder" strain was an indication of *H. influenzae* (68).

H. influenzae colonization of humans, especially toddlers, is transient with a turnover rate of approximately three months (69). As the host clears the predominant residing strain, a successor strain replaces it in the available niche. The intermittent carriage rate of NTHi has been observed to be 32-77% in healthy children at any given time point (70-73). The earliest documented cases of carriage are from four Medieval German adult skeletons dated to 950-1200 A.D. (74). The dental plaque of these

individuals, when analysed with ancient-DNA technology, revealed concomitant carriage of capsulated and non-capsulated strains in the same individuals (74). Historically, the human nasopharynx has been regarded as the primary site of initial colonization, but recent reports show a prevalent presence of the species in the buccal mucosa of the oral cavity as well, lending additional merit to the dental plaque findings (6).

While residing in the upper airways and oral cavity, *H. influenzae* can proceed to cause mucosal infections including otitis media (OM), conjunctivitis, bronchitis and sinusitis if the host's stringent structural barriers and/or immune system is compromised by a preceding viral infection or prolonged episodes of inflammation (75, 76). Occasionally, further dissemination resulting in invasive disease such as meningitis and sepsis is manifested primarily (but not exclusively) by strains belonging to the capsulated serotype b clade (77-80).

Since the introduction of a capsular polysaccharide glycoconjugated vaccine against Hib, which practically eradicated Hib-mediated disease in the places vaccineprograms were employed, researchers have redirected their efforts to NTHi that is currently the most clinically relevant subgroup (77). The rest of this work, including the present investigation, will thus focus on the non-capsulated NTHi.

Genetic Heterogeneity

Margaret Pittman wrote this catching paragraph about *H. influenzae* as early as 1931:

"The bacteria of this group do not form a well characterized bacterial species, and it has long been recognized that individual strains differ from one another in morphology and virulence, in the appearance of the colonies which they form, in their ability to form indole, in power to ferment sugar and to induce hemolysis, in their immunological reactions, and even in their requirements for growth inartificial cultures."(63)

One hallmark of *H. influenzae* is its genetic (and subsequently phenotypic) diversity. This fact is mirrored by the difference in the number of annotated genes in whole genome sequenced strains, ranging from 1688-2355 open reading frames (ORFs) per strain. In parallel with this, the genome sizes vary between 1.79-2.01 mega base pairs (http://www.ncbi.nlm.nih.gov/genome/genomes/165). Recently, the "supragenome" of the species has been estimated to contain up to 4500 unique ORFs, while the core set of genes existing in all strains amounts to only 1485 of said ORFs (81). This means that 2/3 of the "supragenome" of *H. influenzae* is perpetually distributed among countless strains. According to the "Distributed Genome Hypothesis", naturally competent *H. influenzae* strains undergo continuous and dynamic genetic recombination processes during simultaneous colonization of the same host (82, 83). As the organism exclusively takes up intraspecies DNA (under normal circumstance),

naturally competent strains would "have access" to a wide variety of genes throughout time, allowing them to evolve with an altering environment without the burden of maintaining a large genome, thus gaining a fitness advantage (82, 83).

As a parenthesis, it can be noted that the introduction of the capsule biosynthesis loci has seemingly occurred twice throughout evolution in the NTHi background (plausibly via a rare horizontal gene transfer from other Gram-negative bacterial species including *Actinobacillus* spp., *Neisseria meningitides* and *Mannheimia haemolytica* (84)), as capsulated types can be grouped in two distinct lineages (85). Because the capsulated strains are thus largely clonal, the greatest genetic diversity is observed in the NTHi (85).

Due to this high intraspecies variation, researchers have invested substantial efforts on identifying genes and/or strains that are exclusively associated with commensalism and pathogenicity. Recent studies have managed to identify such genetic elements via genome array analysis of a wide range of clinical isolates and avirulent commensal strains from a broad geographical collection (81). A curious finding of this investigation was that the 28 genes identified as primarily pathogen-associated were not any of the already known and well-studied virulence factors such as adhesins or immune evasion factors, but comprised mostly uncharacterized proteins. It remains highly interesting to study the functions of these disease-associated genes.

Finally, it also has to be noted that the genetic heterogeneity of the species has implications for vaccine development, a subject that will receive more attention in a subsequent chapter.

Bacterial Factors Involved in Colonization and Virulence

As previously stated, colonization of the host is a central aspect of any host-dependant bacterial life. In medical microbiology, colonization is commonly equated to pathogenesis. As this may be true in some cases, and while conceding that colonization is a prerequisite for any following episodes of illness, it is certainly not true for all. Respiratory tract illnesses, for instance, originate in the vast majority of cases from bacteria belonging to the normal flora. These opportunistic pathogens that most often only cause asymptomatic colonization include *H. influenzae*, *S. pneumoniae*, *M. catarrhalis*, *S. aureus* and *S. pyogenes* (86-88). However, immature immune systems, defective structural barriers, primary or acquired immune deficiencies, prior viral infections and chronic inflammation may alter the conditions in this microecology, allowing otherwise commensal bacteria to take over and go rogue (89-93). But in order to do so, opportunistic pathogens need to be able to address several obstacles in the process. The *H. influenzae* factors involved in colonization and virulence are briefly reviewed here.

Adherence to Host Tissue

The first step of colonization when arriving in a new host is avoiding mechanical clearance by the mucociliary elevator. At this crucial stage, NTHi induces a decrease in (or arrest of) the ciliary beating, adheres to host cells or the surrounding tissue and forms microcolonies on the epithelium (20, 94, 95).

The importance of the ciliated cells for clearance of *H. influenzae* can be appreciated by reports regarding patients suffering from primary ciliary dyskinesia; *H. influenzae* is the predominant pathogen found in the sputum of these patients (94, 96). *Haemophilus* Protein D is a glycerol-3-phosphodiester phosphodiesterase surfaceexposed lipoprotein (97). It has previously been shown to induce both a decreased ciliary beating frequency as well as detachment of ciliated cells in primary polarized epithelial cells (98), displaying one mechanism by which NTHi compromises the mucociliary epithelium. Other investigations have implicated NTHi-mediated activation of host protein kinase C epsilon in the process of decreased ciliary beating (99).

The Gram-negative endotoxin, also known as LPS or lipooligosaccharide (LOS) in some species including H. influenzae, is a well-characterized virulence factor and potent inducer of TLR-4-dependant inflammation (100, 101). A lipid-A-tethered outer membrane structure, it consists of a triheptose-3-deoxy-D-manno-octulosonic acid backbone that branches off with highly varied subsets of oligosaccharides (101). The LOS has been alternately implicated and refuted in the involvement of H. influenzae-mediated decrease of ciliary beating frequency (102-105). However, as the LOS structure has been shown to be highly variable in NTHi (101, 106), these results need not to be mutually exclusive. NTHi LOS has also been observed to directly bind to epithelial cells, thus potentiating bacterial adherence to host tissue (107, 108). Interestingly, in an experimental human nasopharyngeal colonization model, significant changes were only observed in two out of 16 investigated phase-variable genes (109). One encoded the IgA1-protease and the other was licA, whose gene product incorporates exogenously acquired phosphorylcholine into the LOS to increase the adherence to epithelial cells. The in vivo selection for LicA-producing subpopulations indicates that this mechanism is very important for host colonization.

Type IV pilus, a multicomponent, long filamentous surface structure commonly associated with bacterial conjugation, has also been shown to contribute to NTHi adherence to epithelial cells (110-112). Moreover, there is a number of proteinous adhesins imperative for the direct mucin- or host cell-binding of NTHi, including the autotransporters Hap (113, 114) and Hia (115), lipoproteins *Haemophilus* Protein D (97) and *Haemophilus* Protein E (116), P2 porin (117), outer membrane protein P5 (117-119) and the high molecular weight (HMW) adhesins (120). These proteins belong to vastly different protein families with little, if any, structural resemblance, indicating a highly multifactorial mechanism for the NTHi adherence to the

respiratory tract epithelium. Importantly, it has been frequently observed that the pathogen adheres to sites of epithelial damage, forming microcolonies in places where the cell confluency has been compromised (94, 98, 121). As the basement membrane on the basal side of the epithelium becomes exposed to the lumen during damage, it is highly plausible that bacterial interaction with the host ECM-proteins is imperative for pathogenesis. Several of the adhesins mentioned, including Hap and PE, have been shown to interact with ECM proteins such as Ln, collagen IV, fibronectin and vitronectin (122-125).

There are also conflicting reports about the occurrence of epithelial cell invasion. Some argue it is a pathogen-beneficial mechanism, others claim it to be hostbeneficial (resulting in bacterial death) and yet a third group states it is not occurring at all (126-129). This particular mechanism needs further evaluation before consensus can be reached.

Biofilm-formation

Once adherence to the host tissue is established, many NTHi strains have the capacity to produce biofilms in particular anatomical locations (130). This colonization mechanism is also observed in other otopathogens including *M. catarrhalis* and *S. pneumoniae* (131). Biofilms are highly structured microbial communities consisting of bacterial cells embedded in extracellular protein-, DNA- and polysaccharide-matrices (132). "Fortified" biofilm-associated bacteria display much increased resistance to biological, chemical and physical environmental stresses (including antibiotics and the host immune system) as compared to planktonic microbes, and are believed to be the cause of persistent NTHi infections (132, 133). Indeed, it has been proposed that the majority of prokaryotic life on earth exists in biofilms rather than as dispersed singular cells (134).

Many bacterial factors have been implicated in NTHi biofilm-formation, including double stranded DNA and the previously discussed type IV pili and LOS (111, 135-137). Recently, the entire proteinous content of the extracellular material of the NTHi biofilm was mapped (138). Eighteen proteins, including P2 and P5, were shown to comprise the biofilm-specific extracellular proteome. Bacterial DNA as well as proteins from the cytoplasm, periplasm and the outer membrane were all reported to be present in the biofilm, a fact the authors accredit to bacterial cell lysis being integral to the origination of the community (138). Finally, it should also be noted that several quorum signalling pathways involved in the induction of NTHi biofilm-formation have been elucidated (139, 140).

From a clinical perspective, NTHi biofilms have primarily been implicated in (recurrent) OM with effusion and exacerbations of chronic obstructive pulmonary disease (COPD) (130). It is very plausible that biofilm-associated NTHi can cause chronic infections, as these communities are difficult to eradicate with conventional

antibiotics. Interestingly, one study showed that targeting key factors of the extracellular components of the NTHi biofilm for destruction would collapse the entire biofilm organization and increase clearance of the pathogen from the middle ear of chinchillas (141). Continued research of this persistent mode of bacterial colonization will be highly interesting to follow.

Host Immune Evasion

It is vital for bacteria living in the human body to either avoid, or circumvent, the host immune system during colonization. Investigators have shown how NTHi effectively addresses innate immune effectors such as AMPs, transferrin and nitric oxide (142-145). Moreover, the highly substrate-specific IgA1-protease of NTHi has epidemiologically been implicated in virulence (146). As secretory IgA1 is the main immunoglobulin present in the human airway mucosa, the IgA1-protease may facilitate NTHi persistence in the niche by hydrolysing the main effector of the acquired immunity in the respiratory tract (147). This mechanism has been shown *in vivo* for its upper airways coinhabitant *S. pneumoniae* (148).

Protection against phagocytosis is mediated by the capsule in serotypable strains (149), but recently it was shown that NTHi can produce a "pseudo-capsule" via extensive branching of the LOS that blocks IgM from binding to bacterial surface epitopes (150). Subsequently, neutrophil-mediated killing of NTHi was significantly impaired, providing one mechanistic explanation as to why the excessive presence of neutrophils in the inflamed middle ear does not clear NTHi infections.

The host humoral response has also been shown to be unspecifically activated via a yet unknown factor in NTHi outer membrane vesicles (OMVs) in a "super antigen"-like manner (151). The proliferating lymphocytes would produce antibodies that did not recognize NTHi, resulting in the diversion of the human adaptive immunity.

Evasion of complement-mediated killing is also of utter importance for NTHi colonization and virulence. The evidence shows that NTHi employs two distinct strategies for complement-resistance: i) blocking antibodies and complement components from reaching the surface via LOS-branching and ii) surface sequestration of complement down-regulatory host proteins.

It has been shown that NTHi incorporates host-derived molecules such as sialic acid and phosphorylcholine into its LOS as a mean of camouflage (100, 152). Moreover, alterations of surface glycans have shown to be important in preventing bactericidal antibodies from opsonizing NTHi, thereby preventing complement activation via the classical pathway (150, 153-155). In addition to these, LgtC-mediated modification of the LOS has been shown to delay C4b deposition on the bacterial surface via an unclear mechanism (156, 157). As a testament to their importance, several of these processes have been shown to be activated during exposure to normal human serum (153-155), a scenario that is likely to occur *in vivo* during inflammation-mediated plasma exudation in the airway epithelium.

Nontypeable *H. influenzae* has also been shown to acquire host-derived complement inhibitors, such as C4BP, Factor H and vitronectin, to its surface (158-162). The outer membrane proteins P5 and Protein E have been shown to be important for Factor H- and vitronectin-binding, respectively, whereas the C4BP-binding factor of NTHi has remained elusive. In all instances, it has been shown that NTHi evades complement-mediated killing by these sequestration events, thus increasing its serum-resistance.

Despite the struggle that the bacteria put forth, each specific NTHi strain is normally cleared from the human host within three months after initial colonization (69). The eventual clearance is likely attributed to an overpowering pressure mounted over time by the host acquired immunity.

Niche Adaptability and Nutrient Acquisition

The nasopharyngeal tract is a nutrient-poor *milieu* for bacteria (163). Microbes that colonize this surface rely on sensing and responding to nutrient-scarcity and external environmental changes by activation of an array of scavenging and adaptation systems. It has even been hypothesized that the DNA transformation machinery in *H. influenzae* was primarily evolved as a nutrient uptake system in this barren niche rather than for genetic recombination purposes (164).

As previously stated, *H. influenzae* has lost the genes for *de novo* biosynthesis of heme and NAD. Exogenous NAD uptake is mediated by the outer membrane lipoprotein P4 and the P2 porin (165-167) before the molecule is further metabolized for utilization downstream in the peri- and cytoplasm (168, 169). Porin P2 is the most abundant protein in the outer membrane of NTHi (170), plausibly contributing significantly to maintaining a steady uptake of exogenous NAD. As no specific regulation element has been reported for either P4 or P2, the NAD uptake system is likely perpetually switched on.

This is in stark contrast to transport systems involved in the uptake and metabolism of heme and iron, which are tightly regulated. It was early reported that *H. influenzae* can scavenge human transferrin via an uptake system that is only active in the absence of heme, a scenario that would correlate with the mucosal epithelial environment (171). More recently, the ferric uptake regulator (Fur) regulon of NTHi was shown to contain 73 genes (172), meaning that approximately 4% of the genome was directly regulated by the intracellular Fe²⁺-sensing of Fur. Many of these genes were expectedly involved in iron-utilization but, interestingly, it was shown that iron/heme-independent virulence factors such as the IgA1-protease were also under the regulatory control of Fur. This again suggests that NTHi re-wired several virulence factors to be automatically switched on in iron/heme-depleted environments, *i.e.* in the human nasopharynx.

But the response to iron/heme-deficiency is not exclusively regulated by Fur. The core (identical in a five capsulated and non-capsulated strains) and non-core modulon response to low levels of iron/heme has been mapped *in vitro* and *in vivo* (173). Investigators identified 55 core and 200 non-core ORFs that are up- or down-regulated in the absence of iron/heme, exhibiting how NTHi saves energy by tightly regulating the expression of iron/heme-utilization genes. One factor that has been implicated in sensing heme-availability is the multifunctional Sap-transporter that also recognizes host-derived molecules in the microenvironment and alters the bacterial response accordingly (128, 174).

The urease operon has been shown to be one of the most upregulated genetic elements of NTHi *in vivo* (175, 176). The importance of urease for onset of disease has been demonstrated in several studies, establishing the role of the enzyme in raising the pH in the human respiratory tract microenvironment to facilitate bacterial growth (177, 178). This is in analogy with several other human pathogens in unrelated niches including *Helicobacter pylori* and *Proteus mirabilis*. As human mucins are highly acidic, NTHi is likely dependant on proficient pH-buffering during inflammation and mucus-hypersecretion.

The conditions NTHi encounters in the human nasopharynx and Eustachian tube differ considerably from those in the lower respiratory tract of COPD-patients. The latter environment is characterized by chronic inflammation, excessive presence of phagocytic cells, epithelial damage, failure of the mucociliary elevator, mucus hypersecretion (causing local hypoxia), elevated cvtokine levels and oxidant/antioxidant dysregulation (92). NTHi adapts ably to both these settings, and the genetic elements that allow descension into the bronchia of COPD-patients have been studied (92, 176, 179). Urease, IgA1-protease, anti-oxidants and HMWadhesins are among the factors reported to be upregulated during exposure to this milieu. Moreover, it has been observed that COPD-patient-derived alveolar macrophages are defective in phagocytosis of NTHi but not of *M. catarrhalis* or *S.* pneumoniae (90). These data collectively cast light on the bacterial adaptation and survival in various niches of the human body.

Polymicrobial Interactions

Although NTHi is a proficient colonizer of the human respiratory epithelium, onset of disease most commonly requires prior priming of the host by viral agents (76). For this reason, the polymicrobial interactions that underlie NTHi-virulence are important but somewhat overlooked aspects of pathogenesis.

Clinical evidence shows a prevalent manifestation of viral-NTHi co-infections, with the viruses involved being primarily influenza A, rhinovirus, adenovirus and human

respiratory syncytical virus (180-186). More than 20 years ago, researchers showed how the mucociliary elevator of the middle ear was severely compromised by adenoviral infections for up to 35 days (187). This would surely diminish the host defence against intruding secondary bacterial microbes such as NTHi. Several more molecular mechanisms behind viral facilitation of secondary NTHi-infections have recently been unfolded. Respiratory syncytial virus has been reported to induce downregulation of the host antimicrobial peptide beta-defensin-1, an attenuation of the host immunity that markedly promoted over-growth of NTHi in the nasopharynx (188). Other investigators have shown that rhinovirus attenuated the host TLR-2mediated immune response to NTHi via the adaptor protein IRAK-1 signalling (189). In a third study, researchers reported that if the host was subjected to a primary influenza A virus infection, NTHi could survive in the pulmonary system of mice without many of the genes normally required for stress responses in vivo (190). They also showed that NTHi needed to address excessive oxidative stress brought on by the influenza A virus during co-infection, a condition that the pathogen adapted to by activating the oxidative stress response modulator *iscR*. These studies exemplify molecular mechanisms by which viral infections render the host highly susceptible to secondary NTHi infections and how the bacteria can exploit and adapt to various microenvironments in situ.

But intermicrobial cooperation is not restricted to interkingdom partners. Indeed, NTHi is often isolated with other bacterial otopathogens, including *M. catarrhalis* and *S. pneumoniae*, in clinical settings (180, 191-195). Mixed species experiments have shown that NTHi quorum signalling can induce the biofilm-formation of *M. catarrhalis* and *S. pneumoniae* (140, 196, 197). Other reports suggest that mixed NTHi-*S. pneumoniae* OM are unique clinical entities, separate from single pathogen infections with regard to pneumococcal serotype distribution, suggesting that specific interspecies strain-strain interactions facilitate virulence (198). The mechanisms behind these synergistic infections remains to be more fully elucidated.

Finally, the highly interesting outer membrane vesiculation systems have been shown to play important roles for interspecies cooperation, conferring antibiotic resistance and serum resistance to bacterial coinhabitants in the same niche (199-201). The dispersion of OMVs in the environment may mediate many more polymicrobial interactions, including long-range signalling and nutrient allocation.

Transmission

The molecular mechanisms of bacterial transmission between hosts are the least studied aspect of the cycle of colonization and spread. Indeed, doing a literature search revealed that there are no such investigations published regarding NTHi. However, one recent publication revealed the mechanistic potentiation of *S. pneumoniae* spread by a concomitant influenza A superinfection (202). Authors showed that influenza A virus-mediated mucus-hypersecretion facilitated shedding of

the bacterial pathogen in an infant mouse model. Moreover, they showed that TLR-2 activation decreases this effect, as transmission of *S. pneumoniae* was much increased in *tlr2-l*- mice. Interestingly, a strong bottleneck effect was observed during transmission: colonization was established by a very small number of bacterial cells that transmitted to the new host. As. *S. pneumoniae* and NTHi colonize the same niche and have very similar clinical manifestations, it is plausible that respiratory tract viruses may facilitate the shedding of NTHi in the same manner. However, this remains to be experimentally addressed.

It is known that *H. influenzae* can survive for several days on inanimate surfaces (203). This fact, taken together with the extensive presence of the species in saliva and nasal mucus (204, 205), could potentially explain one route of transmission in, for instance, children's day care centres.

It will be interesting to follow the research in this field to see if additional bacterial factors involved in ensuring shedding from biofilm and transmission to a new host can be identified at the molecular level.

H. influenzae-Mediated Disease

As mentioned, NTHi can cause a range of mucosal infections when given the opportunity. These include OM, conjunctivitis, sinusitis, bronchitis and exacerbations of COPD (75). Moreover, NTHi is observed to occasionally disseminate and cause invasive infections (78, 79). This work will focus on the two most prevalent NTHi-mediated diseases; OM and COPD exacerbations.

Otitis Media

Otitis media, or middle ear inflammation, is a common childhood disease. In large cohort studies, it has been shown that up to 49%-80% of children experience one or several episodes of OM before the age of 3 years (206, 207). The incidence rate has been estimated to an astonishing 709 million cases/year globally (208). In Sweden alone, the incidence of acute OM (AOM) is 200 000 cases per year (209). Otitis media is one the most common reasons, if not the single most common cause, of children seeking medical attention, experiencing hearing loss, receiving antibiotic treatment, and undergoing surgery (41). The socio-economic burden of OM is therefore of considerable significance.

Aetiological agents of OM can be viral, bacterial, or both simultaneously (42). The most common viral agents are respiratory syncytial virus and adenovirus, whereas the most common bacterial agents are *S. pneumoniae*, NTHi and *M. catarrhalis*. It has been repeatedly shown that viral upper respiratory tract infections precede episodes of OM, highlighting the microbial interplay and complexity of disease onset (210), and the molecular mechanisms behind these phenomena are beginning to unravel as previously discussed.

The reason for OM being principally a childhood disease is partly anatomical (Figure 4). The Eustachian tube (also called auditory tube) is shorter and the angle is horizontal in the small, round skull of toddlers, potentiating microbe ascension from the nasopharynx to the middle ear (211). Moreover, the muscles that control the entry to the Eustachian tube are not well-developed in infants. As children reach 4-5 years of age, their skulls elongate, the entry tightens and the Eustachian tube takes on the conformation of later life. Consequently, a marked drop in OM incidence rate is observed between the age groups 1-3 years and 4-6 years (212).

Once pathogens reach the middle ear and proliferate, inflammatory responses will be induced via the host's PRRs such the TLRs and NLRs (41). At this point, inflowing fluids and mucus that clog hearing and prompt otalgia (due to the sheer pressure) will fill and be trapped in the normally gaseous tympanic cavity. In rare instances, the tympanic membrane might succumb to the internal pressure and burst. The clinical symptoms of AOM include fever, otalgia, malaise and occasionally otorrhea (213).



Figure 4. The anatomy of the middle ear. The Eustachian tube (also called autitory tube) seals of the tympanic cavity from the nasopharynx. However, should microbes be able to invade the middle ear from the nasopharyngeal tract, they cause a strong inflammatory reponse resulting in OM. Image used with permission (31).

Most patients can clear the acute infection without treatment, but many require medical attention including antibiotic therapy (209). In some cases, OM can become chronic, resulting in recurrent episodes of middle ear inflammation after the completion of each antibiotic treatment, often caused by the same pathogen. Biofilm-associated NTHi have been implicated in this disease, as NTHi is the primary aetiology for recurrent OM (130, 213, 214). Nontypeable *H. influenzae* is also the reported cause of 55-95% of bacterial AOM in children (75). Considering the extremely high global incidence rate of OM, NTHi is quite plausibly one of the most common bacterial pathogens of man.

Exacerbation of COPD

Chronic obstructive pulmonary disease occurs with a high incidence rate in the world, affecting 5-10% of the population (215). It is estimated that 400 000-700 000 patients suffer from COPD in Sweden (216) and 15.9% of all hospital admissions in the UK are reported to be COPD-related (217). This chronic condition is also associated with a high degree of morbidity and mortality, being the third leading cause of death in USA (218). Collectively, the socio-economical burden of COPD is, in analogy with OM, of striking scope.

Patients suffering from COPD are diagnosed with an irreversible chronic inflammation of the lower airways. The clinical symptoms of COPD include obstruction of breathing, excessive coughing, fatigue, chest tightness and susceptibility to respiratory infections (219). The number one risk factor of developing COPD is cigarette smoking, albeit subjection to air pollution, occupational exposure to dust, gases and fumes as well as genetic factors (such as α 1-antitrypsin deficiency) have been implicated in disease onset (215).

As the pathophysiology of the disease includes the displacement of the mucociliary elevator, the airways of COPD-patients are intermittently colonized by opportunistic pathogens such as NTHi, *S. pneumoniae*, *M. catarrhalis* and *P. aeruginosa* (220). These bacterial species, as well as a series of viruses such as rhinovirus, parainfluenza virus, influenza virus and respiratory syncytical virus, cause acute exacerbations of COPD (218). Exacerbation is associated with abnormal and sudden worsening of respiratory symptoms, including increased dyspnea, cough, sputum purulence and general symptoms of common cold (221). Pathogen-mediated exacerbations are induced by the activation of the host's PRRs after the acquisition of a new strain and the severity of single episodes varies from mild to life threatening (222, 223). NTHi is the primary aetiology of COPD exacerbation, causing approximately 20-30% of the cases (217, 218). Frequent exacerbations have been shown to strongly affect disease progression (including the decline of lung functionality as well as the mortality rate) and often require antibiotic treatment (221).

Investigators studying the pulmonary physiology of COPD-patients colonized by NTHi have made some highly interesting observations. One study showed that in addition to NTHi-specific IgA being significantly lower in NTHi-colonized patients compared to non-colonized patients, a fact that can plausibly be attributed to the NTHi IgA1-protease, the presence of host metalloproteinase-9 (MMP-9) was in average approximately three times higher in the sputum of individuals colonized by the microbe (224). This was true for both the pro- and active form of the enzyme. Moreover, the ratio of the tissue inhibitor of metalloproteinase-1 (TIMP-1) to MMP-9 was 10 times lower in patients colonized by NTHi, suggesting a high activity of MMP-9 *in situ*. As MMP-9 is involved in ECM-rearrangement, the researchers hypothesized that "bronchial colonization by *H. influenzae* may cause structural

changes in the ECM". These findings again suggest that pathogen interaction with the ECM-proteins may be highly important for bacterial pathogenesis.

Other studies have discovered that alveolar macrophages derived from COPDpatients are defective in NTHi phagocytosis. The mechanism behind this dysfunction is currently unclear (90).

Finally, investigators have reported that the presence of AMPs are significantly lower in COPD patients colonized by, or experiencing exacerbations mediated by, NTHi (220). These observations collectively cast light on selected aspects of NTHi-mediated exacerbations of COPD and open the door for evaluation, and potentially inhibition, of the molecular mechanisms behind pathogenesis.

Vaccine Development Against Nontypeable H. influenzae

The first commercial H. influenzae vaccines were not the glycoconjugate Hibvaccines, but a series of vaccines launched in the US during the Spanish flu pandemic of 1918-1919 when H. influenzae was still believed to be the aetiological agent of influenza (61, 225). As pressure was mounting on authorities to provide a treatment for the disease, various medical institutions launched at least five different vaccines based on isolated strains of "Bacillus influenzae" from the airways of influenza patients. This took place more than a decade before Pittman's discovery of capsulated and non-capsulated strains in 1931, so there are no details on what serotypes/NTHi strains were used. However, based on current understanding of epidemiology, they were most probably Hib and perhaps a number of NTHi strains. In any event, most of the vaccines were merely heat-inactivated bacterial preparations that were injected into patients. Several hundreds of thousands of adults were immunized with these heat-denatured H. influenzae lysates. Curiously enough, high efficacies in influenzadisease prevention were unanimously reported for them (61). This could indicate that i) heat-inactivated and adjuvant supplemented bacterial preparations induced a potent immune response in adults (to perhaps the Hib-capsule antigen) and ii) secondary H. influenzae infection following the influenzae A virus infection played a large role in the high mortality observed during the 1918-1919 pandemic.

With the repudiation of *H. influenzae* as the causative agent of influenza, the use of these vaccines were soon discontinued and forgotten (61). However, in the 1990s, with a rise in interest in the medical community to address prevention of bacterial meningitis in children, the protein-conjugated capsular-polysaccharide vaccine against Hib was licenced as the first glycoconjugate vaccine for use in humans (60). This vaccine provided a T-cell-dependant memory cell-inducing protection in toddlers, something that could not be achieved with a capsular polysaccharide-antigen alone.

A highly efficient vaccine, Hib-mediated infections were practically eliminated in the wake of its introduction in national immunization programs (226). In should be noted, however, that Hib-disease is currently still a major problem in areas that have not yet employed the vaccine on a large population scale (227).

Since the achievement of developing a Hib-vaccine, the focus in *H. influenzae* vaccinology has shifted to NTHi. The non-capsulated strains provide a vastly different scientific challenge than Hib due the absence of a singular main surface antigen, which in the case of Hib was the capsule. Moreover, as previously discussed, NTHi strains are highly heterogeneous, making the identification of suitable antigens a partly epidemiological task. In addition to ubiquity, promising antigens need to display conservedness, cell surface-exposure and antigenicity. Currently, there are no licensed NTHi vaccines.

Antigen(s)	Animal Model	Outcome of Immunization	Reference
Hap	Mouse nasopharyngeal colonization	Significantly reduced colonization	(228, 229)
Integration Host Factor (DNABII protein)	Chinchilla middle ear biofilm clearance	Significantly enhanced clearance	(230)
LOS	Chinchilla middle ear and nasopharyngeal clearance	Significantly reduced colonization	(231)
Omp26	Rat pulmonary clearance	Significantly enhanced clearance	(232)
Outer membran vesicles	Mouse nasopharyngeal colonization clearance	Significantly reduced colonization	(233)
P4	Mouse nasopharyngeal clearance	Significantly enhanced clearance	(234)
	Mouse nasal colonization	Significantly reduced colonization	(235)
P5 + PilA	Chinchilla middle ear and nasopharyngeal clearance	Significantly earlier clearance and significant reduction of middle ear biofilm	(236, 237)
Р6	Mouse nasopharyngeal clearance	Significantly enhanced clearance	(238, 239)
Protein D	Chinchilla middle ear clearance Rat middle ear clearance Rat pulmonary clearance	Mixed results	(240, 241)
Protein E	Mouse pulmonary clearance	Significantly enhanced clearance	(116)

Table 1. List of NTHi antigens tested experimentally in vivo for potential as vaccines.
The potential of a series of NTHi antigens for vaccine development has been experimentally investigated (116, 228-241). The bacterial components that have been studied in *in vivo* models are compiled in Table 1. The majority of the tested antigens elicit the desired immune protection, but, with one exception, these findings remain to be verified in translational clinical trials in humans. Moreover, whereas protection against heterologous strains have been addressed in a number of the studies, generally larger strain collections need to be assessed for evaluation of potential prophylaxis in humans.

There is currently one NTHi-component-based vaccine licenced for use in humans, but not against NTHi-mediated disease. *Haemophilus* Protein D has been used as the antigenic protein-carrier for pneumococcal capsular polysaccharides (242). Whereas the efficacy of the vaccine in prevention of invasive pneumococcal disease has been confirmed (243), ambivalent results have been reported regarding its prophylactic capacity with regard to NTHi-mediated infections. One cohort study reported a 33% decrease in NTHi-mediated OM in vaccinated children as compared to an unvaccinated control group (244), but these findings have not been observed in any additional published investigations. On the contrary, other studies have reported that no significant decrease in nasopharyngeal carriage of NTHi has been observed in vaccinated contra unvaccinated children (245-247). This is in stark contrast to the Hib-vaccines, where an abolishment in Hib carriage is observed. As NTHi is an opportunistic pathogen, it is likely that the rate of carriage and incidence of infections correlate. In any regard, the hunt for a more potent NTHi vaccine is currently on going.

Finally, one particularly interesting study merits additional attention. As previously mentioned, bacterial biofilms contain a large amount of extracellular DNA. Goodman and co-workers could show that this DNA is structured in a highly organized three-dimensional grid where bacterial DNA-binding proteins, designated DNABII, appear with regular spatial intervals as the "cornerstones" of every "grid-monomer" (230). Targeting these structural components for immune destruction, researchers could show that immunization with a DNABII-protein known as Integration Host Factor would collapse the entire NTHi biofilm *in vitro* and *in vivo*. This resulted in significantly enhanced clearance of biofilms in a chinchilla middle ear model. Moreover, they could show that biofilm-associated bacteria that were subjected to this treatment displayed a significantly increased susceptibility to antibiotics (141, 230). These findings show that targeting biofilm-components as immunization antigens may yield highly efficient vaccines that could be used in patients already chronically colonized by biofilms.

The Present Investigation

Aims

The aim of this thesis was to at the molecular level study NTHi factors involved in host colonization processes, namely complement-resistance and adherence to the ECM. Newly identified bacterial colonization factors were to be evaluated for potential vaccine development. The specified aims were:

- To in detail characterize the previously reported interaction between *Haemophilus* Protein E and vitronectin.
- To identify novel ECM-binding proteins in NTHi.
- To characterize the interactions between newly identified ECM-binding bacterial proteins and their host ligands.
- To conduct epitope-mapping on newly identified potential antigens.
- To evaluate the effect of immunization with novel bacterial colonization factors in a mouse pulmonary clearance model.

Results and Discussion

Paper I & II: *Haemophilus* Protein E and Protein F delay C9 deposition on the surface of NTHi by interacting with the C-terminal domain of vitronectin.

Nontypeable *H. influenzae* has previously been shown to interact with the complement inhibitory protein vitronectin as a mean to evade the host innate immunity (160). However, the details of this interaction were not elucidated. Moreover, the actual bacterial components that are responsible for vitronectin-binding remain presently unidentified in a series of pathogens (50), suggesting a lack of knowledge of the molecular mechanisms that underlie this virulence trait. By characterizing the protein-protein interaction between Protein E and vitronectin, we aimed to provide new insight into this aspect of bacterial pathogenesis.

In Paper I, we employed site-directed mutagenesis of recombinant Protein E to show that the two positively charged amino acid residues lysine 85 and arginine 86 are responsible for the electrostatic interaction with vitronectin. Moreover, by producing full-length and truncated recombinant human vitronectin in mammalian expression systems, we could identify that Protein E binds to a C-terminal region of vitronectin between amino acid residues 353-363. Importantly, a delayed C9 deposition and MAC-formation was observed in the NTHi wild type strain as compared to the isogenic Δpe deletion mutant. In conclusion, we showed the molecular function of Protein E in the direct vitronectin-mediated inhibition of complement-induced osmolytic killing of NTHi.

It has since been reported that Protein E also interact with Ln and plasminogen, rendering it a highly multifunctional virulence factor (122, 248). Moreover, the crystal structure of Protein E has also been solved (249). The three-dimensional arrangement of the protein revealed that it dimerizes on the bacterial surface, and observations of the concomitant vitronectin- and Ln-binding could be strengthened by the spatially separated binding-sites for the host components (Figure 5).



Figure 5. The crystal stucture of the Protein E dimer. The cartoon and surface-structure representation on the left and right, respectively, show two monomers colored red and blue that dimerize to complete the quaternary stucture of the complex. The vitronectin- and Ln-binding regions are colored in cyan and green, respectively. Image used courtesy of Dr. Birendra Singh.

As important as the Protein E interaction with vitronectin is for NTHi, the isogenic Δpe deletion mutant strain did not display a completely abolished vitronectinbinding, suggesting the presence of additional bacterial components involved in the interplay with the complement inhibitor. The aim of Paper II was to identify these factors.

We ran the NTHi outer membrane vesicle proteome on a 2D-gel and conducted Farwestern blotting with vitronectin to pull down novel vitronectin-binding proteins. This yielded a particularly strong hit to an approximately 30-kDa component. We identified the spot using mass spectrometry and designated the previously uncharacterized protein as *Haemophilus* Protein F (PF), in keeping with tradition of Protein D and E.

Protein F is annotated as a 32 kDa substrate-binding component of an adenosine triphosphate (ATP)-binding cassette (ABC)-transporter and a homolog of the manganese-utilizing YfeA protein of *Yersinia pestis* (250). It has no primary sequence homology to Protein E, ruling out a conserved vitronectin-binding motif in NTHi. However, in analogy with Protein E, we could show that PF sequesters vitronectin on the bacterial surface, inhibits C9 deposition and polymerization, thus down-regulating the MAC-mediated killing and increasing the serum-resistance of NTHi. The simultaneous deletion of Protein E and PF was shown to have a cumulative negative effect on the vitronectin-binding capacity of NTHi. Interestingly, it was observed that PF interacts with the same vitronectin region as Protein E, possibly suggesting a conserved pathogen-binding region on the host molecule as has been reported for Factor H (251). These studies shed light on the multifactorial interactions of NTHi with complement down-regulatory proteins.

Paper III: *Haemophilus* Protein F mediates bacterial adherence to host tissue by interacting with laminin and epithelial cells.

When conducting structural 3D-modeling of PF, we noticed a strong resemblance to streptococcal Ln-binding proteins Lmb and Lbp. This was striking, as the primary sequence homology was very low. We hypothesized that due to the structural properties of PF it could potentially interact with Ln in analogy to Lmb and Lbp.

In Paper III, we could show that PF is a Ln-binding protein that promotes bacterial adherence to immobilized Ln. Isogenic Δhpf knock out mutants (in six different clinical strains) displayed consistently decreased Ln-binding capability as compared to parental wild type strains. Moreover, a cumulative decreased effect in Ln-binding was observed in an isogenic $\Delta pe/hpf$ double mutant.

We could also show that the N-terminus of PF, the same region that interacts with vitronectin, was mediating the binding to Ln. Electronmicrographs of gold-labelled PF indicated that the protein was binding to the LG-domains on the C-terminus of the Ln α -chain. This was highly interesting, as the same binding-region had previously been shown for Protein E (122) as well as for *Mycobacterium leprae* (252), indicating a potentially conserved pathogen-associated motif. This hypothesis is currently under investigation in our lab.

Finally, as many bacterial Ln-binding proteins have been observed to also interact with epithelial cells, we evaluated the PF-mediated bacterial adherence to human epithelial cells. We could show, at both the protein-cell level as well as the cell-cell level, that PF promoted direct bacterial binding to primary human bronchial epithelial cells. We concluded that PF is a novel multifunctional NTHi adhesin that potentiates bacterial colonization of the host by interacting with vitronectin, Ln and epithelial cells.

As ECM reconstruction and NTHi accumulation in sites of disintegrated epithelium have been observed during NTHi infections, the interaction with Ln is indicatively highly important for the pathogenesis of this species (94, 98, 121, 224). This merits further efforts to identify additional Ln-binding proteins, which is currently undergoing in our lab.

One very curious aspect of PF is its presence at the cell surface. As previously mentioned, PF is annotated as a periplasmic substrate-binding component of the ABC-transporter superfamily. Judging by its structure and homology, it is plausibly involved in divalent metal ion transport (Figure 6). It lacks lipidation sites and transmembrane domains but is still consistently detected with highly specific affinity-purified antibodies on the surface of wild type NTHi, and is absent from the surface of the isogenic deletion mutant. Indeed, introducing the *hpf* ORF in a heterologous *Escherichia coli* host results in ascent of PF to the surface of *E. coli*. The mechanism

behind this surface relocation is presently unknown, and we are currently conducting investigations of the phenomenon. It can be mentioned that there are an abundance of reports that show the surface-association of cytoplasmic bacterial enzymes, many of which have been implicated in host-pathogen interactions (253). The term coined to describe these multifunctional factors is "moonlighting proteins". Our data indicated that PF is a moonlighting NTHi protein that exerts diverse functions in the periplasm and on the outer membrane.



Figure 6. Schematic representation of a Gram-negative ABC-transporter. The substrate-binding component (blue), such as Protein F, moves freely in the periplasm. Once it binds its substrate, the protein undergoes a conformational change and gains affinity for the two inner membrane bound permeases (orange and green). It will deliver its cargo to these pores, and the cytoplasmic ATPase component (pink) will help pump the substrate into the cytoplasm by hydrolysing ATP to ADP. Once the substrate-binding protein has delivered its cargo, its conformation opens again, the protein loses affinity for the permeases and is released back into the periplasm (254). Protein F, however, is also consistently detected "moonlighting" on the bacterial cell surface. The mechanism behind the surface-association is currently unclear.

In a recent study, the extracellular biofilm-specific proteins of *S. aureus* were mapped (255). Researchers showed an excessive cell surface-association of cytoplasmic proteins in the early stages of biofilm-formation. They could also show that this mechanism was pH-dependant and suggested that the cytoplasmic proteins adhering to the surface of bacterial cells were crucial for biofilm-formation. The mechanism of release

of cytoplasmic proteins in the environment were not elucidated, but the authors hypothesize that lysis of subpopulations within the bacterial microcommunity is important for the generation of biofilms. As PF has been shown to be a part of the extracellular protein network in NTHi biofilms (138), it is not impossible that a similar mechanism can explain the moonlighting activity of PF. This question remains to be answered.

For being, until recently, uncharacterized, PF is very well-studied protein. As a testament to its importance in NTHi physiology, PF has regularly been detected in several "-omics" studies (Table 2). The protein has been shown to be significantly upregulated during iron/heme deficient growth conditions as well as during bacterial growth in pooled human sputum from COPD patients (173, 176). Moreover, it has been detected as a part the NTHi OMV- and extracellular biofilm-specific proteome (138, 256). Finally, in a seminal study, the PF-operon was shown to be one of 85 non-essential gene elements that are indispensable for bacterial virulence *in vivo* (190). Collectively, these findings show that PF is a highly important host colonization factor and a potential target for antimicrobial therapy.

Investigated Element	Type of Study	Protein F Annotation	Result	Reference
Growth in Fe/Heme restricted media	Transcriptomic	HI0362/yfeA	PF upregulated 1.17-11.74x	(173)
Growth in COPD sputum	Proteomic	NP_438524.1/Yfe A	PF present in 3.932x greater abundance	(176)
NTHi OMV proteome	Proteomic	NTHi0481	PF present (142 proteins in total)	(256)
NTHi biofilm-specific extracellular proteome	Proteomic	HI0362/Q57449	PF present (18 proteins in total)	(138)
Single and IAV co- infection in murine lung	High- throughput sequencing of transposon insertion sites	yfeABCD	PF essential for single and co- infection alike	(190)

Table 2. Detection/role of PF in various NTH	i "-omics" studies.
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Paper IV: Immunization with *Haemophilus* Protein F confers increased mouse pulmonary clearance of NTHi.

As previously stated, a promising antigen to be used as a component in a NTHi vaccine needs to be surface-exposed, antigenic, conserved and ubiquitous in the heterogenic strains. As PF excellently fulfils these criteria, we characterized its antigenicity and investigated the effect of PF-immunization on the pulmonary clearance of NTHi in mice.

In Paper IV, we could show that antibodies raised against the host-interacting Nterminus of the protein were functional in opsonizing our model NTHi strain as well as a heterologous clinical isolate and promote phagocyte-mediated killing of the bacteria in an opsonophagocytic assay.

If PF is ever to be used in a human vaccine, caution has to be observed in order not to induce autoimmunity by forcing an immune response to epitopes that might resemble self-structures. In order to get an indication of the human response to PF without experimental immunization, we assumed that majority of adults have experienced previous bouts of NTHi infections during childhood. Hence, we tested the sera of 60 healthy adult blood donors for presence of anti-PF IgG. In our experimental set up, we could detect circulating anti-PF IgG in the sera of approximately 26% of the individuals, suggesting that PF does not resemble a hostlike epitope and is probably immunogenic in humans. Conducting epitope mapping of PF, we recognized that the human humoral response to PF differed considerably from that obtained in the anti-PF antiserum raised in rabbits using recombinant PF. Human IgG primarily reacted with the host-interacting PF N-terminus, whereas this region was immunoprivileged in rabbits. As NTHi is a human-specific pathogen, it is plausible that the human response confers better protection. To force a humoral response to the N-terminus, mice were immunized four times with KLH-conjugated PF²³⁻⁴⁸ or a negative control peptide. Once the immunization schedule was completed, animals were challenged intranasally with NTHi and the pulmonary clearance of bacteria was examined. Our results showed that immunization with PF²³⁻⁴⁸ caused a significantly increased clearance of NTHi at both tested time points.

These results should be seen in the light of the human-specificity of NTHi. In nature, NTHi is never observed to infect any other species than man. Pfeiffer himself never managed to find an animal model for *H. influenzae* despite inoculating mice, rats, guinea pigs, rabbits, pigs, cats, dogs and monkeys (61). The effect observed in our study can thus conceivably be amplified when translated to human biology. This remains, however, to be experimentally addressed.

In conclusion, immunization with PF will have a three-fold effect. Firstly, blocking a NTHi colonization factor to decrease its adherence to the upper airways. Secondly, opsonizing NTHi with bactericidal antibodies to promote a more rapid clearance in

the case of pathogenesis. And lastly, as PF has been reported to be an integral part of the NTHi biofilm-specific proteome, immunization with PF could potentially help eradicate biofilms as demonstrated for DNABII. We therefore suggest that PF can be a promising candidate as one of the antigens in a multicomponent NTHi vaccine.

Future Perspectives

With new knowledge arise new questions. Without presenting detailed proposals, I would like to highlight some aspects of *H. influenzae* colonization in particular, and bacteriological research in general, that in my opinion need further attention in the future.

Firstly, all available evidence points to a sophisticated bacterial detection of, adaptation to, and selection in, various microenvironments during colonization and pathogenesis. At the same time, the majority of *in vitro* studies are conducted using highly nutrient-rich media developed more than 100 years ago. This is counter-intuitive. Moreover, in the immensely multifactorial *in vivo* studies, it is almost impossible to analyse singular mechanisms at a reductionist molecular level. While conceding that these methods have evidently been effective in uncovering a vast array of bacterial physiological processes, there are plausibly many aspects that have been unnoticed due to the use of irrelevant stress-free culture conditions that may induce down-regulations of important factors. Therefore, I think we will see the field move toward the employment of more sophisticated culture conditions in the future, relevant to the particular biological aspect investigated.

Another interesting issue, in my opinion, is the mechanisms behind moonlighting proteins. The highly common observation of surface-associated cyto- and periplasmic proteins across the entire bacterial kingdom needs to be more systematically investigated. How do proteins relocate? Is the mechanism active or passive? Is it selective or random? And, most importantly, what are the biological roles of these dual locations? As previously mentioned, researchers are gaining new insight into the biological features of moonlighting and this field will hopefully continue to expand.

When NTHi arrives to the site of initial colonization, it has to interact with the other residing prokaryotes. It is plausible that intense and dynamic interspecies exchanges occur. The impact of NTHi colonization on the rest of the normal flora is currently not well studied. It would be of utmost interest to investigate potential changes in the transcriptome and secretome of NTHi and the interacting neighbouring bacteria upon direct cell-cell contact. Does NTHi induce deleterious alterations in the surrounding species in order to prevail in the niche? Or do synergistic interactions occur with regard to the metabolism of, for instance, NAD and heme? Can the normal flora manipulate the gene expression, or the phenotype, of the invading NTHi strain? Simply put, do synergistic, mutualistic or antagonistic interactions occur between NTHi and avirulent members of the human upper airway microbiota? These questions remain largely unknown at the moment but may very well be vital for colonization.

Recently, a large variety of recombinant human Ln isoforms have been commercialized, allowing the study of bacterial interactions with particular Ln variants. In our studies, we could observe that NTHi still interacted with Ln in the absence of Protein E and PF. Using niche-specific Lns, it would be exciting to pull down the additional factors involved in NTHi interaction with the ECM-protein.

The role of PF in NTHi biofilm remains highly interesting to investigate. As the extracellular matrices of bacterial biofilms have consistently been proven to be decidedly coordinated structures rather than gelatinous soups of proteins and DNA, the question that naturally arises is how PF fits into this architecture. And if targeting PF for chemo- or immunotherapy will result in the collapse of already established NTHi biofilm? It would be fascinating to find out.

Lastly, the role of PF in divalent metal ion utilization should be investigated. It is still unclear which function of the multicompetent protein renders it essential for NTHi virulence. By elucidating its role in metal ion utilization, or any other potential role in metabolism, it will be easier to appreciate the importance of its various functions.

In conclusion, many intriguing enigmas of the biology of the humans-specific bacterium *H. influenzae* and its multifunctional colonization factor PF remain to be deciphered. This bodes well for the curious microbiologist.

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Haemophilus influenzae protein E recognizes the C-terminal domain of vitronectin and modulates the membrane attack complex

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Summary

Haemophilus influenzae protein E (PE) is a 16 kDa adhesin that induces a pro-inflammatory immune response in lung epithelial cells. The active epithelial binding region comprising amino acids PE 84-108 also interferes with complement-mediated bacterial killing by capturing vitronectin (Vn) that prevents complement deposition and formation of the membrane attack complex (MAC). Here, the interaction between PE and Vn was characterized using sitedirected mutagenesis. Protein E variants were produced both in soluble forms and in surface-expressed molecules on Escherichia coli. Mutations within PE⁸⁴⁻¹⁰⁸ in the full-length molecule revealed that K85 and R86 residues were important for the Vn binding. Bactericidal activity against H. influenzae was higher in human serum pre-treated with full-length PE as compared with serum incubated with PEK85E, R86D, suggesting that PE guenched Vn. A series of truncated Vn molecules revealed that the C-terminal domain comprising Vn³⁵³⁻³⁶³ harboured the major binding region for PE. Interestingly, MAC deposition was significantly higher on mutants devoid of PE due to a decreased Vn-binding capacity when compared with

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wild-type *H. influenzae*. Our results define a finetuned interaction between *H. influenzae* and the innate immune system, and identify the mode of control of the MAC that is important for pathogen complement evasion.

Introduction

Haemophilus influenzae is a Gram-negative, human respiratory tract pathogen and is categorized as typeable (encapsulated) and non-typeable (unencapsulated) strains. Typeable strains are further classified into serotypes a to f (Meats et al., 2003). Among the typeable strains, H. influenzae type b (Hib) is considered to be most virulent causing bacteraemia, pneumonia and acute bacterial meningitis or occasionally, osteomyelitis, epiglottitis and joint infections (Watt et al., 2009). Non-typeable H. influenzae (NTHi) is most commonly associated with otitis media, sinusitis and chronic obstructive pulmonary disease (COPD) (Murphy, 2003; Sethi et al., 2004, Look et al., 2006). A crucial factor in the H. influenzae pathogenesis involves the initial adherence to the respiratory mucosa. When H. influenzae overcomes the mucociliary escalator and attaches to the epithelium, it may colonize and also damage epithelial cells. This consequently leads to inflammation and deep tissue penetration. H. influenzae usually adheres to and interacts with epithelial cells by many surface exposed proteins including the adhesins high-molecular-weight protein (HMW-1 and HMW-2) (Dawid et al., 1999; Giufre et al., 2008; Gross et al., 2008), Haemophilus surface fibrils (Hsf) (Hallström et al., 2006), H. influenzae adhesin (Hia) (Winter and Barenkamp, 2009), Haemophilus adhesion penetration protein (Hap) (Fink and St Geme, 2003), cryptic Haemophilus adhesin protein (Cha) (Sheets et al., 2008), haemagglutinating (HA) pili (Gilsdorf et al., 1992), Pil A (Jurcisek et al., 2007), protein D (Forsgren et al., 2008) and protein E (PE) (Ronander et al., 2008). Several autotransporters are also H. influenzae virulence determinants and function as adhesins (Poolman et al., 2000). In recent years, characterization of NTHi adhesins has gained much attention since a future vaccine against NTHi is required (Murphy, 2009).
Gram-negative pathogens are usually recognized by the innate immune system comprising complement components, and are finally killed by the membrane attack complex (MAC). However, many respiratory tract pathogens including H. influenzae can overcome the activation of the complement system as well as inhibit the deposition of MAC on the outer membrane (Blom et al., 2009; Hallström and Riesbeck, 2010). H. influenzae inhibits complement activation by interacting with complement C4bbinding protein (C4 BP) that is an inhibitor of the classical and lectin complement pathways (Hallström et al., 2007). Similarly, Haemophilus also binds factor H, which inhibits the alternative complement pathway resulting in marked serum resistance (Hallström et al., 2008). In addition. H. influenzae interacts with several extracellular matrix proteins (ECM) such as vitronectin (Vn), laminin, fibronectin and different types of collagens (Bresser et al., 2000; Fink et al., 2002), which help in adhesion and colonization in the respiratory tract. Among these ECM, Vn is a multifunctional serum glycoprotein that binds to complement complex C5b-6 and complement 9 (C9). Recruitment of Vn to the bacterial surface thus leads to inhibition of the MAC (Singh et al., 2010a). The precise mechanisms and protein-protein interactions involved between bacteria and Vn are largely unknown.

We have isolated and characterized the H. influenzae adhesin PE (Ronander et al., 2008; 2009). A pe-deficient isogenic *H. influenzae* mutant showed a significantly reduced adherence to epithelial cell as compared with wild-type bacteria, and recombinant soluble PE as well as PE-expressing Escherichia coli efficiently bound to epithelial cells. Moreover, PE induces a pro-inflammatory response leading to IL-8 secretion in addition to an increased upregulation of ICAM-1 (CD54) in both cell lines and primary epithelial cells originating from patients with COPD (Ronander et al., 2009). The central core domain of PE (amino acids 84-108) is the active epithelial cell-binding region. Importantly, mice immunized with PE^{84–108} demonstrated a significantly increased pulmonary clearance of NTHi as compared with controls immunized with a non-related peptide (Ronander et al., 2009). Protein E is a highly conserved ubiquitous adhesin among all Haemophilus spp. and homologues of PE also exists in other members of the Pasteurellaceae family (Singh et al., 2010b). Intriguingly, the region PE 84-106 is completely conserved at the amino acid level suggesting an important role in bacterial pathogenesis. These reports indicate that PE may be a valuable vaccine candidate. We also recently reported that PE binds to Vn and contributes to serum resistance (Hallström et al., 2009). The isogenic pe-mutant was unable to bind Vn and was also significantly more serum sensitive. When the PE-Vn interaction was analysed. PE⁸⁴⁻¹⁰⁸ was identified as the active Vnbinding region. Most interestingly, it was shown that the PE–Vn complex was functionally active and inhibited the MAC assembly *in vitro*.

In the present study, we examined in detail the PE⁸⁴⁻¹⁰⁸ active binding region at the amino acid level in order to understand the mode of Vn binding. In addition, the Vn regions bearing the PE-binding properties were characterized. Several variants of recombinant PE⁸⁴⁻¹⁰⁸ were generated by using site-directed mutagenesis and the K85 and R86 residues of PE were found to be involved in the interaction with Vn. In agreement with the observation that heparin blocks the binding of PE to Vn (Hallström et al., 2009), we found that the C-terminal heparin-binding domain (HBD-3) of Vn corresponding to residues 353-363 is mainly involved in this interaction. It is known that PE-dependent binding of Vn to the NTHi surface plays a role in acquiring serum resistance (Hallström et al., 2009; Singh et al., 2010a). Therefore, we directly measured the MAC deposition at the bacterial surface, and a several-fold higher deposition of MAC on the NTHi Ape mutant was observed in comparison with the NTHi wild type. In conclusion, this study shows that the PE-Vn interaction at the molecular level plays a highly important role in H. influenzae serum resistance.

Results

Lysine 85 and arginine 86 of PE are involved in vitronectin binding

To identify amino acid residues responsible for binding to Vn, the Vn binding region of PE (PE⁸⁴⁻¹⁰⁸) was targeted. According to the internet-based tool GOR4 (http://npsapbil.ibcp.fr) PE had 33.8% a-helices, 25.6% extended strand and 40.6% random coils in its secondary structure (Fig. 1A). This estimation was further confirmed using circular dichroism (CD) spectroscopy (Fig. 1E) and smallangle X-ray scattering (SAX). The Vn-binding region PE⁸⁴⁻¹⁰⁸, which contains a helical region and loops was expected to be surface-exposed, at least partially, in order to interact with its ligand Vn (Fig. 1A black background). Thus, based on our previous findings (Hallström et al., 2009; Ronander et al., 2009) and this secondary structure prediction, PE⁸⁴⁻¹⁰⁸ was targeted by site-directed mutagenesis to identify the amino acid residues involved in the interaction with Vn. The charges of amino acid side-chains were altered to opposite charges, or the amino acids were changed to alanine (Fig. 1B).

The mutated PE variants were recombinantly expressed in *E. coli* and purified using Ni-NTA affinity chromatography. The binding of Vn to PE^{WT} (amino acids 22–160) or variants was analysed using ELISA and surface plasmon resonance. The change of the positively charged side-chain of Lys-85 to a negatively charged Glu reduced the binding by 50% (Fig. 1C). Another positively charged



Fig. 1. Amino acids K85 and R86 of PE are involved in binding to Vn.

A. Prediction of the PE secondary structure by the internet-based tool GOR4 (http://npsa-pbil.ibcp.fr). Arrows represent extended strands while tubes represent α -helices. The Vn binding region 84–108 is indicated with a black background.

B. The plan of site-directed mutagenesis of the recombinant His-tagged PE molecules (amino acids 84–108). The side-chains of charged amino acids were either switched to opposite charged side-chains or mutated to alanine.

C. Binding of Vn (Sigma) to PE and variants as revealed by surface plasmon resonance (Biacore). Vn was immobilized on a chip and PE variants (200 μg ml⁻¹) were analysed for binding. The binding of Vn to PE^{WT} in addition to PE^{K85E}, PE^{R86D} and PE^{K85E, R86D} are shown as sensorrarms.

D. ELIŠA showing binding of PE variants to Vn. Approximately 0.01 μ M of each protein was coated on ELISA plates followed by addition of increasing amounts of Vn (Sigma). Bound Vn was detected by using sheep anti-Vn pAb. Mean values of triplicates from three independent experiments were plotted, error bars representing standard deviations and curves are hyperbolic fits. The binding at 5–50 nM Vn concentration was statistical significant when PE^{WT} was compared with mutants ($P \le 0.001$) (ANOVA).

E. CD spectra of PE variants. Five spectra for each protein (5 µM) were recorded and mean values plotted.





Competitor ligands [1µM]

NTHi 3655 A. Unlabelled PEWT blocks binding of [125]-labelled Vn to NTHi 3655 in a direct binding assay as described in detail in Experimental procedures. B. Binding of [125]]-Vn to NTHi 3655 is blocked with PE^{WT}, PE^{K85E}, PE^{R86D}, whereas the double mutant PE^{K85E}, ^{R86D} does not block.

Fig. 2. PE blocks the Vn interaction with

Mean values of triplicates from three independent experiments were plotted and standard deviations were shown as error bars. Differences between the control (without any competitor) and $\mathsf{PE}^{\mathsf{K85E},\ \mathsf{R86D}}$ were statistically significant with PEWT, PEK85E PE^{R86D} ($P \le 0.001$). When PE^{WT} , PE^{K85E} and PERBED were compared these were not significant (ANOVA).

side-chain containing amino acid residue Arg-86 was changed to Asp, and also showed a 48% reduction in binding to Vn. Alteration of the charged side-chains of K85 and R86 in the double mutant PE^{K85E, R86D} (Fig. 1B: mut2) showed defective Vn binding (Fig. 1C). In addition, similar experiments were performed with all PE variants using ELISA, where PE variants were coated on microtitre plates and Vn binding was detected using anti-Vn pAb. Our results revealed that PE^{R85E} and PE^{R86D} decreased Vn binding by 30–50% and that the PEK85E, R86D double mutant of PE lost 70% Vn binding (Fig. 1D). PEWT, PEK85E, PER86D and PE^{K85E, R86D} were analysed for folding pattern by using CD, and these analyses suggested that PE variants were folded (Fig. 1E). All the PE variants shown in Fig. 1B were tested for Vn binding by Biacore and ELISA. Other multiple mutations (Fig. 1B: mut3, mut4, mut5) in the PE molecule showed similar binding pattern to the Vn molecule as shown by PE K85, R86 (data not shown).

Lysine 85 and arginine 86 residues of PE on NTHi surface recognize Vn

The specificity of the binding of Vn to non-typeable H. influenzae (NTHi) 3655 was also verified by a competition assay, where [125I]-Vn binding was blocked with unlabelled PE variants. Vn binding to NTHi was inhibited by unlabelled PE^{WT} in a dose-dependent manner (Fig. 2A). Single mutants of the PE molecule, i.e. PEK85E and PER86D competed with Vn (Fig. 2B), whereas the double mutant PE^{K85E, R86D} did not inhibit Vn binding to NTHi 3655. These results suggested that K85 and R86 were crucial for Vn binding and were involved in the PE-Vn interaction at the bacterial surface. Moreover, since recombinant proteins with alterations of the K85 and R86 side-chains to acidic charged counterparts lost PE-Vn binding, indicating that the interaction may be electrostatic. However, these data solely cannot prove the nature of this interaction.

To analyse the effect of mutated K85 and R86 when expressed at the surface (similarly as expression in NTHi 3655), PE variants were expressed in E. coli. Expression of PE was monitored by flow cytometry (Fig. 3D) followed by binding of recombinant Vn⁸⁰⁻³⁹⁶ that was analysed by transmission electron microscopy (TEM) (Fig. 3A). E. coli (PEWT) displayed a similar binding pattern as compared with the single mutants expressing $\mathsf{PE}^{\text{\tiny K85E}}$ and $\mathsf{PE}^{\text{\tiny R86D}}$ (857–945 gold particles μ m⁻²), whereas the double mutant (PE^{K85E, R86D}) had a five- to sixfold reduced binding (248 particles µm⁻²) to recombinant Vn (Fig. 3A). The compilation of several fields of vision of Vn80-396 binding to E. coli variants is shown in Fig. 3B.

In addition to experiments with recombinant Vn⁸⁰⁻³⁹⁶, the binding of native serum Vn to E. coli expressing PE wild type and variants at the surface was analysed by semiquantitative Western blotting. Bacteria were treated with normal human serum (NHS) diluted 1:25 to 1:500 in PBS containing 2.5% BSA followed by analysis of Vn bound to the bacterial surface using anti-Vn pAb (Fig. 3C). These results demonstrated that E. coli expressing PEWT at the surface acquired Vn binding capacity, whereas E. coli ($PE^{K85E, R86D}$) lost Vn binding and showed a similar Vn binding as the control E. coli devoid of PE (Fig. 3B). Thus, these two data sets indicated that K85 and R86 are both necessary for Vn binding when PE was present at the E. coli surface. This experimental model with E. coli also mimicked the functional role of PE as compared with expression in NTHi 3655.

The C-terminal of Vn (amino acids 353–363) is a major PE binding region

Vitronectin is a 65-75 kDa glycoprotein that contains a 43-amino-acid-long somatomedin-B (SMB) domain, which binds plasminogen activator inhibitor-1 (PAI-1) (Arroyo De Prada et al., 2002; Schroeck et al., 2002). In addition, Vn has four putative haemopexin-like regions and three heparin-binding domains (HBDs) (Singh et al., 2010a), and in the case of PE, heparin is known to inhibit binding to Vn (Hallström et al., 2009). To exclude that the 84 B. Singh et al.



Fig. 3. PE at the bacterial surface binds to Vn by the residues K85 and R86.

A. Binding of gold-labelled Vn⁸⁰⁻³⁹⁶ to *E. coli* expressing PE variants at the surface as shown with TEM.

B. The results analysed from (A) were plotted as a bar diagram. Number of gold particles was determined in 60 independent bacterial profiles and number of particles μ m⁻² was calculated. Mean of triplicates from three experiments are shown and error bars represent SD. The difference between *E. coli* expressing PE^{WT} and PE^{KSE, R800} was statistically significant ($P \le 0.001$). The bar represents 100 nm. C. Serum Vn binds to *E. coli* expressing PE variants. Bacteria (10⁸) were treated with different serum dilutions in PBS containing 2.5% BSA and washed three times in PBS. Total cellular proteins were separated on SDS-PAGE and blotted to a PVDF membrane followed by detection

and washed three times in PBS. Total cellular proteins were separated on SDS-PAGE and blotted to a PVDF membrane followed by detectior with anti-human Vn pAbs. Two bands with Vn (molecular weights 75 kDa and 65 kDa) were detected in Western blot. This experiment was repeated twice and one representative experiment is shown here.

D. Flow cytometry profile of *E. coli* expressing PE variants that were used in these experiments. Filled curve represents control *E. coli* with empty pET16b vector.

SMB domain was involved in the PE–Vn binding, we pre-incubated PE with PAI-1 followed by addition of Vn. Binding was analysed by ELISA and dot blot, but no inhibitory effect of PAI-1 was observed (data not shown). We have recently shown that *Moraxella catarrhalis* Ubiquitous surface protein (Usp) A2 binds to the C-terminus (HBD-3) of the Vn molecule (Singh *et al.*, 2010c). In addition to recombinant Vn molecules used in that particular

study, we created a series of new truncated Vn fragments (Fig. 4A). The protein fragments were purified and their purity and nature was studied by using SDS-PAGE, native PAGE and CD (Fig. 4B–E). All Vn fragments showed monomeric and multimeric forms and folding patterns in non-reducing PAGE and CD respectively. However, the smallest fragment Vn⁸⁰⁻²²⁹ lacked a secondary structure in CD (Fig. 4E).

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Fig. 4. Expression of recombinant Vn fragments and analyses for folding and multimerization.

A. Schematic representation of the different truncated Vn molecules that were expressed in a mammalian expression system. B. Quality control of purified recombinant Vn molecules expressed in HEK293T cells. All protein fragments were purified by Ni-NTA

chromatography and subjected to 15% SDS-PAGE (approximately 2 µg per lane) that was stained with Coomassie blue R250.

C. Non-reducing SDS-PAGE showing the Vn fragments monomers and multimers. Each fragment (5 µg) was loaded in a 4–12% gradient SDS polyacrylamide gel and stained with Coomassie blue R250.

D. A similar gel (as shown in C) was analysed by Western blotting by using anti-Vn antibodies.

E. CD spectra of Vn fragments at a concentration of 5 µM. Five spectra for each Vn fragment were recorded and mean values are shown here.

The purified recombinant Vn proteins (Fig. 4B) were used for binding to whole bacteria and different soluble PE variants. NTHi 3655 wild type (10^{8} bacterial cells) were incubated with Vn fragments ($0.5 \mu g$) and Vn bound to the bacterial surface was analysed by whole-cell protein Western blotting (Fig. 5A). Only Vn⁸⁰⁻³⁹⁶, Vn⁸⁰⁻³⁷⁹, Vn⁸⁰⁻³⁷³ and Vn⁸⁰⁻³⁶³ bound to the NTHi surface, whereas Vn⁸⁰⁻³⁵³ and further shorter truncated fragments of Vn did not interact with NTHi. In the next series of experiments we analysed binding of Vn fragments to purified PE in ELISA. Microtitre plates were coated with PE^{WT} and Vn fragments (O-100 nM) were allowed to bind. Resulting bound Vn fragments revealed that Vn⁸⁰⁻³⁹⁶ and Vn⁸⁰⁻³⁷⁹ displayed similar binding saturation kinetics, while Vn⁸⁰⁻³⁷³ and Vn⁸⁰⁻³⁶³

showed a 20–30% reduced PE binding in comparison with Vn^{80–396} and Vn^{80–379} (Fig. 5B). Further truncation of Vn reduced binding to the PE molecule, indicated that Vn 353–363 is a major PE binding region while a minor interaction may also be involved in Vn 363-373. A comparative binding analysis of PE variants to all Vn fragments was also performed (Fig. 5C). Vn^{80–396}, Vn^{80–379} bound PE^{WT}, PE^{K85E} and PE^{R86D} stronger than Vn^{80–373} and Vn^{80–363}. In contrast, the double mutant PE^{K85E}, R^{86D} significantly lost Vn binding with all Vn fragments (Fig. 5C). Our data thus confirmed that further truncation after amino acids Vn^{80–363} resulted in lost binding to the PE molecule.

Binding of the Vn⁸⁰⁻³⁹⁶ fragments to the NTHi 3655 surface was also analysed for binding specificity and competition with other smaller fragments. NTHi 3655 was



Fig. 5. Vn 353-363 binds to NTHi PE.

A. Binding of Vn fragments to PE-expressing NTHi 3655 wild type as revealed by Western blot. Bacteria (10⁸) were incubated with 0.5 μg of each Vn fragment and washed twice with PBS. Thereafter, total cellular proteins were separated on an SDS-PAGE and blotted onto a PVDF membrane. This experiment was repeated three times, and one representative blot is shown here.

B. Binding of Vn fragments to wild-type PE. Recombinant PE (0.1 μ M) was coated in microtitre plates and increasing amounts of all Vn fragments were added, followed by detection of bound Vn using anti-human Vn pAbs. Data shown are the means of triplicates from three independent experiments and curves shown are hyperbolic fits. The binding of Vn to PE^{WT} at 1.5–100 nM was statistically significant ($P \le 0.001$) when Vn^{80–336} or Vn^{80–379} and other fragments were compared. Differences between Vn^{80–336} and Vn^{80–379}, and between Vn^{80–373} and Vn^{80–379} and between Vn^{80–379} and vn^{80–379}.

C. Comparison of PE variants bound to different Vn fragments. Equimolar amounts (0.1 μ M) of PE variants were coated on microtitre plates and 20 nM of each Vn fragment was allowed to bind. Data represent means of triplicates from three independent experiments and error bars indicate standard deviations. In the Vn groups Vn⁸⁰⁻³⁶⁹, Vn⁸⁰⁻³⁷⁹, Vn⁸⁰⁻³⁷³ and Vn⁸⁰⁻³⁸³ the binding differences between PE^{WT} and PE mutants were statistically significant ($P \le 0.01$) (ANOVA).



Fig. 6. Vn 353-363 amino acids interact to PE at the NTHi surface.

A. Binding of [¹²⁵I]–Vn⁸⁰⁻³⁹⁶ (approximately 300 000 kcpm) at the NTHi 3655 surface and inhibition by using cold Vn fragments. [¹²⁵I]–Vn⁸⁰⁻³⁹⁶ was added to 10⁹ bacterial cells, after extensive washing bound fraction was counted in a gamma counter. In blocking experiments, bacteria were treated with Vn⁸⁰⁻³⁹⁶, Vn⁸⁰⁻³⁹⁸ and Vn⁸⁰⁻³⁹⁵ prior to addition of labelled Vn. [¹²⁵I]–Vn⁸⁰⁻³⁹⁶ binding without any competitor was compared with treated samples.

B. PE^{WT} protein was coated on MaxiSorp ELISA plates (strip plates), [¹²⁵]–Vn⁸⁰⁻³⁹⁶ (approximately 300 000 kcpm) was added, the unbound fractions were washed and bound fractions were counted. In competitive experiments Vn⁸⁰⁻³⁹⁶, Vn⁸⁰⁻³⁹³ and Vn⁸⁰⁻³⁶³ were added to PE-coated wells prior to addition of the [¹²⁵]–Vn⁸⁰⁻³⁹⁶.

In both experiments means of triplicates from three independent experiments (A and B) were plotted and standard deviations are shown as error bars. Differences between control (without any competitor) or Vn^{80-353} were statistically significant with Vn^{80-396} and Vn^{80-363} treatment ($P \le 0.01$). Differences between Vn^{80-396} and Vn^{80-363} were not significant (ANOVA).

allowed to bind [¹²⁵]–Vn^{80–396} and the other smaller unlabelled fragments (Vn^{80–363} and Vn^{80–353}) were used for competition. Results showed that Vn^{80–396} and Vn^{80–363} blocked 40–50% binding of the [¹²⁵]]–Vn^{80–396} molecule, whereas Vn^{80–353} did not inhibit binding (Fig. 6A). Similar results were also observed when PE^{WT} was coated on a microtitre plate and [¹²⁵]]–Vn^{80–396} binding to immobilized PE^{WT} was analysed. Here also Vn^{80–396} and Vn^{80–363} blocked 60–70% of the binding to the [¹²⁵]]–Vn^{80–396} fragment, whereas Vn^{80–353} did not significantly block (Fig. 6B). These both experiments directly indicated that the major PE binding region of Vn is located between amino acids 353 and 363.

In our next set of experiments, Vn fragments were labelled with gold particles and their interaction with whole NTHi 3655 and NTHi 3655 Δpe in addition to *E. coli* expressing PE at the surface was analysed by TEM. As exemplified in Fig. 7A, only the gold-labelled fragments Vn⁸⁰⁻³⁹⁶, Vn⁸⁰⁻³⁷⁹, Vn⁸⁰⁻³⁷³ and Vn⁸⁰⁻³⁶³ interacted with NTHi 3655, whereas the other fragments were unable to bind (all data not shown). As can be seen in the bar diagram (Fig. 7C), 636 gold-Vn⁸⁰⁻³⁹⁶ particles, 798 gold-Vn⁸⁰⁻³⁶³ and Vn⁸⁰⁻³⁶³ and Vn⁸⁰⁻³⁶³ and Vn⁸⁰⁻³⁶³ and this was in comparison with Vn⁸⁰⁻³⁵³ and Vn⁸⁰⁻³⁹⁶ that only bound 125 and 61 particles μ m⁻² respectively. NTHi 3655 Δpe mutants showed a sixfold reduced Vn⁸⁰⁻³⁹⁶ binding with 106 particles µm⁻². The observed minor Vn binding to the PE-deficient NTHi 3655 Δpe could be due to

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other weak Vn-binding surface proteins (Fig. 7A). Moreover, PE-expressing *E. coli* had a similar Vn-binding pattern as compared with NTHi 3655 (Fig. 7B). *E. coli* (PE^{WT}) bound 881 Vn^{80–396} gold particles μ m⁻², whereas the *E. coli* control without PE bound considerably less Vn^{80–396} (45 particles μ m⁻²). Regarding the other Vn fragments, a similar pattern as with NTHi 3655 was observed with *E. coli* (PE^{WT}) (Fig. 7B and D). The TEM data were thus comparable and in agreement with results obtained *in vitro* (Fig. 5B), or *in vivo* at the bacterial surface (Figs 5A and 6A). These results clearly indicated that the C-terminal domain in Vn corresponding to residues Vn 353–363 is a major region for the interaction with PE.

Non-typeable H. influenzae is more sensitive to human serum that is pre-treated with PE

Vitronectin binds C9 and thus inhibits its insertion into the bacterial membrane during MAC assembly (Singh *et al.*, 2010a). When Vn is bound to the bacterial surface, the MAC is inhibited and bacteria have a prolonged survival when exposed to NHS. The PE-expressing wild-type NTHi 3655 was resistant to 10% NHS, and compared with the PE-deficient mutant NTHi $3655\Delta pe$ that was significantly less serum resistant (Fig. 8A). We recently reported that Vn-depleted serum markedly increased *H. influenzae* killing, and that addition of Vn to depleted NHS restored the bacterial serum resistance (Hallström *et al.*, 2009).



Fig. 7. Transmission electron microscopy of NTHi 3655 and *E. coli* expressing PE^{WT}. A. Binding of gold-labelled recombinant Vn⁸⁰⁻³⁹⁶, Vn⁸⁰⁻³⁸³, Vn⁸⁰⁻³⁸³ and Vn⁸⁰⁻³²⁰ to the surface of NTHi 3655.

B. Binding of similar gold-labelled Vn fragments to *E. coli* expressing PE^{wr} at the surface.

C and D. Bar diagrams based upon the experiments delineated in (Å) and (B). Vn molecules (gold particles µm-2) that were bound to NTHi 3655 and E. coli expressing PEWT were determined in 60 independent bacterial profiles and number of particles µm-2 was calculated. Mean of triplicates from three experiments are shown and error bars represent SD. Differences were statistically significant when PE binding to Vn⁸⁰⁻³⁹⁶ and Vn⁸⁰⁻³⁶³ were compared with the other truncated Vn fragments ($P \le 0.001$). Bars in (A) and (B) indicate 100 nm.

Hypothetically, if NHS is incubated with soluble PE, an increased C9 deposition at the bacterial surface may occur due to guenching of Vn, and will be followed by an accelerated MAC-mediated killing. Hence, the Vn-binding capacity of recombinant PEWT or PE variants was analysed by pre-incubation of these recombinant proteins with NHS followed by exposure to NTHi 3655. When purified PE^{WT} was added to NHS, a dose-dependent decrease in bacterial survival was seen in comparison with the recombinant control protein MID⁹⁶²⁻¹²⁰⁰ (Fig. 8B). This effect of



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Fig. 8. H. influenzae is more sensitive to serum that has been pre-treated with PE. A. Serum resistance of NTHi 3655 and NTHi 3655∆pe when incubated with 10% NHS. B. Pre-incubation of NHS with increasing concentrations (1-5 µM) of recombinant PEWT quenches Vn and hence NTHi appears more serum sensitive. MID⁹⁶²⁻¹²⁰⁰ is a protein fragment from Moraxella catarrhalis IgD-binding protein (MID) that was used as a negative control (Nordström et al., 2006). C. NTHi 3655 was incubated with NHS, NHS treated with PE (2 µM) and PE-treated NHS was added to Vn (2 µM)-treated bacterial cells. Differences were statistically significant (5-10 min) when NHS only and NHS+PE or NHS+PE+Vn were compared ($P \le 0.05$), as well as differences between NHS+PE and NHS+PE+Vn ($P \leq 0.001$) (ANOVA). D. PEWT, PEK85E, PER86D showed decrease in serum resistance, whereas PEK85E, R86D did not show any decrease of resistance when added to NHS, PE^{WT}, PE^{K85E} and PE^{R86D} were significantly different ($P \le 0.001$) when compared with control NHS or PEK85E, (ANOVA). No significant difference was observed between control NHS and

In (A)–(D) means of triplicates from three independent experiments are shown and standard deviations are indicated as error bars.

Vn neutralization could be replenished if NTHi 3655 was treated with Vn prior to addition of PE treated serum (Fig. 8C). We also tested whether it was possible to quench Vn by incubating NHS with the different mutated PE variants followed by incubation with whole bacteria. Interestingly, pre-incubation of NHS with PE^{K85E} and PE^{R86D} resulted in significantly higher bacterial killing, whereas experiments with PE^{K85E}. R86D did not reveal any difference in bactericidal serum activity (Fig. 8D). Pre-treatment of NHS with recombinant PE^{WT} and mutated variants thus indicated that PE was able to bind Vn derived from serum, and that in particular residues K85

Vitronectin bound to PE prevents MAC deposition at the surface of H. influenzae

and R86 were responsible for the binding. These results

were in parallel with findings obtained with the direct

binding assays (Figs 2, 3 and 5).

It is well known that Vn inhibits the MAC complex and this property of Vn is successfully utilized by many pathogens to evade the host innate immune response (Singh *et al.*, 2010a). Recently we found that NTHi $3655\Delta pe$ signifi-

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cantly lost Vn binding (as also shown in Fig. 7A and C). and that this PE-dependent Vn interaction is involved in serum resistance (Hallström et al., 2009). To analyse the effect of the PE-Vn interaction on MAC formation, we performed a serum resistance assay and observed the concentrations of C9 polymerized, and measured C9 monomers consumed during MAC deposition. Western blotting using a specific anti-C9 pAb that recognizes both monomeric and polymeric C9, and subsequent densitometric analyses were performed. Under these experimental conditions, MAC deposition on bacteria occurred in a time-dependent manner (Fig. 9A and B). C9 is a 71 kDa monomer that may exist in a monomeric or oligomeric state, associated as a C5b-9 complex in NHS under normal physiological conditions (Podack and Tschopp, 1982). After complete polymerization of C9 (ring closure), a high-molecular-weight band of approximately 400-500 kDa (a C9 oligomer) appears (Podack and Tschopp, 1982; Deng et al., 2007). The disappearance of the C9 monomer in the presence of NTHi 3655∆pe (Fig. 9B) was more evident in comparison with the NTHi 3655 wild type, and an increased C9 oligomerization directly revealed that MAC formation was more efficient in reactions



containing PE-deficient NTHi mutants (Fig. 9A and B). Densitometric measurements of C9 when incubated with the PE-expressing NTHi 3655 revealed that within 10-15 min of incubation 75-90% of C9 was a monomer (Fig. 9C). In contrast, only 40-45% of C9 was remaining as a monomer when exposed to NTHi $3655 \Delta pe$.

We also performed TEM of these serum-treated samples. Samples at 5 and 10 min (as described in Fig. 9A and B) were chosen and analysed by goldlabelled anti-C9 pAbs. Data obtained by TEM also suggested a time-dependent deposition of C9. The number of gold particles reflecting C9 was counted from 60 independent bacterial profiles. At 5 min, 82 gold particles μ m⁻² were found on the NTHi 3655 wild type, whereas 974 gold particles um⁻² were detected on the NTHi 3655 (pe mutant. Similarly, 114 particles μm^{-2} and 1224 particles

2

4

6

Time (min)

8

10

0

0

Fig. 9. MAC deposition at the H. influenzae surface is regulated by PE-dependent Vn binding.

A and B. NTHi 3655 and NTHi 3655∆pe (10⁸ cells) were resuspended in GVB⁺⁺ buffer containing 10% NHS and incubated at 37°C. MAC deposition is seen as increase in C9 oligomer and disappearance of the C9 monomer. The complement reaction was terminated at different time points (5–60 min) by snap freezing of samples. Samples were subjected to 4–12% gradient SDS-PAGE, blotted onto PVDF membrane followed by detection using an anti-human C9 pAb. The experiment was repeated twice. One of the representative blots from triplicates of a single experiment is shown here.

C. The C9 monomer density was measured from triplicate blots of a single experiment. Error bars indicate standard deviations. Statistical significant differences were observed between 10 and 40 min (ANOVA). The experiment was repeated twice.

D. TEM analysis that demonstrates C9 deposition on the surface of NTHi 3655 and NTHi 3655 *pe*. Samples obtained at 5 and 10 min from (A) and (B) were probed with gold labelled anti-human C9 pAb.

E. Estimation of gold-labelled Vn bound to the wild type and PE-deficient NTHi mutant. The curve shows the number of particles µm⁻² evaluated from 60 independent bacterial profiles. Mean of triplicates from three experiments are shown and error bars represent SD.

Bar represents 100 nm. * $P \le 0.05$; ** $P \le 0.01$; *** $P \le 0.001$.

 μm^{-2} were seen after 10 min on the NTHi wild type and the mutant surface, respectively (Fig. 9D and E). These results directly explained why NTHi 3655 Δpe mutants were more serum-sensitive as compared with the wild type, and also that the PE–Vn interaction was highly important for NTHi serum resistance.

In our next set of experiments we analysed the capacity of the PE^{WT} and PE^{K85E, R86D} variant for Vn binding and subsequent effect in MAC formation at the NTHi 3655



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surface. PE^{WT} and PE^{K85E, R86D} were immobilized on CNBr agarose beads to simulate PE as a surface bound protein and checked on a Western blot (Fig. 10A). Control beads were treated similarly without any added protein. The pull-down capacity of these beads was measured by semi-quantitative Western blotting at different serum dilutions. PE^{WT} bound Vn more intensely in comparison with PE^{K85E, R86D} (Fig. 10B). To bind Vn, beads coated with PE (equivalent to 2 μM protein) and control beads were

> **Fig. 10.** PE-treated serum increases MAC deposition at the bacterial surface. A. CNBr agarose beads were immobilized with PE^{WT} and PE^{K85E, R86D} proteins (see *Experimental procedures*). One set of beads were treated similarly without protein to generate control beads. Agarose beads bound PE (5 μg) along with control were resolved in a 12% SDS-PAGE and stained with Coomassie blue R250.

B. Pull-down assay showing binding of Vn by PE-coated beads. PE-coated beads (1 µg) and control beads were added to 100 ul of different serum dilutions in PBS containing 2.5% BSA. Beads were washed extensively with PBS and proteins were separated in SDS-PAGE followed by Western blotting and detection of Vn by anti-Vn antibodies. This experiment was repeated twice; one representative blot is shown here. C-E. Normal human serum was treated with control, PEWT- and PEK85E, R86D-coated beads (2 µM) prior to adding to bacteria. C9 deposition experiments were performed as described in Fig. 9. This experiment was repeated twice. From a single experiment samples, each blot was performed in triplicate; one representative blot out of triplicate is shown here.

F. The C9 monomer density was measured from triplicate blot of a single experiment. The data for 5-15 min are shown. The differences (5-15 min) between control beads and PE^{WT} were statistically significant ($P \le 0.01$). A comparison between PE^{WT} and PE^{K8E, R80D} beads revealed $P \le 0.05$ (ANOVA). Any statistical significance was not observed between control beads and PE^{K8E, R80D}-coated beads.

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added to serum and incubated for 1 h. This serum was subsequently tested for C9 deposition at the NTHi 3655 surface. The results shown in Fig. 10C–E represent C9 deposition on control, PE^{WT} and PE^{K85E, R86D} beads when incubated with serum. Three blots from a typical experiment were scanned for C9 band intensity and are shown in the bar chart (Fig. 10F). The serum incubated with control beads showed a similar C9 disappearance as treatment with beads coated with PE^{K85E, R86D}, whereas serum incubated with PE^{WT}-containing beads had a comparatively faster disappearance of C9 (Fig. 10F). These results directly suggested the involvement of K85 and R86 during Vn interaction.

Discussion

Vitronectin plays a crucial role in many biological systems including cell migration, adhesion, repair angiogenesis (Singh et al., 2010a) and microbial pathogenesis (Leroy-Dudal et al., 2004; Attia et al., 2006; Hallström et al., 2006; Bergmann et al., 2009; Hallström and Riesbeck, 2010; Singh et al., 2010a). It has a multi-domain structural arrangement, where the N-terminal SMB domain is well characterized for its role in several biological processes. The C-terminal HBD-1 (amino acids 175-219), HBD-2 (amino acids 175-219) and HBD-3 (amino acids 348-361) are known to bind to bacterial pathogens (Singh et al., 2010a). Being one of the major regulators of the complement system, Vn has a crucial impact on MAC deposition on the membrane of Gram-negative respiratory pathogens that successfully exploit the MACinhibitory role of Vn (Hallström et al., 2009: Singh et al., 2010a). On the other hand, Gram-positive bacteria have been shown to use the Vn-integrin interaction for cellular invasion and internalization (Bergmann et al., 2009).

Recently we reported that the PE⁸⁴⁻¹⁰⁸ domain is completely conserved in all Haemophilus spp. including 186 NTHi clinical isolates, Hib and other Haemophilus spp. The conserved property of PE among NTHi makes this protein valuable as a future vaccine candidate (Singh et al., 2010b). PE interacts with Vn at a $K_{\rm D} = 4 \times 10^{-7}$ M and protects NTHi from serum-mediated lysis. Vn bound to PE at the NTHi surface has a functional capacity to inhibit the MAC assembly, and this observation supports the involvement of a PE-Vn interaction in NTHidependent serum resistance. Furthermore, a peptide mapping analysis indicated that PE⁸⁴⁻¹⁰⁸ is the active Vn binding region (Hallström et al., 2009). In the present article, we focused on the properties of PE involved in binding to Vn. The PE84-108 region was studied in detail by using a site-directed mutagenesis approach. Interestingly, the PE⁸⁴⁻¹⁰⁸ region harbours K85 and R86, which are significantly involved in the function of PE. Mutation of K85 to the opposite charged side-chain E85, and R86 to D86 resulted in loss of PE binding to Vn. The mutated PE variants were also verified for any defective protein folding and tertiary structures by using CD spectroscopy, where PE^{WT}, PE^{R86D}, PE^{R86D} and PE^{K85, R86D} showed similar helix and beta-sheet patterns (Fig. 1E).

Previously it has been demonstrated that Vn binds to Streptococcus pneumoniae and contributes to bacterial internalization. This interaction can be efficiently blocked in the presence of heparin, indicating that the HBD(s) of Vn are involved in this interaction. Moreover, addition of Vn to epithelial cells significantly increases the adhesion of S. pneumoniae (Bergmann et al., 2009). Another pathogenic organism, Pseudomonas aeruginosa, has a potential binding capacity with Vn and binding of Vn with $\alpha\nu\beta5$ plays a key role in *P. aeruginosa* internalization by A549 cells (Leroy-Dudal et al., 2004; Leduc et al., 2007). H. influenzae binds Vn by the two outer membrane proteins Hsf and PE, which both contribute serum resistance (Hallström et al., 2006; 2009). However, Haemophilus ducreyi binds Vn via the trimeric autotransporter DsrA and thus exhibits serum resistance (Leduc et al., 2009). Many other research reports suggest that Vn potentially plays important roles in the pathogenesis of Yersinia pseudotuberculosis (Gustavsson et al., 2002), Neisseria meningitidis (Duensing and Putten, 1998; Sa et al., 2010) and Candida albicans (Limper and Standing, 1994). In light of the important role of Vn in microbial pathogenesis, there is not much data published that in detail characterize the interaction between Vn and its microbial ligands. The lack of structural data on the HBD(s) of Vn also limits the clear views about the Vn-bacterial interactions. In this study, Vn 353-363 amino acids region was identified as the major PE binding region. We designed a peptide library for the sequence Vn 312-396 and analysed for PE binding, but were not successful in defining the binding site (data not shown). Thus, there might be a secondary structure within the Vn 353-363 region that is involved in the interaction with PE that could not be mimicked by short synthetic peptides. Previously, it has been reported by us that heparin binding region (HBD-3) comprising Vn amino acids 341-368 might have an interaction with the PE molecule (Hallström et al., 2009). These data are in agreement with our previous findings indicating that Vn 353-363 are involved in binding to PE. The detailed alignment of Vn 320-396 has been published by us (Singh et al., 2010c) and reveals that the 353-363 amino acid region of Vn is variable among different mammalian species. This variation might be responsible for selective binding of Vn to pathogens, as reported recently with S. pneumoniae. Interestingly, mouse Vn binds several-fold less efficiently in comparison with human Vn (Bergmann et al., 2009).

Recently we reported that Vn plays a significant role in the survival of NTHi. The Vn-depleted serum decreases

the survival, whereas, addition of Vn restores serum resistance capacity (Hallström et al., 2009). M. catarrhalis also showed similar Vn-dependent serum survival, whereas Vn-depleted serum kills bacteria faster than NHS and addition of Vn restores serum resistance. UspA2 was found as the major Vn-binding protein of M. catarrhalis and UspA2-deficient mutants were highly sensitive to NHS (Attia et al., 2005; 2006). Based upon those earlier findings, we recently examined the UspA2-Vn interaction in detail. Interestingly, UspA2-treated human serum more efficiently killed bacteria as compared with control serum, and this was directly linked to guenching of Vn (Singh et al., 2010c). In parallel to Moraxella UspA2, our results on Haemophilus PE in the present study suggested that the enhanced killing of H. influenzae when NHS was pre-incubated with PE^{WT} as well with the single mutated variants PE^{K85E} or PE^{R86D} was due to the interaction and consequently quenching of the biological activity of Vn.

Previously we have shown that NTHi $3655\Delta pe$ is serum-sensitive and that the Vn-binding capacity significantly decreased in these PE-deficient mutants. The Vn depleted serum killed NTHi 3655 more efficiently in comparison with NHS and addition of Vn restored serum resistance of NTHi 3655 (Hallström *et al.*, 2009). In support of that study we showed in the present study that MAC deposition is significantly higher in NTHi $3655\Delta pe$ mutants in comparison with wild-type counterparts (Fig. 9A–E). Thus, the decrease in Vn binding to NTHi $3655\Delta pe$ is responsible for a considerably higher deposition of MAC.

In conclusion, this study suggests that the role of *H. influenzae* PE is important for Vn binding and consequently serum resistance. Lysine 85 and arginine 86 are located within the active binding region PE^{84–108} and are necessary for the Vn interaction. However, it is not completely evident that PE interacts with Vn by only K85 and R86. A future ultrastructure of the PE–Vn complex may prove details on the interaction. A detailed analysis of the Vn molecule and characterization of the precise amino acids that are responsible for the interaction with PE also remains to be done.

Experimental procedures

Bacterial strains, reagents and cell culture

The NTHi 3655 and an isogenic mutant (NTHi 3655 Δpe) were grown in brain heart infusion (BHI) liquid broth, supplemented with 10 µg ml⁻¹ nicotinamide adenine dinucleotide (NAD) and haemin, or streaked on chocolate agar plates and incubated at 37°C in a humid atmosphere with 5% CO₂. The NTHi 3655 Δpe mutant was grown in the presence of 20 µg ml⁻¹ kanamycin (Ronander *et al.*, 2009). *E. coli* BL21 (DE3) and DH5 α were cultured in Luria–Bertani (LB) broth or on LB agar plates at 37°C in a humid atmosphere containing 5% CO₂.

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The vectors pET26bpe and pET16bpe were as described (Hallström *et al.*, 2009). pET26bpe and variants were supplemented with 50 μ g ml⁻¹ kanamycin and pET16bpe and variants were supplemented with 100 μ g ml⁻¹ ampicillin in LB medium. Human embryo kidney (HEK293T) cells were grown in advanced DMEM (Gibco; Invitrogen, Stockholm, Sweden) with 2 mM L-glutamine, 100 μ g ml⁻¹ streptomycin and 100 U ml⁻¹ penicillin.

Recombinant DNA techniques and site-directed mutagenesis

The construct pET26bpe as described in Hallström et al. (2009) was used for expression and mutation of PE. The nucleotide region (250-324) of the pe gene (CGSHi3655_04936) encodes for PE⁸⁴⁻¹⁰⁸ and was manipulated at several amino acids by using a site-directed mutagenesis approach. The QuikChange site-directed mutagenesis was performed using high fidelity PfuTurbo® DNA polymerase (Stratagene, La Jolla, CA). The primers used for mutagenesis are listed in Table 1. The mutagenesis procedure including PCR and DpnI digestion was performed as described previously (Singh and Röhm, 2008). Mutant vectors were sequenced and finally transformed into E. coli BL21 (DE3) cells. The E. coli cells containing pET26bpe and variants were used for expression and purification, while E. coli containing pET16bpe was used for surface expression of PE (Ronander et al., 2009). Truncated Moraxella IgD-binding protein (MID962-1200) was prepared as described (Nordström et al., 2006).

Recombinant protein expression and purification

The E. coli BL21 (DE3) containing recombinant pET26b vectors were grown in 500 ml of LB medium with kanamycin at 37°C until OD600 to 0.8-1. Expression was induced by 1 mM IPTG and cultivation continued for another 3 h at 37°C. The cells were harvested at 5000 a for 15 min at 4°C. The periplasmic fraction was separated according to a protocol described in Singh and Röhm (2008). Purified proteins were dialysed by several exchanges of HNET buffer (50 mM HEPES, pH 7.5, 150 mM NaCl, 3 mM EDTA and 0.005% Tween-20) for Biacore assay and concentrated by using Centricon cartridge (3 kDa molecular weight cut-off; Millipore, Bedford, MA). Protein concentrations were measured by UV absorbance using a Nano-drop spectrophotometer (Thermo Scientific, Wilmington, DE) in addition to verification by a Bicinchoninic acid (BCA) assay (Pierce, Rockford, IL). The purity and concentrations of expressed proteins was verified on SDS-PAGE stained with Coomassie blue R250.

Expression of Vn fragments in HEK293 cell lines and purification

Vitronectin domains harbouring heparin binding sites have been expressed in HEK293T cells as Vn amino acids 80–396, 80–320 and 80–229 as described in Singh *et al.* (2010c). Furthermore, new constructs corresponding to amino acids 80–379, 80–373, 80–363, 80–353, 80–339 and

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Table 1. Primers used in site-directed mutagenesis of PE amino acids 84-108.

Primer name	Sequence 5'-3'	PE variants
K85E _For	GTTTATCCTGAGCCTGAACGTTATGCACGTTC	K85E
R86D _For	GTTTATCCTGAGCCTAAAGATTATGCACGTTCTGTTCG	R86D
Y87F _For	CCTGAGCCTAAACGTTTTGCACGTTCTGTTCGTC	Y87F
R89A_For	GAGCCTAAACGTTATGCAGCTTCTGTTCGTCAGTATAAG	R89A
R92A_For	GCACGTTCTGTTGCTCAGTATAAGATTTTGAATTGTGC	R92A
Y94F_For	GCACGTTCTGTTCGTCAGTTTAAGATTTTGAATTGTGC	Y94F
K95E_For	GTTCTGTTCGTCAGTATGAGATTTTGAATTGTGC	K95E
N98A_For	TCTGTTCGTCAGTATAAGATTTTGGCTTGTGCAAATTATCATTTAACTC	N98A
N101A _For	GTATAAGATTTTGAATTGTGCAGCTTATCATTTAACTCAAATACGAACTG	N101A
Y102A_For	GTATAAGATTTTGAATTGTGCAAATGCTCATTTAACTCAAATACGAACTG	Y102A
H103A_For	ATAAGATTTTGAATTGTGCAAATTATGCTTTAACTCAAATACGAACTGAT	H103A
Q106A_For	GAATTGTGCAAATTATCATTTAACTGCAATACGAACTGATTTCTATGATG	Q106A
R108A_For	GCAAATTATCATTTAACTCAAATAGCAACTGATTTCTATGATGAATTTTG	R108A
Mut 2_For	GTTTATCCTGAGCCTGAAGATTATGCACGTTC	K85E, R86D (mut 2)
Mut 3_For	GAGCCTGAAGATTATGCAGCTTCTGTTCGTCAGTATAAG	K85E, R86D, R89A (mut 3)
Mut 4_For	GCAGATTCTGTTGCTCAGTATAAGATTTTGAATTGTGC	K85E, R86D, R89A, R92A (mut 4)
Mut 5_For	ATTCTGTTGATCAGTATGAGATTTTGAATTGTGC	K85E, R86D, R89A, R92A, K95E (mut 5)

The mutated bases are underlined in all primers. The forward primer for each mutation is shown here, while reverse primer of same sequence was also used.

80–330, all containing C-terminal 6x-His tags (Fig. 4A) were amplified by using Vn 80–396 as template. The forward primer was inserted with BshT1 (AgeI) and the reverse primer with Acc651 (KpnI) restriction enzyme sites. Amplified and digested inserts were ligated into the pHLsec vector (28). Protein expression and purification was performed as described elsewhere (Singh *et al.*, 2010c).

Circular dichroism (CD)

Purified PE variants and Vn fragments were dialysed against 20 mM phosphate buffer and the concentrations were measured by using Nano-drop at (280 nm) and BCA method (Pierce, Rockford, IL). Finally, $5.0 \,\mu$ M proteins were transferred to a cuvette and CD spectrum was recorded between the wavelengths of 190 nm to 250 nm by using a CD spectropolarimeter J815 (Jasco, Essex, UK).

SDS-PAGE and non-reducing PAGE

Routine SDS-PAGE was performed according to standard laboratory protocols. In gradient separations NuPAGE Gradient 4–12% Bis-tris readymade gels were used (Invitrogen, Carlsbad, CA) and separation was performed according to protocol suggested by the manual. In non-reducing conditions protein samples were loaded with SDS-PAGE loading dye without β-mercaptoethanol.

Flow cytometry (FACS)

E. coli cells were grown in LB medium containing appropriate antibiotics and induced by 1 mM IPTG for 1 h at 37°C. Bacteria (10^8) were resuspended in 100 µl of PBS containing 2.5% BSA and treated with affinity-purified rabbit polyclonal anti-PE antibodies for 1 h at room temperature. After one washing with PBS-BSA, FITC-conjugated anti-rabbit antibodies (Dakopatts) were added (diluted according to the

manufacturer's instructions) and incubated for 1 h at room temperature. Additionally three washes performed with PBS containing 2.5% BSA and finally cells were resuspended in 100 μ I of PBS containing 2.5% BSA. *E. coli* harbouring empty vector was used as a control and treated similarly. Samples were analysed in a flow cytometer (EPICS XL_MCL; Beckman Coulter).

Immobilization of PE on CNBr-activated agarose beads

Cvanogen bromide-activated Sepharose[™] 4 Fast Flow (GE Healthcare Biosciences, Uppsala, Sweden) was used for immobilization of PE variants. Sepharose powder (330 mg ml-1) was washed several times in 1 mM HCl and resuspended in coupling buffer (0.1 M NaHCO₃, pH 8.3 containing 0.5 M NaCl). Protein solutions were also dialysed against coupling buffer and added to the resin (5-10 mg protein ml-1 of resin) and incubated overnight at 4°C in a rolling shaker. Unbound ligands were washed with 5 gel volume of coupling buffer and free groups were blocked by blocking buffer (0.1 M Tris-HCl. pH 8.0) for 2 h at room temperature. Beads were then washed three times by alternating buffers, NaAc, pH 4.0 containing 500 mM NaCl and 0.1 M Tris-HCl, pH 8.0 containing 500 mM NaCl. Finally beads were washed with PBS and stored at 4°C. The protein ligands immobilized were measured by Nano-Drop and BCA methods.

ELISA

Purified PE variants were coated on PolySorp microtitre plates (Nunc-Immuno, Roskilde, Denmark). Coating of PE (0.1 μ M) was performed in 100 mM Tris-HCl, pH 9.0 for 15 h at 4°C. Plates were washed three times with PBS to remove excess unbound protein and blocked with PBS-2.5% BSA for 1 h at 25°C. The Vn was added to the wells in PBS-2.5% BSA and incubated for 1 h at 25°C. In competitor ligand was added to the binding reactions and allowed to bind for 1 h at room

temperature. The unbound Vn fraction was removed by washing with PBS containing 0.05% Tween-20, and bound Vn was detected by polyclonal sheep anti-human Vn antibodies (pAb) (AbD Serotec, Kidlington, Oxford, UK) and secondary horseradish peroxidase (HRP)-conjugated donkey antisheep pAb (AbD Serotec). After four additional washing steps, plates were developed and read at 450 nm in a microplate reader (Multiskan[®] plus, Labsystems, Helsinki, Finland).

Surface plasmon resonance (Biacore)

The interaction between PE variants and Vn was analysed using surface plasmon resonance (Biacore 2000, Uppsala, Sweden). Four flow cells of a CM5 sensor chip were activated, each with 20 µl of a mixture of 0.2 M 1-ethyl-3-(3dimethylaminopropyl) carbodiimide and 0.05 M N-hydroxysulphosuccinimide at a flow rate of 10 µl min-1, after which Vn (10 µl ml⁻¹ in 10 mM sodium acetate buffer, pH 4.0) was injected over flow cell 2 to reach 4000 resonance units (RU). Unreacted groups were blocked with 20 µl of 1 M ethanolamine (pH 8.5). A negative control was prepared by activating and subsequently blocking the surface of flow cell 1. The comparative association kinetics were studied for various concentrations of the PE variants. The flow buffer (50 mM HEPES, pH 7.5 containing 150 mM NaCl, 3 mM EDTA and 0.005% Tween-20) was used for binding. Protein solutions were injected for 600 s during the association phase at a constant flow rate of 30 μ l min⁻¹. The sample was first injected over the negative control surface and then over immobilized Vn. The signal from the control surface was subtracted. The dissociation was followed for 200 s at the same flow rate. In all experiments, 30 µl of 2 M NaCl was used to remove bound ligands during a regeneration step. BiaEvaluation 3.0 software (Biacore) was used for data analysis and SigmaPlot 8.0 to generate final figures.

[125] labelling of Vn fragments and binding assay

The labelling of Vn was performed by the Chloramine-T method (25). For labelling, 0.05 M [¹²⁵]] isotope (Amersham Biosciences, Buckinghamshire, UK) was used per mole of protein. The labelled proteins were purified from unincorporated [¹²⁶] using PD10 columns (GE Healthcare Biosciences, Uppsala, Sweden).

NTHi and mutants were grown in BHI medium to $OD_{A600} = 1.0$ and washed with PBS-2.5% BSA. Approximately, 10⁸ bacterial cells per well were added to microtitre plates and increasing concentrations of [¹²⁵I]–Vn were added. For competition experiments, cold ligands were added to the binding reactions. Bacteria were incubated for 30 min at 37°C and were pelleted at 4200 *g* for 10 min, followed by two washing steps of PBS-2.5% BSA. The unbound radiolabelled protein was removed by three washes with PBS, harvested in a 96-well plate harvester (Tomtec, Hamden, CT), and finally counted in a liquid scintillation counter (Trilux, Microbeta 1450, Perkin Elmer).

In the radioactive ligand direct binding assay, PE (0.1 μ M) was coated on ELISA strip plate (MaxiSorp) and washed three times with PBS to remove excess unbound protein and blocked with PBS-2.5% BSA for 1 h at room temperature.

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[¹²⁵I]–Vn was added to the wells in PBS-2.5% BSA and incubated for 1 h at 25°C. In the competitive assay, competitor ligands were added to the binding reaction prior to addition of labelled Vn and allowed to bind for 1 h at room temperature. The unbound [¹²⁵I]–Vn fraction was removed by washing with PBS containing 0.005% Tween-20. Each single well was separated from strip by mechanical breaking and total radioactivity was counted in a scintillation counter.

Serum resistance assay

NTHi 3655 wild type was tested for impact of PE variants in NHS-mediated killing. Approximately, 10⁵ cells were resuspended in 100 µl of dextrose-GVB (DGVB⁺⁺) buffer, pH 7.3, containing 140 mM glucose, 0.1% (w/v) gelatin, 1 mM MgCl₂, 0.15 mM CaCl₂ and finally 10% NHS. Bacteria were incubated at 37°C and samples (10 µl) were collected at different time intervals and plated on chocolate agar plates. The effect of PE variants on serum proteins was analysed by pre-incubating increasing concentrations of PE variants ($0.5-5 \mu$ M) with 10% serum at room temperature for 1 h and subsequently used in the serum resistance assay. Similarly, the effect of Vn was evaluated by treating bacterial cells with Vn for 1 h at room temperature, prior to addition of NHS. The viable bacterial cells were determined by counting colony-forming units (cfu) after incubation overnight at 37°C.

Transmission electron microscopy (TEM)

Proteins were directly labelled with colloidal gold as described (Roth, 1996). The wild-type H. influenzae and mutants were grown in BHI for 3 h at 37°C. Bacterial solutions were incubated with gold-labelled Vn fragments (100 µg ml-1), fixed in PBS containing 4% paraformaldehyde and 0.1% glutaraldehyde and prepared for electron microscopy as described (Carlemalm, 1990). TEM was thereafter performed as described (Bengtson et al., 2008), and specimens were examined in a JEOL JEM 1230 transmission electron microscope (JEOL, Peabody, MA) at 60 kV accelerating voltage. Images were recorded with a Gatan Multiscan 791 CCD camera (Gatan, Pleasanton, CA). Only particles observed within a distance of 15 nm or less adjacent to the cell surface were counted. This corresponds to the established maximum distance between an IgG and its antigen, or between a protein labelled with gold of this size, and its target. Differences between groups concerning Vn-PE interaction, and complement activation were analysed by the onetailed test for differences between means. The following a priori null hypotheses were tested: full-length Vn does not bind to PE (Fig. 3), or full-length Vn binds to PE double point mutation with the same affinity (Fig. 3), or different Vn constructs exhibit the same affinity to PE (Fig. 7), or NTHi 3655 and NTHi Ape show the same C9 deposition (Fig. 9). The null hypothesis was rejected and statistical significance was assumed when $P \leq 0.05$.

Western blotting and MAC deposition assay

After processing, bacterial cells were boiled with 1× SDS-PAGE (containing β -mercaptoethanol) loading buffer for

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10 min at 95°C and centrifuged for 5 min at 20 000 g. Supernatants (10–15 μ l) were loaded in 12% or 15% SDSpolyacrylamide gels, or 4–12% gradient SDS-polyacrylamide gels (Invitrogen, Carlsbad, CA). The gels were subjected to run at particular current as recommended in standard protocols. Gels were blotted to PVDF membranes. Membranes were blocked with PBS containing 5% milk for 1 h at room temperature. All reactions were performed in PBS containing 5% milk and washing was done with PBS containing 0.05% Tween-20. Vn detection was performed with antihuman Vn antibodies (pAb) (AbD Serotec, Kidlington, Oxford, UK) and HRP-conjugated donkey anti-sheep secondary pAb (AbD Serotec).

Membrane attack complex deposition at the bacterial surface was estimated by C9 polymerization. Usually C9 monomer is a 71 kDa and it polymerizes into 14-16 mer tubules to make a pore at bacterial surface. In NHS C9 exists as a monomer or is associated in an oligomeric form, but when the complement reaction is initiated C9 is polymerized to make the ring closure (cytolytic pore). This C9 oligomer is stable and not dissociated in SDS containing buffer and reducing conditions, thus can be easily detected by SDS-PAGE (Podack and Tschopp, 1982; Deng et al., 2007). Bacterial cells (108) were taken from chocolate agar plates and resuspended into GVB++ buffer followed by addition of 10% NHS and incubation at 37°C. The effect of PEWT and $PE^{K85E, R86D}$ was evaluated by addition of 2 μM protein coated beads in 10% NHS and incubation at room temperature for 1 h. This treated serum was analysed for bactericidal activity and compared with non-coated control beads. The complement activity was stopped at indicated time points by cooling in an ice bath. Samples were boiled with SDS loading buffer (containing β-mercaptoethanol) for 10 min at 95°C and centrifuged for 5 min at 20 000 g. Supernatants were resolved in 4-12% gradient SDS-PAGE gels and blotted on to PVDF membranes. The detection of C9 was performed by using anti-human C9 pAbs (Complement Technology, Texas) and HRP-conjugated donkey anti-sheep secondary pAb as described above. Blots were finally developed with an ECL Western blotting kit (Pierce, Thermo Scientific, Rockford, IL).

Statistical analysis

Statistical analysis was performed by using Student's *t*-test for paired or unpaired data in SigmaPlot 8.0. ANOVA (one-way/two-way) analysis was performed by GraphPad Prism 5 (GraphPad Software, San Diego, CA). *P*-values \leq 0.05 were considered statistically significant. *P*-values are presented as **P* \leq 0.05, ***P* \leq 0.01, ****P* \leq 0.001 and n.s., not significant.

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Haemophilus influenzae acquires vitronectin via the ubiquitous Protein F to subvert host innate immunity

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Summary

Acquisition of the complement inhibitor vitronectin (Vn) is important for the respiratory tract pathogen nontypeable Haemophilus influenzae (NTHi) to escape complement-mediated killing. NTHi actively recruits Vn. and we previously showed that this interaction involves Protein E (PE). Here we describe a second Vn-binding protein, a 30 kDa Yersinia YfeA homologue designated as Protein F (PF). An isogenic NTHi 3655∆hpf mutant devoid of PF displayed a reduced binding of Vn, and was consequently more sensitive to killing by human serum compared with the wild type. Surface expression of PF on Escherichia coli conferred binding of Vn that resulted in a serum resistant phenotype. Molecular analyses revealed that the N-terminal of PF (Lys23-Glu48) bound to the C-terminal of Vn (Phe352-Ser374) without disrupting the inhibitory role of Vn on the membrane attack complex. The PF-Vn complex actively delayed C9 deposition on PF-expressing bacteria. Comparative studies of binding affinity and multiple mutants demonstrated that both PE and PF contribute individually to NTHi serum survival. PF was highly conserved and ubiquitously expressed in a series of randomly selected NTHi clinical isolates (n = 18). In conclusion, the multifaceted binding of Vn is beneficial for NTHi survival in serum and may contribute to successful colonization and consequently infection.

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Introduction

Nontypeable *Haemophilus influenzae* (NTHi) is a Gramnegative human-specific bacterium that is predominantly found as an upper respiratory tract commensal in up to 80% of healthy children (Foxwell *et al.*, 1998). NTHi is occasionally pathogenic and causes conjunctivitis, acute otitis media, sinusitis, and pneumonia. The species is also the leading cause of exacerbations in patients that suffer from chronic obstructive pulmonary disease (COPD) (Thanavala and Lugade, 2011). Furthermore, an increased incidence of invasive disease caused by NTHi has lately been observed (Resman *et al.*, 2011).

Colonization of healthy individuals with NTHi is restricted by the host immune system including the complement system (Zwahlen *et al.*, 1983; Toews *et al.*, 1985). The complement cascade is part of the innate immunity and serves as the first line of defence. It consists of more than 30 distinct plasma and cell bound proteins that form three major pathways: the classical, alternative and lectin pathway (Ricklin *et al.*, 2010). Activation of any pathway results in the formation of the membrane attack complex (MAC) consisting of C5b, C6, C7, C8 and C9, which form a pore in the cell membrane leading to cell lysis. Complement activation also results in an inflammatory process including opsonization, increased phagocytosis and recruitment of leucocytes.

Complement is involved in the mucosal immunity of the nasopharynx, the primary site of NTHi colonization. Activation of complement in the nasal mucosa has been implicated in the inflammatory and immunological processes of several respiratory tract diseases including allergic rhinitis, asthma, nasal polyposis, otitis media and COPD (Stenfors and Raisanen, 1992; Narkio-Makela et al., 1999; Mezei et al., 2001; Marc et al., 2004; Van Zele et al., 2009). Airway inflammation may induce increased mucosal permeability and leads to increased plasma exudation of complement proteins and antibodies into the airway (Stenfors and Raisanen, 1992; Narkio-Makela et al., 1999; Mezei et al., 2001). The role of plasma exudation as the first line of defence against infection and injuries in the respiratory mucosa has been extensively reviewed (Persson et al., 1998). Importantly, airway infection with NTHi has been shown to result in secretion of the proinflammatory cytokines interleukin (IL)-6, IL-8 and TNF- α

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leading to inflammation (Clemans *et al.*, 2000; Sharpe *et al.*, 2011). Moreover, NTHi-specific IgG exudate in the nasal fluid of infected animals effectively limited bacterial load in the nasopharynx (Zola *et al.*, 2009).

The complement system is tightly regulated to prevent self-damage from excessive activation. It is governed by inhibitors including C4b-binding protein (C4BP) in the classical/lectin pathways (Blom *et al.*, 2004), and factor H (FH) in addition to factor H-like protein 1 in the alternative pathway (Ferreira *et al.*, 2010). Clusterin and vitronectin (Vn) inhibit the formation of MAC in the terminal pathway (Morgan, 1999). Importantly, some of these inhibitors are also involved in the prevention of complement-dependent self-lysis at mucosal sites (Narkio-Makela *et al.*, 2001).

NTHi is generally susceptible to complement-mediated killing, which predominantly involves the C-reactive protein- and antibody-dependent classical pathway, although a minor role for the alternative pathway has also been reported (Weiser et al., 1998; Williams et al., 2001; Figueira et al., 2007). Several studies, however, show that most of the nasal mucosa and invasive NTHi isolates are capable of surviving in 5-40% normal human serum (NHS) (Williams et al., 2001; Figueira et al., 2007; Hallström et al., 2010). Serum resistance of H. influenzae is multifactorial and orchestrated by phase variation in lipooligosaccharides (LOS) (Williams et al., 2001; Figueira et al., 2007; Ho et al., 2007; Nakamura et al., 2011; Clark et al., 2012; Langereis et al., 2012) as well as through polymicrobial interactions that delay deposition of the complement components C3 and C4 (Tan et al., 2007). Acquisition of human complement inhibitors including plasminogen, C4BP, FH and Vn has recently also been reported for H. influenzae (Hallström et al., 2006; 2007; 2008; 2009; Barthel et al., 2012). NTHi surface adhesin Protein E (PE) and Haemophilus surface fibrils (Hsf) of serotype b (Hib) have been defined as Vn-binding proteins (Hallström et al., 2006; 2009).

Vn is a multifunctional glycoprotein present in the extracellular matrix (ECM), human plasma and in complex with epithelial cell integrins (Schvartz et al., 1999; Singh et al., 2010a). In human plasma, Vn is in a native monomeric (folded) conformation and multimerizes to its active (unfolded) form in serum (Stockmann et al., 1993). Vn inhibits the formation of cytolytic MAC by occupying the metastable membrane binding site of C5b-7 complex and hinders the insertion of the complex into the cell membrane, and thus prevents the completion of the lytic pore complex C5b-9. In addition, formation of soluble C5b-7 with Vn causes the complex to bind C8 and C9 and form soluble C5b-9, which is haemolytically inactive (Podack et al., 1984; Milis et al., 1993; Sheehan et al., 1995). Furthermore, selective masking of C5b and C8 by Vn has been proposed to interfere with the polymerization of C9 (Su, 1996). Activated (unfolded) Vn inhibits MAC and does not bind to C5b-7 (Hogasen *et al.*, 1992). Intriguingly, the bacterial Vn binding usually does not interfere with the Vn domain involved in inhibition of the MAC (Hallström *et al.*, 2009; Sa *et al.*, 2010).

We previously observed that the reduced Vn-binding capacity of an NTHi mutant devoid of PE was gradually increased when bacteria were incubated with high concentrations of Vn. This observation suggested an additional NTHi Vn-binding factor (Hallström et al., 2009). The goal of the present study was to define the total Vn-binding proteome using NTHi outer membrane vesicles (OMV). An additional outer membrane protein (OMP) was found to be important for NTHi-dependent Vn acquisition, and this newly identified Vn-binding protein we designated as Haemophilus Protein F (PF). In parallel with PE, PF promoted Vn-dependent bacterial adhesion and resistance against complement-mediated killing. The PF-Vn complex was active in delaying MAC formation on the bacterial surface. Importantly, PF was highly conserved and detected in all clinical NTHi isolates analysed.

Results

Nontypeable H. influenzae expresses multiple vitronectin-binding proteins

We have previously shown that NTHi binds Vn through PE (Hallström et al., 2009; Singh et al., 2011). The Vn-binding phenotype was not, however, completely abolished in the isogenic NTHi mutant devoid of PE. Therefore, we hypothesized that NTHi may express alternative Vn-binding OMPs. To identify these putative Vn-binding proteins. OMV fractions of NTHi 3655 were separated by two-dimensional gel electrophoresis (2D-SDS-PAGE) (Fig. 1A) and probed with human Vn in a Far western immunoblotting assay (Fig. 1C). In addition to LOS and DNA, the NTHi OMV contain periplasmic and OMPs (Sharpe et al., 2011). Our OMV proteome profile showed approximately 112 protein spots including the two major OMP P2 and P5. This was in agreement with the established outer membrane proteome of H. influenzae (Link et al., 1997). We found protein spots (~ 16 kDa to ~ 30 kDa) that bound Vn at different intensities (Fig. 1C) compared with the negative blot in the absence of Vn (data not shown). Due to the known difficulties of high molecular weight proteins to enter the IPG strips (Link et al., 1997), we also analysed the immunoblots by one-dimensional SDS-PAGE. Only one Vn-reactive protein band was observed (Fig. 1B and D), and was most likely the ~ 30 kDa Vn-binding protein seen in Fig. 1C. Of note, the ~ 16 kDa Vn-binding spot on Fig. 1C was the previously defined Vn-binding PE (Fig. S1), confirming a successful strategy used in the present study. The recruitment of Vn by NTHi thus involves at least two different proteins.



Identification of Haemophilus Protein F as a novel vitronectin-binding protein of NTHi

We further characterized the ~ 30 kDa NTHi Vn-binding protein using in-gel trypsin digestion and MALDI-TOF analysis. The protein spot was defined as an uncharacterized putative chelated iron-binding protein (GI: 145632610). The peptide ion covered 64% of the total protein sequence with Mascot score of 572, and no other statistically significant matches were identified. We designated this putative Vn-binding protein as Haemophilus Protein F (PF) (Fig. S2), which is a homologue of Yersinia pestis YfeA (74.9% of sequence identity; 86.7% of similarity). PF consists of 293 amino acid residues with a theoretical molecular weight of 32.4 kDa and a pl of 8.75. According to SignalP 3.0 (http://www.cbs.dtu.dk/services/ SignalP/), PF has a predicted 22-amino acid long signal peptide yielding a 30.1 kDa mature protein with the pl of 7.92 as seen in Fig. 1A.

We also wanted to analyse whether the *Haemophilus* PF was surface exposed on intact *H. influenzae*. The wild-type strain NTHi 3655 was mutated by replacing the *Haemophilus* PF gene (*hpf*) with a chloramphenicol resistance open reading frame (ORF) without interfering with the expression of other NTHi OMPs (Fig. S3). Using flow

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Fig. 1. Immunoblotting assay for identification of *Haemophilus* PF as a Vn-binding protein.

A. Separation of NTHi 3655 OMV (100 μg) on a 2D-SDS-PAGE (pH 3–10). Major OMP P2 and P5 are indicated with white arrows.
B. Separation of OMV (10 μg) in 1D-SDS-PAGE.

C. Far western blotting of the 2D gel with human Vn. Signals that represent Vn-binding PE (- 16 kDa, tagged with *) and PF (- 30 kDa, tagged with **) are indicated with black arrows in the Coomassie stained gel (A) and western blot (C). The open arrow indicates a background signal detected on a replica negative control blot (blot not shown). A negative control blot (blot not shown). any the primary and secondary antibodies without Vn. The background signal was attributed to the unspecific protein recognition by the antibodies.

D. Far western blotting of OMV separated on an 1D-SDS-PAGE. The arrow indicates the ~ 30 kDa protein that binds Vn. M, protein marker in kilo Dalton (kDa). A representative gel image and blot of two independent experiments are shown.

cytometry analyses, we found that rabbit anti-PF polyclonal antibodies (pAb) detected PF on NTHi 3655, whereas the protein was absent in NTHi 3655 Δ hpf (Fig. 2B). More importantly, complementing the *hpf* gene restored the PF expression at the bacterial surface to almost 70% of the wild type in NTHi 3655 Δ hpf (pKR7.2) (Figs 2B and S3). In support of these observations, transmission electron microscopy (TEM) using gold-labelled anti-PF pAb showed that PF was abundantly expressed on the outer surface membrane of NTHi 3655 (Fig. 2C), whereas only background staining was detected with the isogenic NTHi 3655 Δ hpf mutant (Fig. 2D). Taken together, our data corroborated that PF is a surface-exposed molecule.

NTHi devoid of PF displays a reduced binding to vitronectin

To ensure that the PF–Vn interaction identified in the Far western immunoblotting assay (Fig. 1C) was not an artifact due to non-native conformation of proteins presented in the 2D-gel electrophoresis, a direct binding assay was done with whole bacteria. At increasing concentrations of Vn (1–400 nM), flow cytometry analyses revealed that knocking out PF expression in the NTHi 3655 Δ hpf mutant

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Fig. 2. Detection of Haemophilus PF as a surface-associated protein on NTHi A and B. The presence of PF at the NTHi surface as measured by flow cytometry. A representative histogram of three independent experiments is shown. Numbers represent the mfi. Left panel (A) illustrates the background binding of the FITC-conjugated goat anti-rabbit pAb, whereas the right panel (B) shows detection by rabbit anti-PF pAb followed by incubation with the FITCconjugated goat anti-rabbit pAb. Based upon the relative mfi, PF expression in the transcomplemented NTHi 3655∆hpf (pKR7.2) mutant was restored to almost 70% of the wild type level, wt. NTHi 3655 wild type: Ahpf. PF-knockout mutant; Ahpf (pKR7.2), hpfcomplemented mutant. C and D. Visualization of PF on NTHi 3655 and NTHi 3655 ∆hpf by TEM. PF was detected by the gold-labelled rabbit anti-PF

detected by the gold-labelled rabbit anti-PF pAb that is visualized as black dots at the NTHi surface (C). Scatter dots presented in (D) was due to background labelling by antibodies on the solid matrix. Representative images of two independent experiments are shown.

resulted in significant reduction of the Vn-binding population (Fig. 3A) and mean fluorescence intensity (mfi) (Fig. 3B) as compared with the PF-expressing NTHi 3655 wild type. Importantly, trans-complementation of *hpf* with pKR7.2 in the mutant restored the Vn-binding to similar levels as the wild type (Fig. 3C).

In addition to the soluble form, Vn is also localized at the surface of epithelial cells or tethered within the ECM together with several other host proteins (Schvartz *et al.*, 1999). To evaluate the role of PF in adherence to immobilized Vn, we incubated both NTHi 3655 and the isogenic PF mutant on Vn-coated glass slides. NTHi $3655\Delta hpf$ adhered less efficiently to the Vn-coated glass slides compared with the wild type NTHi 3655 (Fig. 3D). In conclusion, we proved that the surface expressed PF has the capability to interact with both soluble and immobilized Vn.

Protein F binds vitronectin at the bacterial surface when expressed in Escherichia coli

To demonstrate the direct role of PF in Vn-binding on intact bacteria in the absence of other NTHi Vn-binding proteins, PF was recombinantly expressed at the surface of *E. coli* BL21(DE3), an isolate that does not bind Vn (Hallström *et al.*, 2009) (Fig. 4A). When the PF-expressing *E. coli* was incubated with increasing concentrations of Vn (5–1000 nM), a dose-dependent increase of the Vn-binding population was observed and bacteria were saturated at \geq 350 nM (Fig. 4B). In contrast, control *E. coli* transformed with the empty vector pET-16b bound

less Vn. We further validated the specificity of Vn binding to PF-expressing *E. coli* by the ability of anti-PF pAb to block the interaction. A significantly reduced binding was observed at a dilution of up to 1:50 compared with the control incubated in the absence of antibodies (Fig. 4C). In parallel, the adherence of PF-expressing *E. coli* to immobilized Vn was increased compared with the control *E. coli* (Fig. 4D). Taken together, we showed that PF confers a Vn-binding property to the heterologous host *E. coli* providing further evidence of PF being a significant Vn-binding NTHi OMP.

The vitronectin-binding region is located within Lys23-Glu48 at the N-terminus of Protein F

To pinpoint the Vn-binding domain of PF, a series of polyhistidine-tagged truncated fragments (Fig. 5A) spanning the entire polypeptide of PF (amino acids 12-293) were recombinantly produced in E. coli BL21(DE3). As quantified by ELISA, full-length PF12-293 bound Vn in a dose-dependent manner when incubated with increasing concentrations of Vn (1-50 nM) (Fig. 5B). The dissociation constant (K_D) of the interaction was calculated to 12.82 nM. Thereafter, Vn at pre-saturated concentrations (5 nM and 10 nM) were used for incubation with immobilized full-length and truncated fragments of PF. We found that two fragments encompassing mainly the N-terminus of PF, i.e. PF¹²⁻⁹⁸ and PF¹²⁻¹⁷⁷, showed similar Vn binding to that observed with the full-length PF12-293 (Fig. 5C). An uncharacterized NTHi protein, UHP 09011 (GI: 145632833), which does not bind Vn, was included as a

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Fig. 3. The NTHi mutant devoid of PF has a reduced Vn-binding.

A. Analysis of the Vn-binding population (%) of NTHi 3655 wild type and △hpf mutant at increasing concentrations of Vn (1–400 nM) as measured by flow cytometry.

B. A representative flow cytometry histogram of NTHi-dependent binding of Vn at 100 nM. Numbers represent mfi of the corresponding experiment. A background control without Vn added is also included (black). Three independent experiments were done.

C. Restoration of the Vn-binding phenotype in an *hpf*-trans-complemented mutant. Binding was quantified as mfi by flow cytometry (n = 3). wt, NTHi 3655 wild type; Δhpf , mutant devoid of PF; Δhpf (pKR7.1), trans-complemented control mutant; Δhpf (pKR7.2), *hpf* trans-complemented mutant. For (A) and (C), data represent the mean of three independent experiments and error bars represent standard deviations. Two-way anova was used to calculate the differences in Vn binding between the NTHi 3655 strains. ** $P \le 0.01$; n.s., not significant. D. Adherence on Vn-coated glass slides of NTHi 3655 wild type and the PF-knockout mutant. Bacteria were visualized under light microscope after a standard Gram-staining. Representative images of two independent experiments are shown.

negative control. The specificity of the protein–protein interactions was further confirmed in a competition assay. Pre-incubation of Vn with increasing concentrations of PF¹²⁻²⁹³ or PF¹²⁻⁹⁸ (0.05–5.0 μ M) resulted in a significant decrease of Vn binding to the immobilized PF¹²⁻²⁹³ (Fig. 5D). The fragment PF⁸¹⁻²⁹³ that was devoid of the N-terminus, and the negative control UHP_09011 did not inhibit the interaction.

The exact Vn binding site on the PF molecule was further revealed with a series of synthetic peptides constituting of 23–26 amino acid residues, each with four amino acids overlap (Fig. 5E). Among the peptides, we found that only PF²³⁻⁴⁸ reacted preferentially with Vn both at 5 and 20 nM (Fig. 5F). The K_D value for the interaction was estimated to 16.36 nM (Fig. 5G), which is comparable with the K_D value of PF^{12–293}. In a competition assay (Fig. 5H), pre-incubation of Vn with increasing concentrations of the PF²³⁻⁴⁸ peptide significantly reduced Vn binding to immobilized PF^{12–293}. Neither the peptide PF^{44–68} nor PF^{225–255} competed, however, with the fulllength recombinant PF. To summarize, our data indicated that the N-terminal (Lys23-Glu48) is the major Vn-binding region of PF and that the interaction is specific as revealed by competition assays.

Protein F interacts with the C-terminus of vitronectin

We also wanted to identify the putative PF binding site on the Vn molecule (Fig. 6A). Heparin and plasminogen activator inhibitor-1 (PAI-1) have multiple distinct binding sites throughout the Vn molecule (Lane *et al.*, 1987; Kost *et al.*, 1992; Liang *et al.*, 1997; Gibson *et al.*, 1999). PAI-1 has one primary binding site at the SMB domain of the N-terminus and a secondary binding site at the C-terminus of Vn (Schar *et al.*, 2008a,b) (Fig. 6A). Among the three heparin-binding domains (HBD-1 to 3), only

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Fig. 4. E. coli-expressing Haemophilus PF at the surface acquires Vn binding.

A. Surface expression of *Haemophilus* PF by the heterologous host *E. coli* BL21(DE3) as measured by flow cytometry with anti-PF pAb. Histogram of a single representative experiment from three independent experiments is shown and numbers represent mfi. Background binding by the FITC-conjugated secondary pAb in the absence of vitronectin is included.

B. Percentage of the Vn-binding population in *E. coli* strains (*E. coli*-expressing PF and control *E. coli* containing the expression vector pET-16b only) at various concentrations of Vn (5–1 × 10³ nM) as quantified by flow cytometry. Differences in the Vn-binding population observed between the PF-expressing *E. coli* and control *E. coli* were estimated by a two-way ANOVA.

C. Anti-PF pAb specifically block vitronectin-binding to PF. Data show percentages of the bacterial binding population to 120 nM Vn after pre-incubation with various dilutions of anti-PF pAb followed by flow cytometry analyses. Statistical significances regarding differences between antibody blocking for each strain compared with the conditions without antibody were determined by an one-way ANOVA. For (B) and (C), data represent the mean of three independent experiments and error bars show the standard deviations. * $P \le 0.05$; ** $P \le 0.01$; ** $P \le 0.001$.

D. Bacterial attachment on a Vn-coated glass slide. Adherence was visualized under a light microscope after conventional Gram-staining. A representative image of two independent experiments is shown.

HBD-3 [amino acids (aa) 348–361] is known to overlap with the secondary PAI-1 binding site (aa 348–370) located at the C-terminus of Vn (Kost *et al.*, 1992; Schar *et al.*, 2008a,b). The majority of bacterial surface proteins known to date interact with the C-terminus of Vn (Singh et al., 2010a,b; 2011). Therefore, we investigated the PF binding site on the Vn molecule by using heparin and PAI-1 as blocking agents. We determined that both

Fig. 5. N-terminal Lys23-Glu48 of PF is involved in the Vn binding.

A. Schematic representation of PF recombinant truncated fragments, and (E) synthetic peptides encompassing the entire PF polypeptide. SP, signal peptide.

B and G. Estimation of the K_D value of the (B) PF¹²⁻²⁹³ fragment and (G) PF²³⁻⁴⁸ peptide in interaction with Vn (1–50 nM) by ELISA. The K_D and B_{MAX} values are shown as average values with standard deviations from three independent experiments. Small inserts represent the Scatchard plots.

C and F. Binding of Vn to truncated PF fragments (C), and PF peptides (F).

D and H. Inhibition assays of the PF-Vn interaction by (D) PF fragments and (H) PF peptides. IC₅₀, the 50% inhibitory concentration of each fragment and peptide.

All data represent the mean of three independent experiments and error bars show standard deviations. The increased Vn binding observed for the PF fragments or peptides compared with the negative control (UHP_09011); or inhibition in Vn binding compared with the absence of inhibitor was statistically significant when calculated by the two-way ANOVA test. $*P \le 0.05$; $**P \le 0.01$ and $***P \le 0.001$. n.a., not analysed.



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Fig. 6. Mapping of the PF binding site on the Vn molecule.

A. Schematic representation of the identified active ligand-binding domains of Vn. SP, signal peptide. Recombinant fragments of Vn are shown in black blocks and deleted regions in mutated variants are indicated in grey (Vn deletion mutants).

B and C. Inhibition of the Vn-PF interaction by heparin (6.7 × 10⁻⁶–6.7 × 10⁴ nM) (B) and PAI-1 (0.1–200 nM) (C). Vn (10 nM) was pre-incubated with various concentrations of inhibitors. Reaction mixtures were thereafter incubated with PF¹²⁻²⁹³ immobilized on microtitre plates. IC₅₀, the 50% inhibitory concentration of heparin or PAI-1.

D. Interaction of immobilized PF¹²⁻²⁹³ and PF²³⁻⁴⁸ to Vn variants (10 nM). The non-binding fragment (PF⁸¹⁻²⁹³) and peptide (PF⁴⁴⁻⁶⁸) were included as negative controls. An enhanced binding of truncated Vn fragments to the immobilized PF¹²⁻²⁹³ and PF²³⁻⁴⁸ was statistically significant as calculated by the two-way ANOVA when compared with the negative fragment and peptide respectively. * $P \le 0.05$; ** $P \le 0.01$ and ** $P \ge 0.001$.

E and F. Characterization of the ionic interaction of PF protein fragments (E) and synthetic peptides (F) with Vn (10 nM) in the presence of various concentrations of NaCI (0.1–1.0 M). Vn-binding was measured in ELISA.

All data represent the mean of three independent experiments and error bars show the standard deviations.

heparin and PAI-1 inhibited binding of PF¹²⁻²⁹³ to Vn in a dose-dependent manner (Fig. 6B and C) that indicated a potential overlapping Vn binding site. The results hence suggested that PF targets HBD-3 and the C-terminal PAI-1 binding site of the Vn molecule.

To support the observations outlined above, a series of truncated recombinant Vn fragments that were partially or completely devoid of HBD-3 and the C-terminal PAI-1 binding site were constructed and used to analyse binding to immobilized PF by ELISA (Fig. 6A). As can be seen in Fig. 6D, Vn fragments (n = 4) (comprising aa 80–229, 80–320, 80–330, and 80–353 respectively) did not interact with PF, whereas the other three Vn fragments comprising aa 80–363, 80–373, and 80–396 bound recombinant PF^{12–293} and the synthetic peptide PF²³⁻⁴⁸. Two PF constructs devoid of the Vn-binding regions (PF^{81–293} and PF^{44–68}) were included as negative controls. These data indicated that aa spanning Vn 80–353 were not involved in interactions with PF, whereas the aa Vn 363–396 were crucial.

In order to further determine the PF binding region on the Vn molecule, we produced a series of Vn deletion mutants devoid of the specified C-terminal amino acid residues (Vn Δ 352–362, Vn Δ 362–374 and Vn Δ 352–374) (Fig. 6A; lower part). Both Vn Δ 352–362 and Vn Δ 362–374 bound recombinant PF in ELISA while extensive deletion of 20 amino acids between residues 352–374 (Vn Δ 352– 374) caused a significantly reduced interaction with PF (Fig. 6D).

It is known that the interactions of heparin and PAI-1 at the polycationic C-terminus of Vn is sensitive to ionic strength (Gibson *et al.*, 1999; Schar *et al.*, 2008a,b). In parallel, we found that NaCl (titration 0.05–1.0 M) inhibited Vn binding to the immobilized recombinant fragments PF¹²⁻²⁹³ and PF¹²⁻⁹⁸ (Fig. 6E) and the PF²³⁻⁴⁸ peptide (Fig. 6F). A hyper-physiological concentration of salt (≥ 0.2 M) was required to disrupt and decrease the interaction between Vn and immobilized PF. In conclusion, PF (aa 23–48) directly interacts with the C-terminal part of Vn (aa 352–374) and the binding is based upon ionic interactions. Protein F enhances bacterial serum resistance through the recruitment of Vn

Vn inhibits the formation of the MAC at the terminal lytic step of the complement cascade. Bacteria that recruit Vn would therefore be resistant against the bactericidal activity of NHS (Singh *et al.*, 2010a). Based upon our findings that PF has the ability to bind Vn, we subsequently investigated the role of PF in bacterial serum resistance. Deletion of PF resulted in a greater sensitivity to complement-mediated killing of NTHi $3655\Delta hpf$ compared with the wild type NTHi 3655 (Fig. 7A). Importantly, trans-complementation of the isogenic mutant NTHi $3655\Delta hpf$ with pKR7.2 containing the *hpf* gene restored the serum-resistant phenotype (Fig. 7B).

To reveal the direct role of PF in serum resistance in the absence of other NTHi proteins involved in subverting complement-mediated killing (Ho et al., 2007; Hallström et al., 2009: Clark et al., 2012: Langereis et al., 2012). PF-expressing E. coli was tested with 1% NHS. As shown in Fig. 7C, the presence of PF at the E. coli surface significantly promoted survival in NHS compared with control E. coli carrving the empty expression vector. Interestingly, we found that both the control E. coli and PF-expressing E. coli were equally susceptible to serummediated killing when Vn was depleted from the NHS (Fig. 7C; blue curves). In parallel, a similar increased bactericidal activity was observed with the NTHi wild type and the hpf-deleted mutant (Fig. 7A; blue curves). Importantly, when exogenous Vn (\geq 72 nM) was supplemented we managed to rescue PF-expressing E. coli and NTHi from serum-mediated killing (Figs 7D and S4A). This Vn concentration represented conversion of serum Vn in approximately 1% NHS (400 µg ml-1; 5.3 µM). Heat inactivation of NHS (HIS) abolished the serum bactericidal activity in all experiments (Fig. 7A and C; red curves). In parallel with the data shown in Fig. 4C, blocking the Vn-PF interaction at the bacterial surface with anti-PF pAb further decreased the serum resistance in both E. coli-expressing PF and wild type NTHi (Fig. S4B and C). Taken together, we demonstrated that PF significantly

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Fig. 7. Enhanced serum resistance is exerted by PF-expressing bacteria.

A. Survival kinetics of NTHi in 5% normal (NHS), 5% Vn-depleted serum and HIS.

- B. Restoration of the serum survival phenotype of the hpf-complemented mutant in 5% NHS for 10 min. wt, NTHi 3655 wild type; Δhpf, mutant devoid of PF; \(\Delta\)hpf (pKR7.2), hpf trans-complemented strain.
- C. Survival kinetics of E. coli strains in 1% NHS, 1% Vn-depleted serum and 5% HIS.
- D. Serum resistance of E. coli strains in 1% Vn-depleted serum that was supplemented with various concentrations (36-180 nM) of purified Vn. Bacterial survival was determined at 5 min.
- E. The serum Vn-binding population as measured by flow cytometry.
- F. Survival of E. coli in serum after pre-incubation with various concentrations of HIS (0.1-15%).
- G. Deposition of serum Vn, C4BP and FH on E. coli strains following pre-incubation with 1% HIS.

All data represent the mean values of three independent experiments and error bars indicate the standard deviations. Statistical differences in serum resistance or in the ligand-binding population observed between the strains of NTHi 3655 or E. coli were calculated by the two-way ANOVA. * $P \le 0.05$; ** $P \le 0.01$ and *** $P \le 0.001$, n.s.; not significant.

contributes to NTHi-dependent serum resistance via Vn bindina.

To evaluate whether Vn was the primary complement inhibitor responsible for our observations, we preincubated E. coli-expressing PF and control E. coli with various concentrations of HIS to assess the bacterial recruitment of serum Vn in addition to other potential complement inhibitors. Heat treatment of serum does not abolish the inhibitory activities of the majority of soluble complement inhibitors, including Vn (Zhuang et al., 1996; Kask et al., 2004). We found that following pre-incubation with increasing concentrations of HIS, a dose-dependent Vn binding was observed with the PF-expressing E. coli population (Fig. 7E). This also reflected the increased concentration of serum Vn deposited at the surface of PF-expressing bacteria hence significantly improving the survival in NHS (Fig. 7F). Control E. coli did not exert any enhanced Vn binding or serum resistance.

We previously reported that the recruitment of C4BP, FH and Vn is required for NTHi-dependent serum resistance (Hallström et al., 2007; 2008; 2009). To avoid background binding by other NTHi-derived proteins, E. coliexpressing PF therefore was used to examine the direct interaction of PF with serum C4BP and FH. Recruitment of serum C4BP or FH derived from HIS was indistinguishable between E. coli-expressing PF and control E. coli as revealed by flow cytometry (Figs 7G and S5A) and ELISA (Fig. S5B), and did not correlate with the bacterial survival in serum. Collectively, our data proved that PF-dependent Vn binding at the bacterial surface thus contributes to bacterial serum resistance.

The vitronectin-PF complex inhibits MAC formation and delays deposition of C9 on bacteria

To ensure that the interaction of PF at the C-terminus of Vn would not interrupt or sterically alter the MAC inhibitory role of Vn, we performed a haemolytic assay with sheep erythrocytes and terminal complement proteins C5b-6, C7, C8 and C9. As shown in Fig. 8A, Vn that was preincubated with increasing concentrations of recombinant $\text{PF}^{\text{12-293}}$ (0.25–1 $\mu\text{M}\text{)}$ remained active resulting in inhibition of the cytolytic activity executed by C5b-9. The data indicated an exclusive role of the PF-Vn complex in the inhibition of complement-mediated lysis that was not attributed to recombinant PF12-293 alone.

We further characterized the effect of the PF-Vn complex on C9 deposition and polymerization at the bacterial surface. When NHS was incubated with the mutant NTHi 3655∆hpf devoid of PF expression, an increased C9 polymer deposition (> 250 kDa) was observed after 40 min compared with the NTHi 3655 wild type (Fig. 8B). C9 deposition at the bacterial surface was quantified by flow cvtometry and is shown in Fig. 8C. Moreover, we found that pre-incubation of whole bacteria with increasing concentrations of Vn prevented C9 deposition on NTHi 3655, but was unable to protect NTHi 3655∆hpf (Fig. 8D and E). We therefore concluded that PF-dependent serum resistance requires Vn to delay C9 deposition.

Protein F and Protein E share homologous binding sites on vitronectin and are both important for NTHi serum resistance

PE is hitherto the only Vn-binding protein of NTHi reported and it is also known to contribute to serum resistance (Hallström et al., 2009; Singh et al., 2011). We therefore compared the relative Vn-binding affinity between Haemophilus PE and PF. As shown in Fig. 9A, purified recombinant PE interacted with Vn more effectively than recombinant PF ($P \le 0.01$). The K_D value for the PE–Vn interaction as quantified by ELISA was calculated to 10.13 nM of Vn (Fig. S6), demonstrating a slightly higher affinity for Vn binding compared with PF ($K_D = 12.82$ nM). Pre-incubation of Vn with increasing concentrations (0.05–1.00 μ M) of PE or PF resulted in the depletion of free Vn to immobilized PF or PE respectively (Fig. 9B). As PE was also reported to bind Vn at HBD-3 (Singh et al., 2011), our data thus demonstrated that PE and PF may share a common binding site on the Vn molecule.

We further investigated the relative Vn-binding role of PE and PF using whole bacteria. While NTHi 3655∆hpf and NTHi 3655 Ape individually exerted a low binding of Vn, we found that simultaneous mutation of both pe and

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hpf further reduced the Vn binding by NTHi (Fig. 9C), albeit it was statistically insignificant. A slightly increased sensitivity to serum killing and C9 deposition was also observed with the double mutant (NTHi 3655∆pe/hpf) compared with the single mutants (Fig. 9D and E). The data thus confirmed that both PE and PF are important for Vn binding and the consequent serum resistance of NTHi.

Fig. 8. The active inhibition of MAC formation and delayed C9 deposition by the Vn-PF complex.

A. The haemolytic inhibitory activity of Vn in the presence of recombinant PF¹²⁻²⁹³ (0.25–1.0 µM). C4BP, FH, or buffer was included as negative controls. The difference in the haemolytic activity by MAC between the presence or absence of Vn was statistically significant as revealed by one-way ANOVA.

B. Western blotting analyses of C9 deposition on NTHi 3655 and NTHi 3655 Δhpf when incubated in 10% NHS at various time points. C9 was detected as high molecular weight polymers (> 250 kDa), trimers (~ 250 kDa) and monomers (~ 72 kDa). Activation of complement in NHS results in multimerization of C9 monomers into a trimeric and thereafter a polymeric form deposited on bacterial cells. However, only the polymeric C9 consequently makes up the cell-lytic MAC.

C. Quantification of the C9 polymer deposition as shown in (B) by flow cytometry. Bacteria incubated without NHS were included as a background control.

D. Effect on C9 deposition between strains of NTHi 3655 after pre-incubation (30 min at 37°C) with various concentrations of Vn

 $(0.01-0.5 \,\mu\text{M})$ prior to 30 min incubation with NHS. C9 polymer deposition was analysed by western blotting.

E. Quantification of the C9 polymer deposition as shown in (D) by flow cytometery.

For (C) and (E), data are presented as mean fluorescence intensities (mfi). All numerical data are the mean of three independent experiments and error bars represent the standard deviations. The difference of C9 polymer deposition observed between NTHi 3655 wild type and the PF mutant was statistically significant as determined by the two-way ANOVA. $*P \le 0.05$ and $***P \le 0.001$.

Protein F is a highly conserved Vn-binding protein in NTHi clinical isolates

We wanted to determine whether PF expression, and thus PF-mediated Vn binding, is conserved within the NTHi species. Flow cytometry analyses demonstrated that the majority of NTHi clinical isolates bound Vn (Fig. 10A). In addition, PF was ubiquitously expressed by all the isolates tested (Fig. 10B). Protein sequence alignment of PF from 18 different NTHi clinical isolates exerted 97.9-100% of sequence identity indicating that PF is a highly conserved protein. The sequence alignment is shown in Fig. S7. Importantly, we also found that knocking out hpf in the randomly selected NTHi isolates KR271, KR316, KR336 and KR385 (Fig. S8) resulted in a significantly reduced affinity for Vn (Fig. 10C), and consequently an increased sensitivity to the bactericidal effect of serum (Fig. 10D) when compared with the parental wild types. Taken together, PF is a highly conserved and ubiquitous surface protein mediating NTHi serum resistance.

Discussion

Inhibition of serum bactericidal activity by recruiting Vn is important for NTHi (Eberhard and Ullberg, 2002; Hallström *et al.*, 2009; Ronander *et al.*, 2009; Singh *et al.*, 2011). In this paper we defined PF as a novel Vn-binding protein in addition to the well-characterized PE (Hallström *et al.*, 2009; Singh *et al.*, 2011). Acquisition of Vn by NTHi is thus depending on more than one OMP and represents an important survival strategy to evade complementmediated killing. This multifaceted strategy has been reported to be essential for several serum-resistant pathogens, for example, Vn and FH binding in *Neisseria meningitidis* (Lewis *et al.*, 2010; Sa *et al.*, 2010; Griffiths *et al.*, 2011) and C4BP recruitment in *Yersinia enterocolitica* (Kirjavainen *et al.*, 2008).

Outer membrane vesicles are versatile cargo-containing vehicles that bud off from the outer membrane of Gram-negative bacteria. OMV released by NTHi contain mainly virulence-associated periplasmic and OMPs that are involved in the pathogenesis (Kahn *et al.*, 1983; Wispelwey *et al.*, 1989; Sharpe *et al.*, 2011). Therefore, the use of OMV (instead of whole cell bacteria) for the global screening of NTHi Vn-binding OMPs was merited in this study. Furthermore, the use of OMV excludes potential problems associated with the cytoplasmic compartment of whole bacterial cells that could potentially result in background binding.

All microbial Vn receptors studied to date are surfaceassociated virulence factors (Singh *et al.*, 2010a). Recognition of PF on NTHi and *E. coli*-expressing PF by anti-PF pAb revealed the surface localization in flow cytometry and TEM (Fig. 2). In parallel, a PF homologue of *H. parasuis* (GI: 219871287) (83.1% sequence similarity and 74.7% identity with NTHi PF) was also found at the bacterial surface and shown to be immunogenic in an animal model (Hong *et al.*, 2011).

We presented significant evidence that PF is a Vnbinding factor at the surface of NTHi based upon analyses with NTHi *hpf* mutants (Fig. 3), surface expression of PF in *E. coli* (Fig. 4) and protein–protein interaction assays (Fig. 5). In addition, the capacity of PF-expressing bacteria to bind immobilized Vn implied a potential adhesive role for PF. By using Vn as a bridging molecule for attachment to integrins (e.g. $\alpha_v\beta_3$) on host cells, PF may potentially mediate NTHi colnization at the epithelium of the respiratory tract (Lopez-Gomez *et al.*, 2012). This invasion strategy has also been demonstrated for other mucosal pathogens (Bergmann *et al.*, 2009; Sa *et al.*, 2010; Singh *et al.*, 2010a).

In silico protein structure modelling of PF (Fig. S9) demonstrated that the Lys23-Glu48 region is exposed at the protein surface thus supporting the identification of the Vn-binding domain at the N-terminus of PF (Fig. 5). Here, we also revealed that PF targets the C-terminal region (Phe352-Ser374) of Vn containing the entire HBD-3 and C-terminal PAI-1 binding side (Fig. 6). This is in agree-

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ment with previous work demonstrating that heparin inhibited the *H. influenzae* Vn binding (Eberhard and Ullberg, 2002). Interestingly, lung infection with NTHi triggers the upregulation of PAI-1 as a part of the host early immune defence by recruitment of polymorphonuclear neutrophils (Lim *et al.*, 2011). The role of an interaction between PAI-1 and Vn when bound to the NTHi surface remains, however, to be elucidated.

Antibody-depleted serum suggested that the classical pathway dominated the killing of the PF-knockout mutant

Fig. 9. Involvement of PE and PF in Vn-binding and the serum resistant phenotype of NTHi.

A. Comparison of Vn binding between recombinant PE and PF when measured by ELISA. Purified UHP 09011 was included as a negative control

B. Inhibition assay of Vn binding to the immobilized PE and PF in the presence of increasing concentrations (0.05-1.00 µM) of PF or PE respectively

C. Direct ligand binding assay with NTHi 3655 strains and radiolabellled [1125]-Vn. Radioactivity (count per minute; CPM) represents the concentration of Vn bound to bacteria. The NTHi 3655 \(\Lambda\) hap mutant and E. coli were included as negative controls. Hap is an autotransporter with adhesive properties that binds epithelial cells and laminin/collagen II but not Vn (Fink et al., 2002). D. Survival of NTHi 3655 strains in 5% NHS for 10 min.

E. C9 deposition on NTHi 3655 strains when incubated in 10% NHS for 40 min at 37°C. C9 deposition was quantified by flow cytometry. In panels (C), (D) and (E): wt, NTHi 3655 wild type; Δhpf , mutant devoid of PF; Δpe , mutant devoid of PE; $\Delta pe/hpf$, PE and PF double-knockout mutant; Ahap, mutant devoid of Hap. All data represent mean values of three independent experiments and error bars indicate standard deviations. Differences between proteins or bacterial strains regarding Vn binding and serum survival were calculated by one-way ANOVA. * $P \le 0.05$; ** $P \le 0.01$ and *** $P \le 0.001$ respectively.

(Fig. S4D). However, the relatively greater serum resistance observed in PF-expressing NTHi 3655 occurred via delayed C9 deposition during MAC formation at the terminal lytic step (Figs 7 and 8). Importantly, bacterial survival was related to PF-dependent recruitment of Vn. The

serum resistance was restored after (i) introduction of the hpf gene in the PF-knockout mutant, and (ii) Vn addition to Vn-depleted serum. Active interaction of cell-associated PF with Vn in the fluid phase suggested a potential role of PF to scavenge Vn from either the blood stream during



Fig. 10. PF is a conserved Vn-binding protein important for serum resistance in NTHi clinical isolates

A. A series of NTHi clinical isolates (n = 18) were analysed for Vn binding. Binding (mfi) was measured by flow cytometry. Background binding represents experiments performed in the absence of Vn

B. Detection of PF expression on NTHi clinical isolates by anti-PF pAb in western blotting. Grey and black arrows indicate purified recombinant and native PF respectively. CB, Coomassie blue stained SDS-PAGE; WB, western blotting. A representative gel image and blot of two independent experiments is shown. C. Vn binding of PF-knockout mutants of NTHi KR271, KR316, KR335 and KR385 clinical isolates. Binding was measured by flow cytometry.

D. Survival of NTHi clinical isolates and their respective mutants devoid of PF in 5% NHS for 10 min.

In panel (C) and (D): wt, NTHi wild type; Δhpf , PF mutant. For panel (A), (C) and (D) data represent mean values of three independent experiments and error bars indicate standard deviations. Differences in Vn-binding and serum survival between the NTHi wild types and PF mutants were calculated by a two-way ANOVA. * $P \le 0.05$; ** $P \le 0.01$ and *** $P \le 0.001$.

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invasive sepsis or from the pericellular ECM during respiratory tract infection. We found that PF binding did not interfere with the C5b-C9 inhibitory domain (Met51-Thr310) of Vn (Sheehan *et al.*, 1995) (Fig. 8). We therefore excluded the possibility of disruption of the MAC inhibition site or a conformational change of the Vn molecule after binding to PF. Delayed MAC formation on serum-resistant NTHi isolates can also be attributed to delayed deposition of antibodies or C3b and C4b (Figueira *et al.*, 2007; Ho *et al.*, 2007; Clark *et al.*, 2012; Langereis *et al.*, 2012), and host deficiencies in C3, C4 and C5 production (Toews *et al.*, 1985). We do not exclude the possibility that PF may also affect the activation or deposition of other components important in the classical complement pathway.

A functional similarity between PE and PF was observed regarding binding to the HBD-3 and C-terminal PAI-1 binding sites of Vn (Singh et al., 2011). The Vn-binding domain derived from both PE (aa 84-108) (Hallström et al., 2009; Singh et al., 2011) and PF (aa 23-48) has, however, only 20.5% sequence similarity (Fig. S6), indicating that unique interactions are responsible for Vn binding. The relatively higher affinity for Vn by PE compared with PF implies an important role of PE in Vn-dependent serum resistance of NTHi, whereas PF may serve as an alternative/additional candidate. This is probably a back-up strategy for NTHi to consistently acquire Vn, and this dual Vn-binding strategy may have an impact on bacterial pathogenesis. NTHi 3655 devoid of either PF or PE resulted in decreased Vn-binding, serum resistance, and finally increased C9 deposition (Fig. 9C-E) when compared with the wild type. However, despite mutation of both the hpf and pe genes resulted in a decreased Vn binding it was not statistically significant compared with the pe single mutant. Thus, in our hands PE most likely plays a major role for vitronectin-mediated NTHi serum resistance.

We found that PF is a highly conserved Vn-binding protein in NTHi clinical isolates (Fig. 10). Since PF is also surface exposed and immunogenic, it may therefore serve as a potential vaccine candidate. Although immunization with PE has previously been proven to be protective in a mouse pulmonary clearance model (Ronander *et al.*, 2009), simultaneous targeting of both PE and PF, in addition to other conserved OMP (e.g. Omp6 and PD), may result in a more effective vaccine strategy (Pichichero *et al.*, 2010).

PF has also been designated as adhesin B (Lu *et al.*, 1998), HfeA (Harrison *et al.*, 2005) or YfeA-like protein (Whitby *et al.*, 2006), and belongs to the TroA superfamily with homology to YfeA from the *yfeABCD* operon of *Y. pestis* that is involved in Fe^{2+} , Mn^{2+} and Zn^{2+} uptake (Bearden and Perry, 1999). However, the role of PF in iron uptake has not yet been fully characterized. Importantly, PF expression was reported to be upregulated

during iron/haem-restricted growth conditions, and in bacteria incubated with sputum derived from COPD patients (Whitby *et al.*, 2006; Qu *et al.*, 2010). *hpf* was also one of several NTHi genes shown to be essential for a delayed NTHi clearance in a pulmonary mouse model (Gawronski *et al.*, 2009). Moreover, PF orthologues are known virulence determinants in *Y. pestis* when causing bubonic plague (Bearden and Perry, 1999), in *Y. pseudotuberculosis* by mediating polymyxin B resistance and colonization (Arafah *et al.*, 2009), and also contribute to iron-uptake and consequently infection by *Shigella flexneri* (Runyen-Janecky *et al.*, 2003).

In summary, NTHi-dependent acquisition of Vn is multifactorial, and here we identified PF as a novel Vn-binding protein. The important role of PF in serum resistance and adhesion warrants further characterization for a full understanding of NTHi pathogenesis. Our findings may also be an impetus for development of a future vaccine against NTHi.

Experimental procedures

Bacterial strains

Clinical isolates (*n* = 18) (Table S1) and laboratory strains of NTHi 3655 (Table S2) were grown on chocolate agar or cultured in brain heart infusion (BHI) broth supplemented with NAD (Sigma-Aldrich, St Louis, MO) and hemin (Merck, Darmstadt, Germany) (each at 10 μ g ml⁻¹) at 37°C in a humid atmosphere containing 5% CO₂. *E. coli* BL21(DE3) (Novagen, Darmstadt, Germany) and DH5 α (Stratagene, Santa Clara, CA) were cultured in Luria–Bertani (LB) broth or on LB agar. Kanamycin (Km) and ampicillin (Amp) (Merck) were used at 50 μ g ml⁻¹ and 100 μ g ml⁻¹, respectively, when indicated.

Construction of an NTHi hpf deletion mutant and trans-complementation of hpf

To delete the hpf gene in NTHi, the upstream (621 bp) and downstream (997 bp) flanking regions were amplified from NTHi 3655 genomic DNA with primers listed in Supporting information (Table S3). Primers used for the cloning of the flanking regions contained overhangs that are complementary to the chloramphenicol acetyltransferase gene (cat) (EMBL BAA78807). The cat gene was amplified from the plasmid pLYS (Novagen). A linear knockout vector fragment (2279 bp) was synthesized by using the two flanking regions containing cat overhang together with the cat amplicon as a template in an overlapping PCR reaction. The PF promoter was kept intact to obviate a possible polar effect caused by disruption of the PF operon by the cat promoter. The vector also included the H. influenzae uptake signal sequence (AAGTGCGGT). A schematic map for the hpf knockout is shown in Fig. S10. Competent NTHi was prepared and transformations were performed as previously described (Poje and Redfield, 2003). NTHi 3655 and NTHi 3655∆pe were transformed with the vector resulting in the strains NTHi 3655∆hpf and NTHi 3655∆pe/hpf respectively. Transformants were selected on

chocolate agar plates containing chloramphenicol (Cm) (5 μ g ml⁻¹) for NTHi 3655 Δ *hpf* and combination of Cm and zeocine (Zeo) (15 μ g ml⁻¹) for NTHi 3655 Δ *pe/hpf*.

Trans-complementation of the hpf gene in NTHi 3655∆hpf was achieved by genomic homologous recombination following a strategy described previously (Jones et al., 2002; Ünal et al., 2012). The pKR2 plasmid which contains the intergenic region from the locus CGSHi3655_06559 of NTHi3655 was used as starting material. A Km resistance cassette (nptl) amplified by PCR using the Primers Km_F and Km_R (Table S3) was inserted into the vector pKR2 using the restriction sites Sphl and Sall leading to the vector pKR7.1. The hpf gene was amplified by PCR with primers PFprom F and PF293 R from genomic DNA and cloned into pKR7.1 linearized by EcoRV and Sall yielding the plasmid pKR7.2. The plasmids pKR7.1 and pKR7.2 contain either the promoter of hpf (pKR7.1) or hpf driven by its endogenous promoter (pKR7.2) followed by a Km resistance gene cloned in reverse direction (Fig. S10). Plasmids pKR7.1 and pKR7.2 were used as template for PCR amplification with the primers KS36 and KS37 to generate the constructs for transformation. The subproduct of pKR7.1 was used to transform NTHi3655 Appf in order to generate the control strain NTHi3655∆hpf (pKR7.1). The PCR product derived from pKR7.2 was used to transform NTHi3655 Δ hpf yielding the strain NTHi3655 Δ hpf (pKR7.2) carrying hpf complemented in trans. Insertion of the elements in the CGSHi3655_06559 locus was confirmed by PCR using the primers PFprom R and RecNTHi3655 F located upstream the insertion point of the construct transformed. Resulting NTHi 3655∆hpf (pKR7.1) and NTHi 3655∆hpf (pKR7.2) mutants were selected on a chocolate agar containing chloramphenicol (Cm) (Sigma) (5 µg ml-1) and Km (15 µg ml⁻¹). All mutants were characterized and confirmed by DNA sequencing, growth curve analysis, SDS-PAGE, western blotting, and flow cytometry.

Sequence analysis of hpf in clinical isolates

For assessment of the *hpf* gene in clinical isolates, the gene was amplified from bacterial genomic DNA with primers For_PF_flank and Rev_PF_flank (Table S3) covering the upstream (CGHi3655_02304) and downstream flanking region (intergenic region, located at nucleotides 1034–1186). Purified PCR products were subjected to DNA sequencing. ORFs of PF variants were deduced from the sequenced DNA. Multiple sequence alignment was performed using Lasergene[®] MegAlign (DNAstar, Madison, WI).

Recombinant protein expression, peptides and antibodies

Full-length and truncated ORF of PF (GI: 145632610) were amplified from the genomic DNA of NTHi 3655 with site specific primers listed in *Supporting information* Table S4. The expression vector pET26b (Novagen) was used for cytoplasmic recombinant protein production, whereas pET16b (Novagen) was used for bacterial cell surface expression of the target protein. Resultant recombinant constructs were thereafter transformed into the chemically competent host *E. coli* DH5 α . The authenticity of cloned DNA sequence was verified by DNA sequencing (MWG Eurofins, Ebersberg,

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Germany). Using the pET26b backbone constructs, recombinant proteins were produced in E. coli BL21(DE3) and harvested as previously described (Su et al., 2010). The polyhistidine-tagged recombinant proteins were purified from crude extracts using immobilized metal affinity chromatography-based Histrap[™] FF Crude (GE Healthcare Biosciences, Uppsala, Sweden) according to the manufacturer's instructions. Protein concentrations were estimated at absorbance 280 nm with Nanodrop (Thermo Scientific) and analysed by SDS-PAGE. For expression of surface-bound PF, the E. coli expression culture was induced overnight, and used in a serum killing assay and flow cytometry analyses. A rabbit anti-PF pAb was prepared as previously described (Hallström et al., 2009). To increase the specificity, the rabbit anti-PF antiserum was subsequently affinity purified as previously described with PF12-293 coupled to CNBr-activated Sepharose[™] (GE Healthcare Biosciences).

A series of peptides (23–25 aa long) encompassing the entire PF protein sequence (excluding signal peptide aa 1–22) were also synthesized (Innovagen, Lund, Sweden).

Two-dimensional gel electrophoresis

Outer membrane vesicles of NTHi 3655 are enriched with periplasmic- and outer membrane-associated proteins (Schaar et al., 2011; Sharpe et al., 2011). In the following experiments, we thus used the OMV extract as the protein source of bacterial OMP in order to reduce contamination from the cytoplasmic compartment. The OMV of NTHi 3655 were prepared from overnight cultures and separated by 2D-gel electrophoresis as previously described with some modifications (Schaar et al., 2011). For the first dimension separation by isoelectric focusing (IEF), 125 µl of solubilized sample (100 μ g) was removed and applied to a precast 7 cm pH 3-10 IPG gel strip (Immobiline Drystrips, GE Healthcare Biosciences) by in-gel sample rehydration method. Using IPGphor II electrophoresis unit (GE Healthcare Biosciences), IEF was performed with the following programme: 30 V for 0.30 hr (15 Vhr), 500 V for 0.30 hr (250 Vhr), 1000 V for 0.30 hr (500 Vhr), 2600 V for 0.15 hr (650 Vhr), 3500 V (875 Vhr) and 5000 V for 1.00 hr (5000 Vhr) yielding a total voltage of 7291 Vhr, with maximum current of 50 µA per strip. Thereafter, the strips were equilibrated as described elsewhere (Schaar et al., 2011) and run in a second dimension of gel electrophoresis on a 12% (v/v) SDS-polyacrylamide gel at 70 V for 15 min and 150 V for 40 min.

Protein spots on gels were visualized by silver staining (Bio-Rad) for image analysis and alternatively with Coomassie brilliant blue staining for protein sequence identification with mass spectrometry. Gels were scanned on a Chemi-Doo[™] XRS+ System (Bio-Rad) and analysed with the Image Lab[™] Software (Bio-Rad). Targeted Coomassie stained protein spots were manually excised and protein identification by Matrix-assisted laser desorption/ionisation-time of flight mass spectrometry (MALDI-TOF-MS) was performed by Protein Analysis Service available at Alphalyse (Odense, Denmark).

SDS-polyacrylamide gel electrophoresis

One-dimensional (1D) SDS-polyacrylamide gel electrophoresis (SDS-PAGE) was performed according to the Laemmli

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method (Laemmli, 1970). Briefly, OMV (10 µg), purified protein (10 µg) or whole cell lysates $[3 \times 10^7 \text{ cfu} (colony-forming$ units)] in PBS were heat-denatured at 95°C for 10 min inSDS-reducing sample buffer (50 mM Tris-Cl pH 6.8, 2% SDS,6% Glycerol, 1% β-mercaptoethanol, 0.004% bromophenolblue). Thereafter, samples were resolved in 12% (v/v) SDSpolyacrylamide gel at 150 V for 60 min. Protein bands on gelswere visualized by Coomassie blue staining and scanned asdescribed above. For detection of Vn binding or PF expression, proteins on gels were subjected to western blotting.

Western blotting

Proteins separated in SDS-PAGE were transferred to a 0.45 µm Immobilon-P[™] polyvinylidene fluoride (PVDF) membrane (Millipore) at 15 V for 16 hr. Membranes were blocked with blocking buffer [5% (w/v) skimmed milk in PBS] and washed with PBS containing 0.05% (v/v) Tween 20 (PBST). For detection of Vn-binding proteins, membranes were probed with human Vn (Sigma) (3 μ g ml⁻¹) in blocking buffer for 1 hr at room temperature. Bound Vn was detected with a sheep anti-human Vn pAb (1:2000 dilution) (Abd Serotec, Oxford, UK) and horseradish peroxidase (HRP)-conjugated donkey anti-sheep pAb (1:4000 dilution) (AbD Serotec). For detection of PF, blots were incubated with rabbit anti-PF pAb (1:1500 dilution) followed by HRP-conjugated swine antirabbit pAb (1:1000 dilution) (Dako). Reactive antibodies were detected with enhanced chemiluminescence (ECL) western blotting substrate (Pierce, Rockford, IL). Chemiluminescence signals on membranes were visualized on a ChemiDoc™ XRS+ System.

Bacterial binding to Vn-coated glass slides

Glass slides (Menzel-Gläser, Thermo Scientific, Wilmington, DE) were incubated with 6 μ g Vn (Sigma) for 1 hr at 37°C for immobilization. Bacteria were grown in broth until mid-log phase (OD₆₀₀ = 0.5) and incubated on the Vn-coated glass slide for 1 hr at room temperature. Unbound bacteria were fixed by flaming and visualized by Gram-staining. Slides were prepared according to a standard protocol and analysed in an Olympus BX51 microscope at 400× amplification.

Transmission electron microscopy

Negative staining and TEM were used to visualize the localization of PF at the bacterial surface. NTHi 3655 and NTHi 3655 Δhpf were incubated with a colloidal thiocyanate goldlabelled anti-PF pAb and visualized by a Jeol JEM 1230 electron microscope (JEOL, Tokyo, Japan) operated at 60 kV accelerating voltage was used. Images were recorded with a Gatan Multiscan 791 CCD camera (Gatan, Pleasanton, CA).

Serum bactericidal assay

NHS and Vn-depleted serum was prepared as previously described (Hallström *et al.*, 2009). The NHS used in the present study was reactive against NTHi as revealed by western blotting (data not shown). Susceptibility of NTHi and *E. coli* BL21(DE3) to complement-mediated killing was tested

as previously described (Yi and Murphy, 1997). The optimal concentration of serum used was as previously suggested (Hallström et al., 2009; 2010) and had been empirically determined for NTHi (5% NHS) and E. coli BL21(DE3) (1% NHS). Aliquots of 1.5×10^3 cfu in 150 µl of DVBS-BSA [2.5 mM Veronal buffer, pH 7.3, 1 mM MgCl₂, 2.5 mM CaCl₂, 2.5% (w/v) glucose and 10% (w/v) bovine serum albumin] were added with appropriate dilutions of serum and incubated at 37°C at the indicated times with gentle shaking. At 0 min (T₀ sample) and indicated times (T_t sample) aliquots of 10 μ l of the reaction mixtures were plated on chocolate agar for NTHi or LB agar containing Amp for PF-expressing E. coli and incubated at 37°C. Percentage of bacterial killing was expressed as $(T_0 cfu - T_t cfu)/T_0 cfu) \times 100$. Three individual experiments, each performed in triplicates, were done. Control experiments were also performed as described above using heatinactivated serum (HIS) (pretreated at 56°C for 30 min).

Expression of Vn fragments in HEK293 cell lines and protein purification

Vn fragments (spanning aa 80–396, 80–320, 80–229, 80–379, 80–373, 80–363, 80–353, 80–339, and 80–330) were expressed in HEK293T cells and purified as previously described (Singh *et al.*, 2011). In addition, Vn^{80–396} with internal deletions of amino acids 352–362, 362–374, and 352–374, were produced in the present study. The protein expression, purification and CD spectra analysis were performed as described elsewhere (Singh *et al.*, 2011).

Direct protein binding assay

Human Vn (Sigma) was labelled with 0.05 M iodine (Amersham Biosciences, Buckinghamshire, UK) per mol protein using the chloramine-T method (Nordstrom *et al.*, 2004). Bacterial colonies were resuspended in PBS to $OD_{600} = 0.5$, washed and resuspended in PBS-2% BSA. [I¹²⁵]-labelled-Vn (80 kcpm/sample) was added to bacterial suspensions containing 1×10^7 cfu and incubated for 1 hr at 37°C. After two washing steps with PBS-2% BSA, iodine-labelled protein bound to bacteria was measured in a gamma counter (Wallac, Espoo, Finland). *E. coli* Top10 was included as a negative control. All direct binding assays were performed in triplicates at three independent occasions.

Flow cytometry analyses

PF expression at the bacterial surface was detected using flow cytometry. NTHi from mid-log phase cultures or PF-expressing *E. coli* BL21(DE3) transformants induced overnight were resuspended in PBS containing 1% BSA (blocking buffer), and adjusted to $1 \times 10^{\circ}$ cfu ml⁻¹. Aliquots containing 5×10^{7} cfu were incubated with anti-PF pAb for 1 hr, washed and pelleted by centrifugation. Thereafter, bacterial suspensions were incubated with FITC-conjugated swine anti-rabbit pAb (Dako) for 30 min.

For ligand binding at the bacterial surface, suspensions were incubated with increasing concentrations of Vn [1–400 nM for NTHi or $5-1 \times 10^3$ nM for *E. coli* BL21(DE3) expressing PF] for 1 hr at 37°C. After washing, bound ligands were detected using sheep anti-human Vn pAb (dilution 1:50),

followed by FITC-conjugated donkey anti-sheep pAb (dilution 1:150) (Abd Serotec). Serum C4BP and FH were detected with mouse anti-C4BP and anti-FH monoclonal antibodies (mAb) followed by FITC-rabbit anti-mouse pAb (Abd Serotec). For detection of C9 deposition at the bacterial surface, bacterial suspensions were incubated with 10% NHS as described in the C9 deposition assay as described below. Thereafter, cells were washed and surface-deposited C9 were detected using goat anti-C9 pAb (CompTech) (dilution 1:100) followed by FITC-conjugated donkey anti-goat pAb (Abd Serotec). Unspecific binding by secondary antibodies was assessed by omitting the primary antibodies, ligands or NHS. Bacteria were analysed in a flow cytometer (EPICS XL-MCL flow cytometer; Coulter, Hialeah, FL). Ten thousand events were examined for each sample. The positive population was gated at 5% of the background binding population in the control.

Enzyme-linked immunosorbent assay (ELISA)

The interaction of Vn with PF was analysed by ELISA. This assay was done by using PF recombinant proteins or synthetic peptides as indicated (Fig. 5A and E). Either the indicated PF synthetic peptides or recombinant PF proteins diluted at 50 nM in coating buffer (100 mM NaH₂CO₃, pH 8.6) were immobilized overnight on Polysorb microtitre plates. After washing, Vn fragments as outlined in Fig. 6A were added to each well at the indicated concentrations followed by incubation for 1 hr at room temperature. Thereafter, bound Vn was detected with sheep anti-human Vn pAb followed by HRP-donkey anti-sheep pAb. Signals of antigen–antibody complexes were developed with 20 mM tetramethylbenzidine and 0.1 M potassium citrate. Finally, the reactions were stopped with 1 M H₂SO₄. Signals developed were read at an absorbance of 450 nm. All experiments were performed in triplicates.

Haemolytic assay

The MAC inhibitory activity of PF-bound Vn was analysed in a haemolytic assay with sheep erythrocytes as previously described with some modifications (Hallström et al., 2009). Briefly, sheep erythrocytes resuspended to 2×10^8 cells ml⁻¹ in DVBS containing 0.1% gelatin (DGVB). Aliquots of 100 µl sheep erythrocytes were pre-incubated with 2 µg ml-1 C5b-6 (CompTech, Tyler, TX) for 30 min at 37°C. In a separate preparation, 0.5 µM Vn (Sigma) was pre-incubated with increasing concentrations of purified recombinant PF12-293 $(0.25-1.0 \ \mu\text{M})$ for 15 min at 37°C in a final volume of 100 μ l. Thereafter, the mixture was added with 2 µg ml-1 C7 (CompTech), 0.2 µg ml⁻¹ C8 (CompTech) and 2 µg ml⁻¹ C9 (CompTech), and further incubated for 30 min. After pre-incubation, the C5b-6 coated erythrocytes were added to the Vn-C7-9 mixture and incubated for 30 min at 37°C. Non-haemolysed erythrocytes were pelleted and the amount of haemoglobin representing the lysed erythrocytes was measured at 541 nm. The relative MAC inhibitory activity was presented in the percentage of total haemolysis.

C9 deposition assay

The effect of PF on MAC formation at the bacterial surface was assayed on the basis of C9 deposition on bacterial cells

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as previously described with some modifications (Singh *et al.*, 2011). Briefly, bacteria resuspended to $OD_{eo0} = 0.1$ in DGVB were pre-incubated with various concentrations of Vn. This was followed by incubation with 10% NHS in a final volume of 100 µl at 37°C at the indicated time points and mixtures were incubated on ice to stop the reactions. Thereafter, 10 µl aliquots were mixed with LDS sample buffer (Invitrogen, Carlsbad, CA) and boiled at 70°C for 10 min. Total bacterial cell proteins were separated on a 4–12% Bis-Tris gradient gel (Invitrogen) as described by the manufacturer. This was followed by western blotting with goat anti-C9 pAb and HRP-conjugated donkey anti-goat pAb (Abd Serotec). In parallel, C9 deposition at the bacterial surface was quantified by flow cytometry as described above.

Statistics

One- or two-way ANOVA tests were used to compare the difference between more than two experimental groups when indicated. Differences were considered statistically significant at $P \le 0.05$. Statistical analyses were performed using Graph-Pad Prism[®] version 5.0 (GraphPad Software, La Jolla, CA).

Ethics statement

Normal human sera were obtained from healthy adult volunteers at Skane University Hospital, Malmo, Sweden, and prepared according to the general rules established by the Swedish Government (Svensk författningssamling 1988:534, Ändringsförfattning 2012:256; Riksdagen, Riksgatan 1, SE-111 28, Stockholm, Sweden). Ethical permit (M193-11) was obtained from Malmö/Lund District Court (Djurförsöksetiska nämnden, Tingsrätten, Byggmästaregatan 2, SE-222 37 Lund, Sweden). This body approved the protocol used in this study.

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Conflict of interest

Conflict of interest between authors are not reported.

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Supporting information

Additional supporting information may be found in the online version of this article at the publisher's web-site.

Haemophilus influenzae Protein F Mediates Binding to Laminin and Human Pulmonary Epithelial Cells

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The mucosal pathogen nontypeable Haemophilus influenzae (NTHi) adheres to the respiratory epithelium or, in the case of epithelial damage, to the underlying basement membrane and extracellular matrix that, among other proteins, consists of laminin. We have recently identified protein F, an ABC transporter involved in NTHi immune evasion. Homology modeling of the protein F tertiary structure revealed a strong resemblance to the streptococcal laminin-binding proteins Lbp and Lmb. Here, we show that protein F promotes binding of NTHi to laminin and primary bronchial epithelial cells. Analyses with recombinant proteins and synthetic peptides revealed that the N-terminal part of protein F contains the host-interacting region. Moreover, protein F exists in all clinical isolates, and isogenic NTHi Δhpf mutants display significantly reduced binding to laminin and epithelial cells. We thus suggest protein F to be an important and ubiquitous NTHi adhesin.

Keywords. ABC transporter; adhesion; laminin; nontypeable *Haemophilus influenzae*; protein F; pulmonary epithelial cells; respiratory tract infection; virulence.

Nontypeable Haemophilus influenzae (NTHi) is a gram-negative opportunistic pathogen that colonizes the nasopharynx of humans. It is one of the leading causes of bacterial respiratory tract infections, such as acute otitis media in children and bronchitis, as well as exacerbations in patients with chronic obstructive pulmonary disease (COPD) [1–3]. To prevent clearance by the ciliated respiratory tract mucosal epithelium an important initial step of bacterial colonization is adherence to host tissue [4]. In patients with disruption of the epithelial integrity caused by viral infections, mechanical damage, or chronic inflammation, the underlying basement membrane is exposed

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to the lumen and forms a viable attachment site for pathogens [5].

The heterotrimeric glycoprotein laminin (Ln; approximately 800 kDa) is one of the major constituents of the extracellular matrix (ECM) and the basement membrane. It is involved in an array of physiological functions, such as cellular proliferation, migration, and structural scaffolding in tissues [6–8]. During bacterial pathogenesis, Ln is frequently targeted for adherence by respiratory tract pathogens [9–16]. NTHi has previously been reported to bind Ln via the *Haemophilus* adhesion and penetration protein (Hap) and protein E (PE) [17, 18]. In parallel, Hap and PE interact with host epithelial cells [19, 20]. These interactions are thus multifactorial and involve several adhesins that mutually promote bacterial host-adhesion.

We recently identified a virulence factor in NTHi that we designated as protein F (PF; Su et al, unpublished data). PF (approximately 30 kDa) is annotated as a bacterial metal-binding receptor, belonging to the adenosine triphosphate-binding cassette (ABC) transporter family. During structural analysis of PF, we noticed a strong structural resemblance to the

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Ln-binding proteins of *Streptococcus pyogenes* (Lbp) and *Streptococcus agalactiae* (Lmb) [21, 22]. The structural similarity raised a hypothesis regarding the potential Ln-binding capacity of PF. In the present study, we show that PF is a novel surface-exposed Ln-binding protein that also promotes NTHi adherence to host epithelial cells. This is the first report that shows a direct interaction between a gram-negative ABC transporter protein and host components.

METHODS

Bacteria, Eukaryotic Cells, Culture Conditions, and Reagents

NTHi 3655 wild-type (wt), Δhap , and Δpe mutants, and clinical isolates KR217, KR314, KR315, KR336, KR385 obtained by nasopharyngeal swabbing from patients with upper respiratory tract infection, were cultured as described elsewhere [17]. Chloramphenicol (10 µg/mL) was used for selection of Δhpf mutants. Escherichia coli BL21 (DE3; Novagen) and DH5a (Invitrogen) were cultured in Luria-Bertani broth and solidphase medium supplemented with kanamycin (50 µg/mL) or ampicillin (100 µg/mL). The type II alveolar A549 (American Type Culture Collection [ATCC] CCL-185) and bronchial epithelial NCI H292 (ATCC CRL-1848) cell lines were cultured in F-12 and Roswell Park Memorial Institute medium (Gibco), respectively, with 10% fetal calf serum. Primary bronchial epithelial cells were obtained from a healthy adult donor with no history of lung disease. This protocol was approved by the Swedish Research Ethical Committee in Lund (FEK 413/ 2008), and written consent was obtained from the closest relatives. Cells were maintained in BEGM medium (Clonetics). All bacteria and human cells were cultured at 37°C with 5% CO2. Anti-PF polyclonal antibodies (pAb) were raised in rabbits and affinity purified using recombinant PF (rPF) or PF peptides [23].

Structural Modeling and Bioinformatic Analyses

Modeling of PF was performed using the Swiss-Model automated server against homologous templates available in the Protein Data Bank (PDB; available at: http://www.rcsb.org). Homologs were analyzed with BLAST (available at: http:// www.ncbi.nlm.nih.gov/), and multiple alignment was performed using ClustalW (available at: http://www.ebi.ac.uk/). Three-dimensional models were prepared using PyMOL (available at: http://www.pymol.org/).

Construction of NTHi Δhpf Mutants

Upstream (621 bp) and downstream (997 bp) flanking regions of the *hpf* gene (CGSHi3655_02309) from NTHi 3655 genomic DNA (GenBank accession number AAZF00000000) and *cat* (chloramphenicol acetyltransferase; BAA78807) from pLysS (Novagen) were amplified (Supplementary Table 1). Thereafter, we produced a linear knockout vector, inserting *cat* between the flanking regions, using an overlap extension polymerase chain reaction (Supplementary Figure 1). The *hpf* gene in NTHi 3655, clinical isolates, and the isogenic NTHi $3655 \Delta pe$ was mutated as described elsewhere [24].

Recombinant Proteins and Peptides

We amplified full-length and truncated fragments of NTHi 3655 *hpf*, using specific primers (Supplementary Table 1). Restriction sites were introduced as indicated. pET26(b)+ (for purified recombinant proteins) or pET16b (Novagen; for surface expression of PF) were used and transformed into *E. coli* DH5 α , followed by transformation into *E. coli* BL21 (DE3). Recombinant proteins were produced as described previously [25]. Peptides (approximately 25 amino acids long) spanning the entire mature PF and overlapping with neighboring peptides were from Innovagen. The radiolabeled peptidebinding assay was performed as described elsewhere [26].

Flow Cytometry and Transmission Electron Microscopy

For flow cytometry, stationary phase bacteria (10⁹ colony forming units/mL) were washed and resuspended in phosphate buffered saline (PBS) containing 1% bovine serum albumin (BSA) followed by addition of Ln (Engelbreth-Holm-Swarm murine sarcoma basement membrane, Sigma-Aldrich). The samples were incubated at 37°C for 1.5 hours. Bacteria were thereafter washed in PBS plus 1% BSA and were incubated with rabbit anti-Ln pAb (Sigma-Aldrich). After washing, fluorescein isothiocyanate–conjugated swine anti-rabbit pAb (Dako) were added. Finally, bacteria were analyzed by flow cytometry (EPICS XL-MCL, Beckman Coulter). Ten thousand events were measured for each sample. Transmission electron microscopy (TEM) was performed as described previously [17].

Bacterial Adherence Assays

Adherence of NTHi to immobilized Ln was studied by bacterial probing over Ln-coated glass slides [17]. NTHi adherence to mammalian cell lines and primary cells was analyzed using [³H]-thymidine-pulsed bacteria as described elsewhere [19]. Blocking was conducted with 20 µg of antibodies per well.

Enzyme-Linked Immunosorbent Assay (ELISA)

Laminin binding to rPF fragments was studied with an indirect ELISA as described previously [17]. Recombinant truncated PF fragments (50 nM) and synthetic PF peptides (1 μ M) were immobilized on polysorp microtiter plates (Nunc). Blocking was done with PBS containing 5% milk for 1 hour at room temperature. For the binding-inhibition assay, Ln (10 nM) was preincubated with PF peptides (0–5 μ M) at 37°C for 1.5 hours. The suspension was thereafter transferred to plates coated with rPF¹²⁻²⁹³, and the binding was measured as described above. In the direct cell ELISA, cells were grown to confluency in 96-well plates (Nunc) and fixed with 2% formaldehyde for 30 minutes at room temperature. Thereafter, cells were incubated with PF and, subsequently, rabbit anti-PF pAb. Horseradish peroxidase-conjugated anti-rabbit pAb (Dako) was used as the secondary layer for all ELISA.

Statistical Analyses

We used the Student *t* test and Mann–Whitney *U* test for statistical analyses of 2 parametrical sets of data and 2 nonparametric sets of data, respectively. Two-way analysis of variance was used for statistical analyses of several sets of data. Statistical analyses were performed using GraphPad Prism 5 (GraphPad Software). A *P* value of $\leq .05$ was considered statistically significant.

RESULTS

PF Is Surface Associated and Has a Tertiary Structure Similar to That of Streptococcal Laminin-Binding Proteins

We have recently identified PF, an NTHi protein involved in serum resistance (Su et al, unpublished data). The hpf gene encodes an uncharacterized 30-kDa iron-chelating protein and is one of 4 structural genes in an ABC transporter operon. The operon is present in all 20 available NTHi genomes in GenBank and is highly conserved (>98% identity; data not shown). To experimentally examine the subcellular localization of PF and determine whether it is exposed on the bacterial surface, we constructed an isogenic NTHi 3655 Ahpf mutant (Supplementary Figure 1A). The physiological fitness of the mutant did not differ from the parental strain with regard to the whole cell protein profile, but a slightly slower growth was observed in soluble medium (data not shown). Transcription analysis showed that the other genes in the operon remained actively expressed in the isogenic mutant (data not shown). By use of TEM, gold-labeled anti-PF pAb recognized PF at the surface of NTHi 3655, but it did not recognize the PF-deficient mutant (Figure 1A). Since the affinity-purified anti-PF pAb were highly specific for PF (Supplementary Figure 1B), this experiment suggested that PF is a surface-exposed protein.

To elucidate the characteristics of the protein, we analyzed the tertiary structure of PF by using homology modeling. The manganese-binding protein MntC of cyanobacteria Synechocystis species (PDB code 1XVL) was found to be the closest structure-solved homolog with a 343-bit alignment score (52.1% identity/67.0% similarity) and was therefore used as the modeling template. The superimposition of PF with MntC is shown in Figure 1B. The structural model revealed that PF is constituted by distinct N- and C-terminal globular domains that are interlinked by a long helix backbone. A metal-binding active site is present between both lobes. Since the Ln-binding proteins Lbp, of S. pyogenes (PDB code 3GI1), and Lmb, of S. agalactiae (PDB code 3HJT), belong to the same bacterial ABC transporter family, we superimposed PF with their crystal structures. Interestingly, despite the weak primary sequence similarity (Lbp, 28.3% identity/46.9% similarity; Lmb,

28.0% identity/46.3% similarity), we observed a strong resemblance in the secondary structure and folding of PF and Lbp/ Lmb (Figure 1*C*) [21].

PF Is a Ubiquitous Laminin-Binding Protein of H. influenzae

The structural resemblance between PF and Lbp/Lmb prompted us to investigate the potential Ln-binding capacity of PF. To study the PF-Ln interaction at the bacterial surface, we incubated NTHi 3655 wild-type and the PF-deficient NTHi 3655 Δhpf with increasing concentrations of soluble Ln. The unbound Ln fraction was thereafter washed away, and bacterial Ln binding was analyzed by flow cytometry. Our data showed that the isogenic Δhpf mutant bound significantly less Ln as compared to the NTHi 3655 wild-type ($P \leq .05$; Figure 2A and 2C).

To prove the Ln-binding property of PF at the bacterial surface, we introduced the *hpf* open reading frame into a heterologous *E. coli* host. After induction of *hpf* expression, we observed a Ln-binding phenotype in the PF-producing *E. coli*, compared with the control *E. coli* containing the empty plasmid (Figure 2*B* and 2*C*). Further analysis of the Ln interaction at the bacterial surface was conducted using TEM. The isogenic NTHi 3655 Δhpf showed a marked reduction in Ln binding as compared to the parental strain (Figure 2*D*). Moreover, co-localization of gold-labeled Ln (10 nm) and PF (detected by gold-labeled anti-PF pAb; 5 nm) was observed at the surface of both PF-producing NTHi 3655 and *E. coli* (Figure 2*E*).

To study the functional bacterial binding to immobilized Ln, which is more representative of the in vivo basement membrane, Ln was coated on glass slides and incubated with various samples of NTHi and *E. coli* adjusted to the same cell density (optical density at 600 nm = 1.0). NTHi 3655 Δhpf displayed a markedly weaker adherence to immobilized Ln as compared to the NTHi 3655 wild-type (Figure 2*F*). In parallel, PF-producing *E. coli* adhered to the Ln-coated glass slides, whereas the control *E. coli* did not. Taken together, the co-localization of PF and Ln at the surface of NTHi 3655, in addition to the PFdependent Ln binding of heterologous host *E. coli*, suggest that PF is directly involved in Ln binding at the bacterial surface.

We further wanted to evaluate the Ln-interacting role of PF in a series of clinical NTHi isolates from patients with upper respiratory tract infection. PF-deficient mutants were produced in 5 clinical isolates, and Ln binding was assessed by flow cytometry (Figure 3A). A consistent decrease in Ln binding was observed in all strains in the absence of PF. The results indicate that PF is important for the NTHi-Ln interaction in several clinical isolates.

PF Contributes to the Multifactorial NTHi-Dependent Laminin Binding

Currently, there have been 2 reports regarding Ln-binding proteins in NTHi [17, 18]. To evaluate the relevance of PF in



Figure 1. Subcellular localization of protein F (PF) and the structural model of the protein. A, Gold-labeled anti-PF polyclonal antibodies recognize PF at the nontypeable Haemophilus influenzae (NTHi) 3655 surface but not at the surface of the isogenic PF-deficient NTHi 3655 $\Delta h \rho f$ mutant, as revealed by transmission electron microscopy. B, Superimposed model of PF (grey) with the template MntC of Synechocystis species (Protein Data Bank code 1XVL) shown in green (52.1% identity and 67.0% similarity). C, Superimposition of PF (grey) with Streptococcus pyogenes laminin-binding protein (blue) and Streptococcus agalactiae laminin-binding protein (yellow). N- and C-terminal domains are indicated, with the metal ion-chelating active site residues shown in sticks, situated between both domains. Figures were prepared using PyMOL.

comparison to PE and Hap for the NTHi-Ln interaction, NTHi 3655 wild-type and knockout mutants were incubated with soluble Ln, and binding was measured with flow cytometry (Figure 3*B*). Our data demonstrated that all mutants had decreased Ln binding as compared to their wild-type counterparts, showing that PE, Hap, and PF contribute to the Ln interaction.

The N-terminal Region Lys23-Glu48 of PF Interacts With the C-terminus of the Laminin $\alpha\text{-Chain}$

To study the PF-Ln interaction at the molecular level, a series of recombinant PF (rPF) fragments were produced (Figure 4Aand Supplementary Table 2). The putative Ln-binding region of PF was determined by incubating immobilized truncated rPF fragments with increasing concentrations of Ln. Recombinant PF¹²⁻²⁹³ and rPF¹²⁻⁹⁸ bound significantly better when compared to the nonbinding fragment rPF⁶¹⁻¹⁴⁴ ($P \le .001$; Figure 4*B*). The N-terminal rPF¹²⁻⁹⁸ also exhibited saturable and dose-dependent Ln binding, whereas no binding to the negative control fragment rPF¹²⁹⁻¹⁷⁷ was detected (Figure 4*C*).

The Ln-binding region of PF was further analyzed in detail by using 13 synthetic peptides that span the entire PF molecule (Figure 5A). PF²³⁻⁴⁸ was identified as the major Lnbinding region ($P \le .001$) in an indirect ELISA. Moreover, PF²³⁻⁴⁸ exhibited saturable and dose-dependent binding to Ln, whereas the non-Ln-binding peptide PF¹²⁴⁻¹⁴⁸ did not (Figure 5B). The specificity of the interaction with Ln was evaluated with a binding inhibition assay in which Ln was preincubated with various PF peptides prior to addition to



Figure 2. Protein F (PF) mediates laminin (Ln) binding at the surface of bacteria. *A* and *B*, Flow cytometry analysis shows significant differences in the Ln binding of nontypeable *Haemophilus influenzae* (NTHi) 3655 wild-type and NTHi 3655 Δhpf (*A*) and PF-producing *Escherichia coli* (*B*) acquire previously nonexistent Ln binding, as measured by mean fluorescence intensity/bacterium. *E. coli* containing an empty plasmid was used as a control. The mean of 3 separate experiments is plotted, and error bars indicate the standard error of mean (*A* and *B*). Statistical analysis was performed using 2-way analysis of variance, and $P \leq .05$ was considered statistically significant. $**P \leq .01$: $***P \leq .001$. *C*, Histogram of a data set from *A* and *B*. *D*, Transmission electron microscopy shows a marked decrease of bound Ln (gold-labeled) at the surface of NTHi 3655 Δhpf as compared to the parental wild-type strain. *E*, Arrows point to bound gold-labeled Ln (10 nm) that is colocalized with PF detected by gold-labeled anti-PF polyclonal antibodies (5 nm) at the bacterial surface. The bar indicates 100 nm. *F*, PF promotes NTHi 3655 and *E*. *coli*-PF adherence to immobilized Ln in a basement membrane-mimicking setting, whereas non-PF-producing bacteria are seen to exhibit markedly less binding to the Ln-coated surface.

microtiter plates coated with rPF¹²⁻²⁹³ (Figure 5C). The peptide PF^{23-48} inhibited Ln binding to rPF^{12-293} , proving that the PF^{23-48} -Ln interaction was specific. Although $PF^{184-209}$ bound Ln (Figure 5A and 5*B*), the peptide did not inhibit Ln binding to immobilized rPF^{12-293} (Figure 5C). The N-terminal peptide PF^{23-48} was thus identified as the main Ln-binding region.

To determine the PF-binding domain of the Ln molecule, TEM with Ln and gold-labeled rPF^{12-293} was conducted. As

seen in Figure 5*D*, PF consistently bound to the C-terminal part of the cruciform Ln molecule. When the length of the molecule was measured, PF appeared to bind to the C-terminal globular domains (LG1–5) of the Ln α -chain (Figure 5*D* and 5*E*). We conclude that the N-terminal region of PF interacts with the C-terminus of the Ln α -chain, as revealed by the peptide mapping approach and TEM.



Figure 3. Protein F (PF) mediates laminin (Ln) binding in clinical isolates and contributes to the nontypeable Haemophilus influenzae (NTHi)– dependent Ln binding by an interaction analogous to that of PE and Hap. A, Flow cytometry analysis show that 5 clinical isolates display markedly decreased Ln-binding capacity when PF is knocked out, compared with the PF-producing parental wild-type (wt) strains. B, Comparison of PF-, PE-, and Hap-deficient mutants shows that the absence of all 3 proteins causes a decline in Ln binding (Ln concentration, 2.5 nM). The mean of 3 separate experiments is plotted in both graphs, error bars indicate the standard error of mean, and statistical differences are between wild-type and mutants. Statistical analysis was performed using analysis of variance, and $P \leq .05$ was considered statistically significant. * $P \leq .05$, ** $P \leq .01$.



Figure 4. Recombinant protein F (rPF) binds laminin (Ln) via a region in PF¹²⁻⁹⁸. *A*, Results of Coomassie-stained sodium dodecyl sulfate polyacrylamide gel electrophoresis, showing the affinity purified recombinant truncated PF-fragments. *B* and *C*, rPF¹²⁻⁹⁸ contains the Ln-binding region and binds to Ln in a dose-dependent and saturable manner. Statistical analysis was performed using the Mann–Whitney *U* test. The mean of 3 separate experiments is plotted in *B*. In *C*, 1 set of data from 3 similar experiments is shown. Error bars indicate the standard error of mean in all experiments.



Figure 5. Protein F (PF)²³⁻⁴⁸ interacts with the C-terminus of the cruciform laminin (Ln) macromolecule α -chain. *A*, To more precisely define the Ln-binding region of PF, a series of synthetic peptides spanning the entire PF molecule were produced. Two potential Ln-binding regions, PF²³⁻⁴⁸ (consistent with the data from the recombinant fragments) and PF¹⁸⁴⁻²⁰⁹, were identified using an indirect enzyme-linked immunosorbent assay. *B*, Ln binds to PF²³⁻⁴⁸ and, to a lesser degree, PF¹⁸⁴⁻²⁰⁹ in a dose-dependent and saturable manner. *C*, Only PF²³⁻⁴⁸ binds specifically to Ln and inhibits the interaction with recombinant PF (rPF¹²⁻²³³). *D*, Electron microscopy showing gold-labeled rPF¹²⁻²³³ binding to the Ln α -chain C-terminus (as indicated by white arrows). *E*, Schematic representation of the Ln hetereotrimer, showing the α -, β -, and γ -chains that form the cruciform molecule. The PF-binding α -chain C-terminal globular domains are indicated by an arrow. Statistical analysis was performed using the Mann–Whitney *U* test, and error bars indicate the standard error of the mean for all experiments. The mean of 3 separate experiments are shown in *A* and *C*, whereas 3 separate experiments were performed with the same outcome for *B*, of which 1 set of data is shown. *P* ≤ .05 was considered statistically significant. ****P* ≤ .001.

PF Mediates Adhesion of NTHi to Pulmonary Epithelial Cells via a Conserved N-Terminal Domain

Since several bacterial Ln-binding proteins, including the aforementioned PE, Hap, Lmb, and Lbp, have been reported to interact with epithelial cells [15, 19, 20, 27, 28], we analyzed the cell-adhesive capacity of PF. Epithelial cells were immobilized on a solid surface and incubated with increasing concentrations of rPF¹²⁻²⁹³ (Figure 6A). Recombinant PF bound to both NCI H292 and A549 cell lines in a dose-dependent and saturable manner. To evaluate the influence of PF on bacterial adhesion, NTHi 3655 wild-type and the PF-deficient Δhpf mutant were pulsed with [³H]-thymidine and incubated with pulmonary epithelial cells, and adhesion was measured after washing (Figure 6B). The adherence of NTHi 3655 Δhpf to NCI H292 and A549 was reduced by 32.9% and 64%, respectively, compared with the PF-producing wild-type. Importantly, the isogenic mutant also showed a significant reduction (36.6%) in binding to primary bronchial epithelial cells obtained from a healthy adult donor ($P \le .05$; Figure 6B).

To investigate the putative cell-adhesive region of PF, cell lines were probed with iodine-labeled synthetic PF peptides, followed by washes and measurement of binding (Figure 6C). Interestingly, the Ln-binding region PF²³⁻⁴⁸ (Figure 5A) also displayed the strongest binding to epithelial cells. Moreover, the bacterial-host cell interaction was significantly reduced ($P \le .05$) when bacteria were preincubated with anti-PF²³⁻⁴⁸ pAb but not with the control anti-PF⁴⁴⁻⁶⁸ (Figure 6D). These results thus suggest that NTHi uses the same N-terminal PF region for both binding to Ln and attachment to epithelial cells.

Bioinformatic analyses and a detailed database search using BLAST showed that PF orthologs exist in a number of other pathogens. The partial alignment of PF^{23-48} with these is shown in Figure 7*A*. The orthologs present in several members of the family Pasteurellaceae and in other pathogens, such as *Eikenella corrodens* and *Yersinia pestis*, have highly similar N-termini (Figure 7*A*). In addition, *S. pyogenes* and *S. agalactiae* Lbp/Lmb N-terminal sequences display partial similarity to PF^{23-48} (Figure 7*B*). The 3D model of the celland Ln-binding region was analyzed in detail and revealed to comprise 2 β -sheets and 1 α -helix, connected by 3 loops (Figure 7*C*). Amino acid residues Lys23 and Lys25 in loop-1, Gln34, Asp35, Gln38, Asn39, and Asn43 in the α -helix, and Thr46 and Glu48 in β -sheet 2 are exposed on the protein surface, as predicted by the model (Figure 7*D* and 7*E*). Thus,



Figure 6. Protein F (PF) promotes nontypeable *Haemophilus influenzae* (NTHi) adherence to pulmonary epithelial cells via the N-terminus. *A*, A direct enzyme-linked immunosorbent assay shows that PF binds to fixed and immobilized NCI H292 and A549 in a saturable and dose-dependent manner. *B*, Deletion of *hpf* results in a marked decrease in the adhesion of NTHi to pulmonary epithelial cell lines, as well as to primary bronchial epithelial cells from a healthy adult donor. *C*, PF²³⁻⁴⁸ shows the strongest binding to both cell lines, as revealed by a radiolabeled peptide adherence assay. *D*, The NTHi interaction with A549 cells can be inhibited by epitope-specific anti-PF²³⁻⁴⁸ polyclonal antibodies but not with anti-PF⁴⁴⁻⁶⁹. The mean of 3 separate experiments is used for all panels, and error bars indicate the standard error of the mean in all experiments. The Student *t* test was used for statistical analyses in panel *B* and for analysis of variance in panel *D*. *P* ≤ .05 was considered statistically significant. **P* ≤ .05, ****P* ≤ .001. Abbreviation: NS, not significant.



Figure 7. Bioinformatic analyses of protein F (PF)²³⁻⁴⁸. *A*, PF homologs are present in other pathogens of the Pasteurellaceae family, including Haemophilus parainfluenzae, Haemophilus somnus, Haemophilus parasuis, Haemophilus ducreyi, Pasteurella dagmatis, Pasteurella multocida, Aggregatibacter aphrophilus, Aggregatibacter segnis, Actinobacillus actinomycetemcomitans, as well as *Yersinia pestis* and Eikenella corrodens belonging to other families. Partial alignment shows the highly conserved nature of the laminin (Ln)–binding region. *B*, Alignment of PF²³⁻⁴⁸ with Lbp and Lmb of *Streptococcus pyogenes* and *Streptococcus agalactiae*, respectively. *C*, A cartoon of PF²³⁻⁴⁹ with numbered β-sheets, α-helix, and loops with their side chains shown as a ball-and-stick model. *D*, The surface of the PF molecule. The Ln-binding region (encircled) is shown with colors indicating the charge distribution (blue represents positively charged and red negatively charged residues at the surface). E, The PF²³⁻⁴⁸ surface (encircled in C) is zoomed in. Figures were prepared using PyMOL.

the structural analysis shows that the experimentally defined host component-binding region Lys23-Glu48 contains several conserved and charged residues that are accessible for interaction with Ln and host epithelial cells.

DISCUSSION

NTHi has recently been reported to bind to ECM proteins such as Ln and vitronectin [17, 18, 29]. Here, we report that the NTHi ABC transporter protein PF to be a novel Ln- and cell-binding adhesin. ABC transporters constitute a large family of proteins that play important roles for bacterial pathogenesis [30]. Although previous reports have suggested similar proteins to be implicated in interactions between gram-negative bacteria and the host [31–35], this is the first study that shows a gram-negative ABC transporter protein directly mediating bacterial adherence to host components.

NTHi primarily colonizes the upper respiratory tract of humans and causes a range of airway infections. A significant rise in the occurrence of invasive NTHi infections has recently been observed, possibly due to a niche opening after the introduction of a capsule polysaccharide-conjugate vaccine against H. influenzae serotype b [36, 37]. To disseminate from the primary site of infection, bacteria need to colonize and breach the basement membrane barrier. Pathogen interactions with ECM proteins are particularly important after epithelial disruption (such as that due to chronic inflammation and viral infections), which leaves the basement membrane exposed [5]. This is supported by the fact that NTHi infections are commonly identified in patients with COPD or during coinfections with viral pathogens [3, 38, 39]. Other respiratory tract pathogens, such as *S. pneumoniae*, *S. pyogenes, Moraxella catarrhalis*, and *Mycobacterium* species, have also been reported to interact with Ln and thereby augment their virulence [9–16]. Binding to ECM components such as Ln is thus most likely an important mechanism for bacterial pathogenesis.

In the present study, NTHi Δhpf mutants were shown to lose a significant part of their Ln-binding capacity as compared to the parental PF-producing wild-type strains. Moreover, expression of hpf in the heterologous host *E. coli* verified the role of PF as a Ln-binding protein, and colocalization of PF and Ln could be observed at the surface of PF-producing bacteria. In addition, rPF was shown to bind to both alveolar and bronchial epithelial cell lines, and NTHi 3655 Δhpf had a significant reduction in adherence to the cells as compared to the parental wild-type strain. Since NTHi 3655 Δhpf also displayed a significant loss of binding to primary cells, our collective data indicate that PF may be clinically relevant for the course of infection by promoting bacterial adherence to the host epithelium and basement membrane during colonization.

The NTHi adhesins PE (a 16-kDa lipoprotein) and Hap (a 155-kDa autotransporter) have previously been shown to interact with Ln [17, 18]. These proteins share no structural or primary sequence similarity with PF or each other, demonstrating the multifariousness of the NTHi-Ln interaction. We show that PF contributes to NTHi Ln binding to approximately the same degree as PE and Hap (Figure 3*B*). Furthermore, our data show that PF interacts with the C-terminus of the Ln α -chain (Figure 5*D* and 5*E*), the same region that PE binds to [17]. Because PE and PF are not similar, these data could possibly hint at a general pathogen-specific binding region in the Ln molecule. The presence of additional Ln-binding factors in NTHi remains to be elucidated.

PF (also annotated as a Y. pestis yfeA homologue) has been identified among other H. influenzae proteins to be upregulated during physiological conditions [40, 41]. Culturing NTHi in sputum obtained from COPD patients resulted in a 3.9-fold increase in PF concentration. Moreover, during heme- and iron-restricted growth, H. influenzae Rd KW20 was reported to enhance the hpf-operon transcription up to 11.8-fold. As the iron-concentration is low in the epithelial mucosa [42], an iron-dependent upregulation of PF may enhance the adhesive capacity of NTHi during the initial colonization of the host. These data suggest that PF is important for NTHi in vivo, possibly mediating adherence and metal ion transport.

The 30-kDa PF shares a strong structural resemblance with streptococcal Ln-binding proteins Lbp and Lmb (Figure 1*B*), al-though the primary sequence similarities are relatively weak. Preliminary unpublished data from our laboratory indicate that PF chelates Zn^{2+} via an interaction analogous to that of Lbp/Lmb [21, 22]. Assuming that the proteins share the same Ln-binding mechanism, the interactions could either be mediated by several separate but conserved sites in the tertiary structure, or they could be confined to a small local region that is highly similar in both proteins. Concurrent Zn^{2+} -binding can be an additional mechanism, as has been suggested for *S. pyogenes* Lbp [21], since Ln is also known to bind Zn^{2+} [43]. However, we found that the PF-Ln interaction was unaffected by the divalent cation binding of PF (data not shown), suggesting that the mechanism behind the PF Ln binding is a protein-protein interaction.

In conclusion, PF is characterized as a novel NTHi adhesin that mediates binding to the basement membrane glycoprotein Ln, as well as to human pulmonary epithelial cells. Our data shed light on the host colonization strategies of the multifaceted and important respiratory tract pathogen NTHi and will be the impetus for further studies.

Supplementary Data

Supplementary materials are available at *The Journal of Infectious Diseases* online (http://jid.oxfordjournals.org/). Supplementary materials consist of data provided by the author that are published to benefit the reader. The posted materials are not copyedited. The contents of all supplementary data are the sole responsibility of the authors. Questions or messages regarding errors should be addressed to the author.

Notes

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Brief report

Impact of immunization with Protein F on pulmonary clearance of nontypeable *Haemophilus influenzae*

ABSTRACT

NTHi-vaccine candidate.



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1. Introduction

Nontypeable Haemophilus influenzae (NTHi) is a mucosal pathogen and a prevalent cause of exacerbations in patients with chronic obstructive pulmonary disease (COPD), the second most common cause of bacterial acute otitis media in children and occasionally disseminates from the respiratory tract to produce invasive disease [1,2]. Otitis media is one of the most common reasons for children seeking medical care, receiving antibiotic treatment and, in the case of reoccurrence, undergoing surgery [3]. While still remaining the foremost cause of recurrent otitis media, there are indications that NTHi is rising to also become the premier acute otitis media-causing pathogen following the introduction of pneumococcal vaccines [1,4,5]. There have been contradictory reports regarding the effect of the NTHi Protein D (PD)-conjugated pneumococcal vaccine on the carriage rates of NTHi in children, although NTHi-mediated acute otitis media was observed to decrease by approximately 30% [6-9]. The great socioeconomic burden of otitis

Nontypeable Haemophilus influenzae (NTHi) is one of the main aetiologies of childhood bacterial infec-

tions as well as exacerbations in COPD patients. Currently, no licensed NTHi vaccine exists. In the present

study, we evaluated the potential of the conserved and ubiquitous surface protein Haemophilus Pro-

tein F (PF) as a vaccine candidate. Our results show that incubation of NTHi with anti-PF antibodies

significantly increased the opsonophygocytosis of human promyelocytic leukemia cell line-derived granulocytes, leading to efficient killing of the bacteria ($P \le 0.05$). The presence of anti-PF IgG titers in healthy

adults (n = 60) was investigated, and we found that 26% of healthy blood donors carried antibodies with

the main antigenic epitope being PF²³⁻⁴⁸. Finally, mice immunized with PF²³⁻⁴⁸ attained a significantly

increased capacity to clear NTHi as compared to a control group immunized with a peptide derived from

Moraxella catarrhalis β -lactamase ($P \le 0.05$). Taken together, our results indicate that PF is a potential

media, in parallel with exacerbations in COPD patients, warrants the continued pursuit of a more potent NTHi vaccine.

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We recently identified NTHi Protein F (PF), a surface-exposed and highly conserved protein mediating bacterial adhesion to host tissue and complement-evasion [10,11]. Protein F belongs to the ATP-binding cassette (ABC)-transporter superfamily, members of which have shown promise as potential vaccines candidates [12]. Indeed, a multiantigen-vaccine containing the *Staphylococcus aureus* ABC-transporter component MntC, a PF-homologue, is currently undergoing human clinical trials [13].

To study the potential of PF as a vaccine candidate, we tested the promotion of opsonophagocytosis of NTHi by raising rabbit anti-PF antibodies, investigating the presence of anti-PF antibodies in healthy adults and, finally, evaluating the prophylactic effect of PFimmunization *in vivo* in a murine pulmonary clearance model [14]. Our results indicate that PF may be a potential vaccine candidate against NTHi.

2. Material and methods

Bacterial strains, antibodies and recombinant constructs. NTHi3655 and KR403 were routinely cultured in brain heart infusion broth and chocolate agar plates as described [10,15]. A human promyelocytic leukemia cell line (HL60) was propagated in RPMI 1640 supplemented with 10% fetal bovine serum (Fetalclone

Abbreviations: Abs, antibodies: CFU, colony forming units: COPD, chronic obstructive pulmonary disease: NTHi, non-typeable Haemophilus influenzae: OPA, opsonophagocytic assay: PD, Protein D; PF, Protein F.

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Fig. 1. Immunization of rabbits with Protein F elicits a humoral immune response against specific epitopes and the resulting Abs promote opsonophagocytosis of NTHi. (A) PF^{44-60} , PF^{44-60} , PF^{45-60} ,

I; HyClone, Logan, UT), 1% L-glutamine and penicillin/streptomycin (Invitrogen). Recombinant PF, PD and affinity-purified rabbit anti-PF and anti-*Moraxella catarrhalis* Ubiquitous Surface Protein A (UspA) antibodies (Abs) were raised as previously described [10,16,17].

Indirect enzyme-linked immunosorbent assay (ELISA). Synthetic peptides spanning the entire PF molecule (1 μ M), recombinant full-length PF and PD and Escherichia coli BL21 lysate (100 ng/well) were immobilized on PolySorp (Nalge Nunc International, Rochester, NY) microtiter plates followed by ELISA as previously described [10]. Affinity-purified rabbit anti-PF polyclonal antibodies (pAbs) or sera obtained from healthy blood donors aged 18–65 years (n = 60) were used as primary Abs. Sera reacting with the *E. coli* BL21 lysate were excluded from analysis to prevent cross-reactivity with impurities in the recombinantly expressed proteins (n = 10). Titers were quantified as previously described [16].

Opsonophagocytic assay (OPA). A standardized OPA [18] was employed to determine the functionality of rabbit anti-PF Abs. Briefly, differentiation of HL60 cells into granulocytes was induced by suspending cells into culture medium containing 1.25% dimethyl sulfoxide for 5 days. Baby rabbit complement (Pel-freez Biologicals Rogers, AR) was used as a standardized source of complement. Bacteria were pre-incubated with various titers of affinity-purified Abs for 30 min at RT on a mini-orbital shaker. Subsequently, bacteria and HL60 cells were washed twice and diluted to 10^5 and 10^7 CFU/ml, respectively, and incubated together with baby rabbit complement for 45 min on a mini-orbital shaker at 37° C, 5% CO₂. Fifty µl of the final reaction mixture was plated out and incubated overnight in 37° C, 5% CO₂. Experiments with a nonspecific killing (in the absence of Abs) \geq 30% were excluded.

Pulmonary clearance model, BALB/c mice (n=69) at 8 weeks of age were subcutaneously immunized with 50 µg keyhole limpet hemocyanin (KLH)-conjugated PF²³⁻⁴⁸ peptide or the KLHconjugated control peptide M. catarrhalis B-lactamase²⁶⁻⁴⁵. The first immunization was done with Freund's complete adjuvant. After four weeks, mice were immunized three more times at 7 day-intervals using alum as adjuvant (under the guidelines of the Ethical Committees on Animal Experiments, the Swedish Board of Agriculture, Freund's incomplete adjuvant is not allowed to be administered more frequently than once every fourth week. Alum, however, is allowed for administration once a week.). The animals were challenged with 10⁶ CFU of NTHi3655 via the intranasal route seven days post-final immunization and bacterial load in the lungs were quantified as described previously [19]. As a mean of normalization, the highest (n=1) and lowest (n=1) values for each group were excluded. Mouse sera were collected and ELISA verified the presence of anti-PF Abs. Three animals that displayed weak/no humoral response to the antigen were excluded from the analysis. Statistics. Student's *t* test and two-way analysis of variance (ANOVA) was used for statistical calculations of two and several sets of data, respectively. Statistical analyses were performed using GraphPad Prism 5 (GraphPad Software). A *P* value of \leq 0.05 was considered statistically significant.

3. Results and discussion

We have previously shown that *Haemophilus* Protein F (PF) binds to the host extracellular matrix protein laminin, the complement-regulator vitronectin and to epithelial cells via an N-terminal region comprising amino acid residues Lys23-Glu48 (PF²³⁻⁴⁸) [10,11]. To determine the immunodominant regions of PF, we raised and affinity-purified rabbit anti-PF pAbs using full-length recombinant protein. We analyzed the resulting IgG directed against synthetic peptides and identified PF⁴⁴⁻⁶⁸, PF⁴⁰⁻¹²⁸ and PF²⁵⁻⁴⁸, to contain the most immunodominant epitopes (Fig. 1A). Interestingly, we found only low titers of Abs directed towards PF²³⁻⁴⁸, whereas the region (PF⁴⁴⁻⁶⁸) adjacent to PF²³⁻⁴⁸ was one of the most immunogenic.

To test the functionality of the anti-PF Abs, anti-PF⁴⁴⁻⁶⁸ IgG directed against the N-terminal part of PF was affinity-purified from sera obtained from rabbits immunized with full-length PF. Moreover, we immunized rabbits with KLH-conjugated PF²³⁻⁴⁸ to produce specific rabbit anti-PF²³⁻⁴⁸ Abs that was also subsequently affinity-purified. The PF N-terminus is interesting from a host-interaction point of view since it targets epithelial cells and proteins derived from the extracellular matrix. Antibodies against this region would thus be effective in both blocking the host interaction as well as opsonizing the bacteria. As shown in Fig. 1B and C, both affinity-purified pAbs significantly promoted antibody-dependent opsonophagocytosis of NTHi3655 (Fig. 1B) and the heterologous strain KR403 isolated from the subepithelial compartment of a palatine tonsil [15] (Fig. 1C) as compared to affinity-purified anti-M. catarrhalis UspA IgG that was included as a negative control.

To investigate the presence of specific anti-PF IgG in a healthy adult population, sera obtained from healthy blood donors (n = 60) were analyzed. Since almost all humans are exposed to NTHi during childhood, we expected to observe the natural serological anti-PF IgG response in humans. Approximately 26% of the healthy blood donors were positive for anti-PF IgG (IgG concentration > highest concentration of IgG directed against the *E. coli* lysate control) (Fig. 2A), which corresponded to the anti-PD IgG levels (26% of donor sera) in our experimental set up. Surprisingly, no correlation between carriage of anti-PF and anti-PD IgG was found, indicating that humans may have different and individual serological responses to NTHi with regard to these antigens

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Fig. 2. Antibodies directed against Protein F exist in 26% of healthy blood donors, and primarily react against the epitope comprising PF²³⁻⁴⁸. (A) Sera from healthy adult blood donors (n = 60) were tested for IgG reactivity against recombinant PF and PD at a dilution of 11.000. Those sera reactive to total cell lysate of *E. coli* BL21, which was the expression host of recombinant PF and PD. were excluded from the analysis to prevent a false positive bias (n = 10). (B) No correlation was found between high titers of anti-PF and anti-PD IgG ($R^2 = 0.0431$, two-tail Pearson's r). (C) Protein F-reactive sera (n = 8) were tested for epitope recognition. Three sera reacted only against full-length PF (data not shown). The remaining five sera had the highest IgG-titers against PF²³⁻⁴⁸, which was unexpected since a low immunogenicity of this region was observed in rabbit. All experiments were repeated three times and the mean ± SEM is shown. In panel A, a statistical analysis was performed using two-tail Student's t test, and $P \le 0.05$ was considered statistically significant. " $P \ge 0.001$.

(Fig. 2B). Immunization with both proteins may thus complement each other, potentially in a multicomponent vaccine including other immunogenic proteins such as Protein E [19].

To delineate the immunodominant epitopes, donor sera with high concentrations of anti-PF Abs (defined as two-fold higher concentrations than to the *E. coli* lysate negative control) were selected (n = 8) for further analysis. Three of these sera reacted only against full-length PF, suggesting that the Abs recognized epitopes not encompassed by the shorter, presumably linear, peptides (data not shown). Surprisingly, however, we found that the most immunodominant epitope in the remaining five sera was PF²³⁻⁴⁸ (Fig. 2C), which was in stark contrast to the epitope reactivity of the rabbit anti-serum (Fig. 1A). This confirmed that animals and humans do not have the same serological response to antigens with regard to epitope immunodominance [20]. The fact that NTHi is a human-specific pathogen may explain why humans, but not rabbits, develop Abs towards the PF N-terminus that interacts with host components.

Finally, we investigated the effect of PF immunization in a mouse pulmonary clearance model. PF²³⁻⁴⁸ that binds to host components, and to which the highest IgG concentration was found in humans, was chosen as the antigen. Mice were subcutaneously immunized four times with KLH-conjugated PF²³⁻⁴⁸ or the negative control peptide β -lactamase²⁶⁻⁴⁵ derived from *M. catarrhalis* (Fig. 3). After 6 weeks, mice were challenged intranasally with NTHi3655 and



Fig. 3. Mice immunized with Protein F have a significantly increased clearance of NTHi. Mice immunized with PF^{23-48} or the negative control peptide M. catarrhalis B-lactanase²²⁻⁶⁴ were challenged intranasally with NTHi 3655 and bacterial loads in the lungs were quantified at 0 h (baseline control), 3 h and 5 h post-challenge. Statistical analysis was performed using one-tail Student's t test, and $P \le 0.05$ was considered statistically significant. P < 0.05.

clearance was monitored after 3 and 5 h. Our results revealed that mice immunized with the Pr²³⁻⁴⁸ peptide exhibited significantly better NTHi clearance at both tested time points as compared to the control group ($P \le 0.05$).

4. Conclusions

The efficiency of anti-PF Abs in promoting opsonophagocytosis, in addition to an increased pulmonary clearance of NTHi in mice immunized with PF, suggests that this highly conserved and ubiquitous surface-protein would be a potential vaccine candidate against NTHi. Our results will thus be the impetus for further studies.

Conflict of interest statement

None.

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