Covalent bonding of heme to protein prevents heme capture by nontypeable Haemophilus influenzae

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Nontypeable Haemophilus influenzae (NTHi) are Gram-negative pathogens that contribute to a variety of diseases, including acute otitis media and chronic obstructive pulmonary disease. As NTHi have an absolute requirement for heme during aerobic growth, these bacteria have to scavenge heme from their human hosts. These heme sources can range from free heme to heme bound to proteins, such as hemoglobin. To test the impact of heme structural factors on heme acquisition by NTHi, we prepared a series of heme sources that systematically vary in heme exposure and covalent binding of heme to peptide/protein and tested the ability of NTHi to use these sources to support growth. Results from this study suggest that NTHi can utilize protein-associated heme only if it is noncovalently attached to the protein.

Abbreviations
AOM, acute otitis media; BHI, brain heart infusion; COPD, chronic obstructive pulmonary disease; CXXCH, Cys-X-X-Cys-His; cyt, cytochrome; E. coli, Escherichia coli; Hib, Haemophilus influenzae type b; Ht-C12A, Ht C12A cyt c; Ht-C15A, Ht C15A cyt c; Ht-DM, Ht C12A/C15A cyt c; Ht, Hydrogenobacter thermophilus; LB, Luria broth; MP-11, microperoxidase 11; NTHi, Nontypeable Haemophilus influenzae; ppIX, protoporphyrin IX.
the highest known binding affinities to heme [11]. During heme acquisition, a conformational change in HxuA induces structural changes in hemopexin, lowering its affinity for heme and resulting in heme release to the HxuC receptor [10]. In addition to the HxuABC system, the HemR and Hgp/Hup systems have been linked to free heme acquisition and heme acquisition from hemoglobin in *H. influenzae* type b (Hib), an encapsulated strain of *H. influenzae* [12–14].

While most studies have focused on hemoproteins such as hemoglobin and hemopexin, one study demonstrated that horse cytochrome (cyt) c failed to provide an adequate heme source to Hib [15]. Unlike hemoglobin and hemopexin, the heme group in cyt c proteins, including human cyt c (which shares 89% sequence identity with horse cyt c), is covalently attached to the protein via a Cys-X-X-Cys-His (CXXCH) motif (Fig. 1). Whether horse cyt c failed to interact with Hib’s hemophore(s) or the hemophore(s) were unable to extract heme due to the covalent linkages was not addressed in that study.

In this work, we sought to further our understanding of the effect of heme-binding mode and solvent exposure on heme scavenging by the pathogenic bacteria NTHi. As a heme source, we used single (C12A or C15A) and double (C12A/C15A) site-directed mutants of *Hydrogenobacter thermophilus* cyt c in which the Cys residues of the C12XXC15H motif (Fig. 1) are mutated to Ala [16]. *Ht* cyt c was selected as an ideal model system for this study, because its inherent stability allows the double mutant protein to maintain a folded structure with heme bound in the absence of thioether linkages to heme [17]. This property allowed us to determine the effect of covalent attachment on heme availability within one protein scaffold. To determine whether heme that is solvent-exposed but has covalent bonds to two Cys can serve as a heme source for NTHi, we supplemented NTHi cultures with microperoxidase-11 (MP-11), an 11-mer peptide with covalently bound heme [18]. This series of heme sources (*Ht* cyt c, its mutants, MP-11, and hemin) systematically varies in heme exposure and covalent binding to a peptide, providing a means to test the impact of heme structural factors on heme acquisition.

**Materials and methods**

**Mutagenesis, expression, and purification**

Plasmid pSCH552 (ampicillin resistant) was used as a template for site-directed mutagenesis of *Ht* cyt c [19]. The pSCH552 plasmid contains the *Thiobacillus versutus* cyt c550 signal sequence inserted in frame and upstream of the *Ht* cyt c gene, directing excretion to the periplasm for maturation (covalent heme attachment) in *Escherichia coli* (E. coli). The two single mutants, *Ht* C12A cyt c (*Ht*-C12A) and *Ht* C15A cyt c (*Ht*-C15A), were prepared using the QuikChange II kit (Agilent, Santa Clara, CA, USA) according to the manufacturer’s instructions and confirmed by DNA sequencing. The plasmid pEST202 encoding *Ht* C12A/C15A cyt c (*Ht*-DM) was a gift from Stuart Ferguson [17]. *Ht* wild-type, C12A, and C15A cyt c were expressed and purified as previously described for the wild-type protein [19]. *Escherichia coli* BL-21(DE3) cells were cotransformed with the plasmid harboring the cyt c gene and the pEC86 plasmid (chloramphenicol resistant), which contains the cyt c maturation genes *cmABCDEFGH* [20]. The pEC86 plasmid was a gift to Kara Bren from Linda Thöny-Meyer. *Ht*-DM was expressed and purified as described [17], with the following modifications. The *E. coli* cells were grown on Luria broth (LB) agar plates, and single colonies were used to inoculate 25 mL LB (supplemented with 50 µg·mL⁻¹ ampicillin). Small cultures were grown for 16 h (180 rpm, 37 °C) and used to inoculate large cultures (in 1 L LB, 140 rpm, 37 °C). Protein expression was induced with 1 mM IPTG when the optical density of the cultures at 600 nm (OD₆₀₀) reached ≈0.6; cultures were incubated for an additional 3 h before harvesting by centrifugation at 5000 g for 15 min.

**NTHi growth experiments**

Nontypeable *H. influenzae* cultures were grown on brain heart infusion (BHI) medium (BD) supplemented with

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**Fig. 1.** The heme and CXXCH motif from *Hydrogenobacter thermophilus* cyt c (Protein Data Bank ID: 1YNR). One or both covalent linkages between peptide and heme (arrows) are absent in the single mutants (*Ht* C12A cyt c, *Ht* C15A cyt c) and the double mutant (*Ht* C12A/C15A cyt c), respectively. The His residue (His16) coordinates to the heme iron.
content: 

30 μM NAD⁺ (Sigma, St. Louis, MO, USA) and either water (negative control) or a single heme source. NTHi lack the enzymes required to synthesize the essential cofactor NAD⁺ and therefore must uptake exogenous NAD⁺. NTHi 86-028NP is a pediatric isolate (a gift from Lauren Bakaletz, The Research Institute at Nationwide Children's Hospital) [21]. In the first set of NTHi growth experiments, heme sources were 15 μM hemin (Sigma), 15 μM purified human hemoglobin (Sigma), or 15 μM wild-type Ht cyt c, Ht-C12A, or Ht-DM. A small 500 μL aliquot was removed from each 10 mL sample immediately after the supplements were added, but prior to inoculation, to be used as blanks for the OD readings. The cultures were incubated at 37 °C, shaken at 180–200 rpm, and monitored for changes to the OD₄₉₀ every 2 h. In a second set of NTHi growth experiments, three different concentrations of hemin (1 μM, 0.1 μM, and 0.01 μM) and Ht-DM (1 μM, 0.1 μM, and 0.01 μM) were used to supplement BHI.

When using Ht-C12A as a heme source, the proportion of heme covalently bound to the protein had to be considered. Prior work on Ht-C12A revealed that ~14% of the sample had noncovalently bound heme, with the remainder covalently bound [17]. To consider this factor, we supplemented BHI with a ‘high’ concentration of Ht-C12A (6–7 μM) to test whether these conditions provided sufficient ‘free’ noncovalently bound heme for NTHi to grow [16]. We also grew NTHi in BHI supplemented with a ‘low’ concentration of Ht-C12A (0.6 μM). As positive controls, we grew NTHi in BHI supplemented with 6–7 μM hemin and 0.6 μM hemin. All heme sources were diluted in 1 mL of water and added to 5 mL of sterile BHI and 30 μM NAD⁺. Growths were incubated (37 °C, 180–200 rpm) for 24 h. Readings (OD₄₉₀) were recorded after 12 h of growth and then every 2 h.

Nontypeable H. influenzae were also grown on BHI using 15 μM hemin, 15 μM MP-11 (Sigma), or water (negative control) as its heme source. The NTHi were cultured as described above, and the OD₄₉₀ was monitored every 2 h.

**Results and Discussion**

Nontypeable H. influenzae have a strict requirement for heme as their source of iron, thus providing biological motivation for their complex and redundant heme acquisition and uptake systems [4,10,12–14,22]. Hemophores play an important role in those systems. Secreted into the extracellular space or tethered to the bacterial surface, hemophores serve as the initial heme ‘scavengers’ by stealing heme from hemoproteins directly or by lowering the hemoprotein’s affinity for heme. Cyt c is ubiquitous hemoproteins, highly conserved across species, and found in most living organisms. Defined, in part, by their two covalent thioether linkages to heme, cyt c exhibit a diverse array of cellular functions in such biological arenas as energy transduction and apoptosis [23]. Our Ht cyt c variants, with one, two, or zero covalent linkages to heme, provide a means to test what requirements and limitations there are on heme acquisition by NTHi with respect to the mode of heme binding to polypeptide.

To determine the ability of NTHi to scavenge heme from Ht cyt c and variants with one or two thioether linkages removed, we cultured NTHi in BHI, supplemented with 30 μM NAD⁺ and (a) sterile water (negative control); (b) 15 μM hemoglobin; (c) 15 μM hemin (positive control); (d) 15 μM wild-type Ht cyt c; (e) 15 μM Ht-C12A; or (f) 15 μM Ht-DM. For all experiments, hemin was used as a positive control. Hemoglobin contains noncovalently bound heme [24]. Growth was tracked by measuring the OD at 490 nm every 2 h. NTHi showed minimal growth when BHI was supplemented with sterile water, wild-type Ht cyt c, or Ht-C12A (Fig. 2). In comparison, NTHi grew to log phase and approached stationary phase by hour twelve when BHI was supplemented with hemoglobin, hemin, or Ht-DM (where both thioether linkages were absent) (Fig. 2). Two additional independent growth experiments using the same parameters and supplements yielded similar results (Fig. S1), which suggests that NTHi hemophore(s) can access and extract heme from Ht-DM.

In a second growth study, we determined that 1 μM hemin supports NTHi growth, while 0.1 μM hemin is insufficient for NTHi growth. We observed similar growth trends for the equivalent concentrations of Ht-DM (Fig. 3), indicating that heme noncovalently bound to this protein scaffold supports growth as well as hemin. Two additional independent growth
experiments using the same parameters and supplements yielded similar results (Fig. S2), which suggest that both of these heme sources (hemin and Ht-DM) are similarly accessible to NTHi hemophore(s). In contrast to NTHi, which we found require 0.1–1.0 μM hemin for unrestricted growth, Hib requires a minimum of 0.03 μM free heme for unrestricted growth, indicating different heme requirements for these bacteria in culture [15]. Further, we predict that NTHi cultured under different conditions will have different heme requirements, not surprisingly as H. influenzae has been shown to exhibit different heme requirements in the presence and absence of oxygen [25]. As NTHi are pathogens (causing AOM in the inner ear and bronchitis and COPD in the lungs) and commensal organisms (in the nasal cavity), these nuances in NTHi’s heme requirements may come into play as the bacteria live and thrive in different environments.

In Ht-C15A, only the Cys12 residue is able to covalently link to the heme group, while in Ht-C12A, only the Cys15 residue covalently links to the heme. However, it has been shown that the Cys15-to-heme covalent bond does not form in 100% of the Ht-C12A protein sample, when overexpressed in E. coli [16]. Specifically, Tomlinson and Ferguson showed that the heme group in Ht-C12A is covalently bound in 86% of the sample, while the heme in Ht-C15A is bound in 100% of the sample [16]. To determine whether the small proportion of noncovalently bound heme in Ht-C12A could support NTHi growth, we cultured NTHi in the presence of each single mutant at high concentrations. Results from our previous experiment suggested that NTHi requires between 0.1 μM and 1 μM accessible heme to grow. Assuming ~14% of Ht-C12A contains noncovalently bound heme, we estimated that we would need to supplement BHI with between 0.7 μM and 7 μM of Ht-C15A to observe growth similar to what was seen using excess hemin. As such, we grew NTHi in the presence of either high (6–7 μM) or low (0.6 μM) concentrations of hemin, Ht-C12A, or Ht-C15A [16]. NTHi cultures reached log phase in less than 12 h when supplemented with high and low concentrations of hemin, but did not grow in the presence of the negative control, high or low concentrations of Ht-C15A, or low concentrations of Ht-C12A. When supplemented with high concentrations of Ht-C12A, NTHi grew slowly, but eventually reached similar OD₄₉₀ levels to the hemin-supplemented growths by 24 h (Fig. 4). Two additional independent growth experiments using the same parameters and supplements yielded similar results (Fig. S3). The logarithmic growth phase was not reached until approximately 18 h. The delay in growth may have been the result of the noncovalently bound heme in Ht-C12A being less accessible to NTHi than the equivalent concentration of hemin.

Microperoxidase-11 (MP-11) is a heme peptide derived from proteolysis of horse heart cyt c [18]. The heme in MP-11 is covalently attached to an 11-mer peptide via the two Cys residues contained within the

![Fig. 3. Growth curves, monitored at 490 nm, are similar for NTHi cultured in BHI supplemented with 1 μM hemin and 1 μM Ht-DM (C12A/C15A); lower concentrations (0.1 μM or 0.01 μM) of hemin or Ht-DM do not support NTHi growth.](image)

![Fig. 4. Nontypeable Haemophilus influenzae growth curves, monitored at 490 nm after 12 h of growth. NTHi cultures in BHI were supplemented with high (6–7 μM) or low (0.6 μM) concentrations of hemin, Ht-C12A, or Ht-C15A. NTHi grew with both concentrations of hemin and grew slowly with high concentrations of Ht-C12A.](image)

![Fig. 5. Nontypeable Haemophilus influenzae growth curves, monitored at 490 nm; NTHi cultures in BHI were supplemented with 15 μM hemin, water, or 15 μM MP-11.](image)
CXXCH motif, and the heme iron is coordinated to a single His residue donated by the same sequence. If heme uptake requires binding to a hemophore or receptor protein in a ‘tight’ pocket, as described for the heme transport protein HbpA, even a small attached peptide could prevent proper binding/fitting of the heme [26]. Additionally, the coordinated His of MP-11 may not be readily displaced, because it is positioned in the covalently bound peptide; this His–heme interaction could also hinder its interaction with a hemophore. In order to determine whether MP-11 can serve as a heme source for NTHi, we repeated the initial NTHi growth experiment, using hemin (positive control), MP-11, or water (negative control) as the heme source. Results from the experiment suggest that the covalent attachment of the heme to the 11-mer peptide prevents MP-11 from serving as a heme source for NTHi (Fig. 5). Two additional independent growth experiments using the same parameters and supplements yielded similar results (Fig. S4).

Conclusions

Nontypeable H. influenzae are known to colonize in the nasopharynx of humans. Considering their requirement for heme, NTHi must acquire heme from human proteins such as hemoglobin or complexes such as heme bound to hemopexin. Cyt c, containing two covalent linkages to heme, have failed to serve as a heme source for H. influenzae. Here, we showed that even a single thioether bond between heme and polypeptide is sufficient to prevent heme scavenging from Hp cyt c mutants by NTHi. However, NTHi hemophores can interact with and extract heme from Hp-DM, which contains zero covalent linkages to heme. We also confirmed that the C15-to-heme bond does not form 100% of the time in Hp-C12A, allowing for heme scavenging by NTHi from high concentrations of this variant. Finally, we showed that MP-11, which is similar in structure to heme, but covalently tethered to an 11-mer peptide, does not function as a heme source for NTHi. Results from this study suggest that the only accessible protein-associated heme to support NTHi growth is noncovalently bound heme.

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Author contributions

VS, BN, SS, JP, BK, MFH, and KG collected experimental data and assisted with data analysis. VS, KLB, and LVM designed the study, analyzed data, and prepared the manuscript. KLB and LVM provided reagents.

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**Supporting information**

Additional Supporting Information may be found online in the supporting information tab for this article:

**Fig. S1.** Two additional NTHi growth curves, monitored at 490 nm, show that NTHi grows in BHI supplemented with hemoglobin, hemin, or Ht-DM (C12A/C15A), but does not grow with water, wild-type Ht cyt c, or single mutant Ht-C12A.

**Fig. S2.** Two additional NTHi growth curves, monitored at 490 nm, show that NTHi grows in BHI supplemented with 1 mM hemin or Ht-DM (C12A/C15A), but does not grow with lower concentrations of hemin or Ht-DM.

**Fig. S3.** Two additional NTHi growth curves, monitored at 490 nm after 12 h of growth.

**Fig. S4.** Two additional NTHi growth curves, monitored at 490 nm after 12 h of growth.