Molecular Signatures of Non-typeable *Haemophilus influenzae* Lung Adaptation in Pediatric Chronic Lung Disease

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Non-typeable *Haemophilus influenzae* (NTHi), an opportunistic pathogen of the upper airways of healthy children, can infect the lower airways, driving chronic lung disease. However, the molecular basis underpinning NTHi transition from a commensal to a pathogen is not clearly understood. Here, we performed comparative genomic and transcriptomic analyses of 12 paired, isogenic NTHi strains, isolated from the nasopharynx (NP) and bronchoalveolar lavage (BAL) of 11 children with chronic lung disease, to identify convergent molecular signatures associated with lung adaptation. Comparative genomic analyses of the 12 NP-BAL pairs demonstrated that five were genetically identical, with the remaining seven differing by only 1 to 3 mutations. Within-patient transcriptomic analyses identified between 2 and 58 differentially expressed genes in 8 of the 12 NP-BAL pairs, including pairs with no observable genomic changes. Whilst no convergence was observed at the gene level, functional enrichment analysis revealed significant under-representation of differentially expressed genes belonging to Coenzyme metabolism, Function unknown, Translation, ribosomal structure, and biogenesis Cluster of Orthologous Groups categories. In contrast, Carbohydrate transport and metabolism, Cell motility and secretion, Intracellular trafficking and secretion, and Energy production categories were over-represented. This observed trend amongst genetically unrelated NTHi strains provides evidence of convergent transcriptional adaptation of NTHi to pediatric airways that deserves further exploration.

Understanding the pathoadaptative mechanisms that NTHi employs to infect and persist in the lower pediatric airways is essential for devising targeted diagnostics and treatments aimed at minimizing disease severity, and ultimately, preventing NTHi lung infections and subsequent chronic lung disease in children.

**Keywords:** non-typeable *Haemophilus influenzae*, NTHi, RNA-seq, transcriptomics, comparative genomics, convergence, bacterial evolution, adaptation
INTRODUCTION

Non-typeable *Haemophilus influenzae* (NTHi) is a Gram-negative bacterium that frequently colonizes the respiratory tract mucosa of humans. NTHi is part of the upper airway flora and is found in 20–50% of healthy children and 20–30% of healthy adults (Slack, 2015). The upper airways may act as a reservoir for seeding the lung environment (Fothergill et al., 2014), where NTHi switches to an opportunistic pathogenic lifestyle, a transition driven by multiple, poorly understood factors (Duell et al., 2016).

NTHi facilitates the colonization of other bacterial species in the lower airways (Van Eldere et al., 2014) and is associated with exacerbations in several respiratory diseases (Duell et al., 2016). Indeed, its presence in the lower airways is associated with increased risk of future development of bronchiectasis (Wurzel et al., 2016). In children with chronic secretory lung disease (COPD), bronchiectasis, or COPD with bronchiectasis (Martinez-Garcia et al., 2011; Sriram et al., 2018), resulting in increased airway inflammation (Tufvesson et al., 2015). NTHi also has the capacity to form biofilms (Murphy and Kirkham, 2002; Swords, 2012) resulting in a decreased susceptibility to antibiotics and increased inflammatory and defense responses in the host (Starner et al., 2006). Unsurprisingly, NTHi is a pathogen with increasing public health recognition (Van Eldere et al., 2014; Slack, 2017).

NTHi has evolved several adaptive mechanisms that enable the bacterium to survive and persist in the human host, including modification of its genome via homologous recombination and phase variation (Erwin et al., 2005; Power et al., 2012). Modulation of gene expression through phase variation enables NTHi to rapidly adapt to and colonize new environments (Wong and Akerley, 2012). Analysis of the NTHi genome has identified several phase-variable virulence factors involved in persistence, nutrient uptake, cellular adherence, and host immune response evasion (Pettigrew et al., 2018), with direct modulation of gene expression in response to environmental stimuli and host milieu (Baddal et al., 2015). NTHi is naturally competent and can undergo high rates of recombination, resulting in greater genetic and phenotypic diversity than serotypeable *H. influenzae* (Connor et al., 2012); however, unlike other chronic respiratory infections (Smith et al., 2006; Price et al., 2013), no significant gene loss or gain has yet been observed in chronic *H. influenzae* infections (Pettigrew et al., 2018).

Identifying signals of NTHi adaptation to the lung would aid in unmasking the mechanisms facilitating pathogenesis, which is critical for improving diagnosis, treatment and prevention of NTHi-associated lower airway diseases. However, the mechanisms involved in the transition from a commensal state in the nasopharynx (NP) to a pathogenic lifestyle in the lower airways is not well understood (De Chiara et al., 2014; Pettigrew et al., 2018).

In this study, we investigated genome- and transcriptome-wide differences between 12 isogenic NTHi pairs obtained from the NP and lower airways (via bronchoalveolar lavage; BAL) of patients with bronchiectasis or CSLD. We hypothesized that NTHi isolated from the airways of patients with lung disease would exhibit characteristic genetic or transcriptional profiles that signaled a change from a commensal to a pathogen.

MATERIALS AND METHODS

Isolate Collection

NP swabs and BAL specimens were collected concurrently from Australian children (mean, 3.2 years; range: 1.1–13.9 years) diagnosed with bronchiectasis or CSLD (Table 1) at participating hospitals in the Northern Territory and Queensland (Hare et al., 2010). All children had an active lower airway infection, as defined by >10^4 colony-forming units of respiratory bacteria/mL BAL fluid (Hare et al., 2010). Following collection, specimens were transported on ice in 1 mL skim milk, tryptone, glucose and glycerol broth (STGGB) (Gibson and Khoury, 1986; Hare et al., 2010) and stored at −80°C before processing. All subsequent cultures were also stored in STGGB at −80°C. Following selective culture on chocolate blood agar (CBA) supplemented with bacitracin, vancomycin, and clindamycin (Oxoid, Scoresby, VIC, Australia), NTHi was isolated and confirmed as previously described (Hare et al., 2010; Price et al., 2017). Antibiotic sensitivity was determined by disk diffusion as per the Calibrated Dichotomous Susceptibility (CDS) testing guidelines (8th Edition) (Bell et al., 2016) for ampicillin, amoxicillin-clavulanate, cefaclor, ceftriaxone, chloramphenicol, tetracycline, and azithromycin.

Whole-Genome Sequencing and Comparative Genomics

DNA extraction was performed as previously described (Price et al., 2015). Whole-genome sequencing (WGS) was carried out at the Australian Genome Research Facility (Parkville, VIC, Australia) using the Illumina (San Diego, CA, United States) HiSeq 2500 platform. As a criterion for our study, all isolate pairs were required to be highly genetically similar (i.e., isogenic). To identify isogenic NP-BAL pairs from each patient, multilocus sequence typing (MLST) (Jolley and Maiden, 2010) followed by phylogenetic analysis was performed. Further details on isolate selection is described in Supplementary Methods. Based on this analysis, we selected 12 paired NP-BAL isolates from 11 patients for further investigation; two genetically distinct NP-BAL pairs were selected from one patient (designated 60373_P1 and 60373_P2).

To catalog the mutations separating each NP-BAL pair, SPANDx v3.2 (default settings) (Sarovich and Price, 2014) was used. Where possible, genetic variants were also confirmed using transcriptomic (RNA-seq) data. For each analysis, the NP isolate draft assembly was used as the reference for variant calling. Further details are provided in Supplementary Methods.
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### TABLE 1
Summary of genomic and transcriptomic differences between paired, isogenic NTHi isolates obtained from the nasopharynx (NP) and lower airways (bronchoalveolar lavage; BAL) of 11 Australian children with chronic lung disease.

| Patient ID | Site of isolation | Isolate no. | No. SNPs | No. indels | No. differentially expressed genes (up/down) | Genes with transcriptional and genetic alterations |
|------------|------------------|-------------|----------|------------|-----------------------------------------------|--------------------------------------------------|
| 60051      | NP               | 1           | 0        | 1          | 58 (4/54)                                      | hsdM3                                            |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 60068      | NP               | 3           | 0        | 1          | 54 (1/53)                                      | –                                                |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 60069      | NP               | 2           | 0        | 1          | 0                                              | –                                                |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 60294      | NP               | 1           | 0        | 0          | 0                                              | –                                                |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 60295      | NP               | 1           | 1        | 0          | 46 (38/8)                                      | –                                                |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 60316      | NP               | 1           | 0        | 0          | 0                                              | –                                                |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 60362      | NP               | 1           | 0        | 0          | 15 (14/1)                                      | –                                                |
|            | BAL              | 2           |          |            |                                               |                                                  |
| 60370      | NP               | 2           | 0        | 0          | 2 (2/0)                                        | –                                                |
|            | BAL              | 2           |          |            |                                               |                                                  |
| 60373_P1   | NP               | 1           | 0        | 1*         | 5 (3/2)                                        | hsdM3                                            |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 60373_P2   | NP               | 3           | 0        | 0          | 4 (4/0)                                        | –                                                |
|            | BAL              | 4           |          |            |                                               |                                                  |
| 65001      | NP               | 2           | 0        | 3          | 0                                              | –                                                |
|            | BAL              | 1           |          |            |                                               |                                                  |
| 65290      | NP               | 3           | 1        | 0          | 4 (4/0)                                        | –                                                |
|            | BAL              | 4           |          |            |                                               |                                                  |

SNP, single-nucleotide polymorphism; indel, insertion-deletion. * The BAL strain from this patient contains a mixture of two loss-of-function indel mutations in hsdM3 at a 40:60 ratio (see Figure 1).

### NTHi Liquid Media Growth Conditions for RNA Harvest

Due to the reliance of NTHi growth on the presence of X (haemin/haematin) and V (nicotinamide-adenine-dinucleotide) blood factors (Holt, 1962), we initially attempted to culture isolates in brain heart infusion (BHI) broth supplemented with Haemophilus Test Medium (HTM; Oxoid), and subsequently, a HTM-supplemented artificial sputum medium formulated to mimic the mucosal environment in the airways of cystic fibrosis patients (Fung et al., 2010; Price et al., 2017). However, due to inconsistent optical density (OD) readings and poor growth or flocculation of some or all isolates in these media, we investigated an alternative liquid medium for the growth of NTHi for RNA-seq (see Supplementary Methods for further details). CBA is widely used for culturing NTHi and other fastidious bacteria (Artman et al., 1983). Thus, we used this medium as the basis for formulating liquid chocolate medium (LCM). Formulation of LCM and growth conditions are provided in Supplementary Methods.

### RNA Extraction

All isolates were cultured in LCM and in duplicate to account for any technical replicability issues. RNA was extracted using TRIzol® as described in Supplementary Methods. After DNA digest, absence was verified using real-time PCR assays that detect H. influenzae DNA (Aziz et al., 2017) and defibrinated horse blood DNA (Humair et al., 2007). For further details see Supplementary Methods.

### Differential Expression (DE) Analysis

RNA-seq processing and DE analysis was performed as described in Supplementary Methods. Briefly, RNA-seq was carried out using the Illumina HiSeq 2500 platform. In total, 24 isolates were sequenced in duplicate, generating 48 transcriptomes. To identify molecular signatures of lung adaptation on a within-patient basis, DE analysis was first conducted by comparing BAL with NP isolates from individual patients. Subsequently, convergence analysis was performed by comparing BAL with NP isolates from all 12 isolate pairs. The following additional analyses with different sample sets were performed to ensure robustness: (a) those NP-BAL pairs with DE detected in the within-patient analysis (eight in total), and (b) the three NP-BAL pairs with > 45 DE genes (60051, 60068, and 60295). For all convergence analyses, the two NP-BAL pairs from patient 60373 were treated either as independent NP-BAL pairs or grouped as a single patient. DE was conducted using edgeR (v3.18.1) (Robinson et al., 2010), implemented in R (v3.4.1) (R Core Team, 2014). For all analyses, DE was determined using the quasi-likelihood method (glmQLFit function).
Genes with significant DE were annotated using Clusters of Orthologous Groups (COG) (Galperin et al., 2015), with categories retrieved from a previous study (Santana et al., 2014) with minor corrections (Supplementary Table S1). For genes with multiple COG categories, all assigned categories were included in the analysis. To assess for significant enrichment of functional groups, COG categories of the reference genome 86-028NP were compared with those identified in the within-patient DE analyses. Statistical assessment comparing the proportion of categories was performed in R using a two-tailed Fisher's exact test with a false discovery rate corrected p-value of ≤ 0.05.

RESULTS

Comparative Genomic Analysis of Paired Isolates

To assess diversity and to identify potential clustering of NP or BAL isolates, a maximum parsimony phylogenetic tree was constructed using 124,262 biallelic, core-genome SNPs identified among the paired isolates and a global NTHi isolate set comprising 157 strains (Supplementary Figure S1). Phylogenetic analysis confirmed that pairs from individual patients clustered closely together yet were distinct from other NP-BAL pairs. Consistent with previous studies (De Chiara et al., 2014; Price et al., 2015; Pettigrew et al., 2018), there was no evidence of NP- or BAL-specific clades. Isolate pairs were identical sequence types (STs), with 5 of the 12 pairs possessing novel STs (ST-1906 to ST-1910; Supplementary Table S2). Patient 60373 had two distinct isolate pairs (NP-BAL pairs 60373_P1 and 60373_P2) that were separated by >17,000 SNPs, demonstrating the presence of at least two NTHi lineages colonizing this patient. This finding was also reflected in the MLST data, with 60373_P1 isolates belonging to ST-1910 whereas 60373_P2 isolates were ST-139 (Supplementary Table S2). Antibiotics are routinely used to treat exacerbation events in pediatric chronic lung disease, and resistance to clinically relevant antibiotics can arise during such infections. In the current work, no NTHi isolate exhibited resistance toward any of the tested blood factors (Holt, 1962), commonly used culture media for this fastidious bacterium typically includes lysed blood in a nutrient-rich agar base (e.g., CBA) (Artman et al., 1983), or HTM-supplemented artificial media after isolation, these mixtures suggest that the hsdM3 SSR undergoes a rapid mutation rate, with mutations potentially occurring during laboratory passage.

Liquid Chocolate Medium for Reproducible NTHi Growth

As NTHi has an absolute growth requirement for X and V blood factors (Holt, 1962), commonly used culture media for this fastidious bacterium typically includes lysed blood in a nutrient-rich agar base (e.g., CBA) (Artman et al., 1983), or BHI base supplemented with HTM (Coleman et al., 2003). However, we found uniform and reproducible growth of NTHi in HTM-supplemented BHI problematic; variable growth rates were observed both among isolates and between technical replicates, indicating inconsistent bacterial growth. Additionally, the microbial load was insufficient for RNA-seq, even after 24 h incubation. We also observed inconsistent OD readings between replicates and a tendency toward flocculation in certain strains. Similar issues were observed with HTM-supplemented artificial sputum medium, which is designed to mimic cystic fibrosis sputum (Fung et al., 2010; Supplementary Figure S2A).

To overcome these issues, we developed liquid chocolate medium (LCM), a medium that is straightforward to formulate, inexpensive and produces high-quality NTHi cellular density for downstream analysis. LCM yielded NTHi cultures with reproducible growth, consistent growth rates, little to no flocculation, and sufficient microbial load to support sequencing of late-log phase RNA. LCM was thus used to determine the growth curves of four isolates to identify an appropriate culture duration for RNA extraction. A 7.5-h post-inoculation time point
| Patient ID | 86-028NP locus tag | Gene  | NP to BAL genotype | Amino acid change (WT length) | WT isolate | Mutation effect in BAL isolate | Affected protein | Protein function | COG category |
|------------|------------------|-------|-------------------|-----------------------------|------------|-------------------------------|-----------------|-----------------|-------------|
| 60051      | NTHi1838         | hsdM3*| AAGACG            | Glu12fs (556)               | NP         | Loss-of-function frameshift; premature stop yields 13aa product | Type I restriction enzyme HindVIIP M protein | DNA binding; site-specific DNA-methyltransferase (adenine-specific) activity | Defense mechanisms |
| 60068      | NTHi0782         | hgpB* | GCCAACCAA         | Thr35fs (992)               | NP         | Loss-of-function frameshift; premature stop yields 12aa product | Hemoglobin-haptoglobin binding protein B | Receptor and transporter activity | Inorganic ion transport and metabolism |
| 60069      | Intergenic       | –     | GATTATT           | –                           | NP         | Intergenic (62 bp upstream of NTHi0369 [hxuC]) | –               | –               | –           |
| 60295      | NTHi0782         | hgpB  | C                 | Arg407Lys (992)             | NP         | Non-synonymous                | Hemoglobin-haptoglobin binding protein B | Receptor and transporter activity | Inorganic ion transport and metabolism |
| 60373_P1   | NTHi1838         | hsdM3*| CGAGA            | Glu12fs (556)               | NP         | Loss-of-function frameshift; premature stop yields 16aa or 24aa product | Type I restriction enzyme HindVIIP M protein | DNA binding; site-specific DNA-methyltransferase (adenine-specific) activity | Defense mechanisms |
| 65001      | NTHi0472         | lic3A*| TTTGA             | Ser18fs (320)               | NP         | Loss-of-function frameshift; premature stop yields 36aa product | CMP-Neu5Ac–lipooligosaccharide alpha 2–3 sialyltransferase | Transferase activity; transfer of glycosyl groups | Cell envelope biogenesis |
| NTHi0225   | ompP2            | C     | CAGCAGTGGG       | Ala269_Gly27 ValGlyAla (365) | NP         | In frame insertion of ValGlyAla; 368aa product | Outer membrane protein P2 | Porin activity | Cell envelope biogenesis |
| NTHi0315   | hsdS2            | A     | ACTTACAGCCGAG     | Thr173_Leu1 76del (390)     | NP         | In-frame deletion of 4aa | Type I restriction-modification system specificity protein | DNA binding and modification | Defense mechanisms |
| 65290      | NTHi1084         | bamA  | T                 | Glu508Gly (800)             | NP         | Non-synonymous                | Protective surface antigen D15 | Cell outer membrane assembly; protein insertion into membrane | Cell envelope biogenesis |

*No genetic variants were found between the 60294, 60316, 60362, 60370, and 60373_P2 pairs; only pairs with genetic differences are shown. bWild-type isolate is defined as the isolate which produces full length protein of similar length to 86-028NP reference. cMutation found in a phase-variable region. dThe BAL strain from this patient contains a mixture of two loss-of-function indel mutations in hsdM3 at a 40:60 ratio (See Figure 1). BAL, bronchoalveolar lavage; COG, cluster of orthologous groups; NP, nasopharynx; WT, wild type.
was found to represent late-log phase growth in all four cultures tested (Supplementary Figure S2B). This time point, which was chosen to reflect non-logarithmic bacterial growth rates in vivo (Yang et al., 2008; Kasi et al., 2017), was subsequently used for RNA extraction of all isolates.

Within-Patient DE and Comparison With Genetic Mutations

Culture and RNA extraction of 24 isolates was performed in duplicate. On average, 89.2% (87.1–92.0%) of sequence reads aligned uniquely to the 86-028NP reference genome, and of these, an average of 70% (68.2–74.8%) were assigned to known genomic features (Supplementary Table S2). Due to this high mapping percentage, within-patient DE analysis was performed using 86-028NP as the reference genome. This approach also greatly simplified comparisons among patients. DE was observed between eight NP-BAL pairs, ranging from 2 to 58 DE genes per pair. Three pairs (60051, 60068, and 60295) had 15 or fewer DE genes (Supplementary Dataset S1), and the remaining four pairs (60069, 60294, 60316, and 65001) did not exhibit any DE.

In total, 189 DE genes, representing a wide functional spectrum, were detected across the eight NP-BAL pairs. Eight genes were DE in three pairs (comE, comM, glgC, ligA, pilB, pilC, smf and tnaA), 39 were DE in two pairs (arcC, arcF, asnA, comABCDF, dmsC, glgABPX, glpK, hitA, hsdMS3, licBC, mao2, nagAB, nanAEK, NTHI0229, NTHI0232, NTHI0235, NTHI0646, NTHI0647, NTHI1183, NTHI1243, nrfCD, pckA, pilA, sdaA, siaT, and tnaB), and the remaining 87 were DE in single NP-BAL pairs (Figure 2 and Supplementary Dataset S1). However, only ten genes exhibited concordant downregulation (mao2, NTHI0646, NTHI0647, pckA, sdaA) or upregulation (asnA, hitA, hsdMS3, hsdS3, and tnaB) in the BAL isolates. In all other DE genes, discordance in the directionality of expression was observed, including the eight shared DE genes in three NP-BAL pairs (Figure 2). Additionally, genes in the 60295 pair were predominantly upregulated, whereas genes were predominantly downregulated in pairs 60051 and 60068. Overall, there was little overlap amongst patients.

There was also minimal correlation between the number of genetic mutations and the number of DE genes detected between NP-BAL pairs (Table 1). For example, the 60295 NP-BAL pair exhibited 46 DE genes, but only a single non-synonymous SNP (HgpBArg407Lys) was observed in the BAL isolate. In contrast, no DE was detected between the 60051 NP-BAL pair (Supplementary Dataset S1), despite three indels being identified in the BAL isolate, including a loss-of-function frameshift in the phase-variable sialyltransferase gene, lic3A, which truncates the Lic3A protein from 320 to just 36 residues (Table 2). Two NP-BAL pairs (60294 and 60316) contained no mutations and no DE genes. Conversely, two NP-BAL pairs (60051 and 60373_P1) contained both genetic and transcriptional alterations affecting a single phase-variable gene, hsdM3 (Table 2 and Supplementary Dataset S1). For both pairs, hsdM3 and hsdS3 were upregulated in the BAL isolates, with an average log2 fold change of 2.5 and 4.6, respectively.
FIGURE 2 | Heat map of differentially expressed (DE) genes identified between the NP-BAL pairs from 11 pediatric chronic lung disease patients. In total 134 non-redundant DE genes were detected across eight NP-BAL pairs. DE was observed in eight NP-BAL pairs; the remaining four pairs contained no DE. Genes were predominately upregulated in the 60295 NP-BAL pair but predominately downregulated in the 60051 and 60068 NP-BAL pairs. The direction of log₂ fold change of DE genes in BAL is color-coded as follows: upregulation (blue), downregulation (green), no DE (tan).

Additionally, *hsdR3* was upregulated by 2.2-fold in 60373_P1. However, both BAL isolates from 60051 and 60373_P1 encode truncated HsdM proteins of 13 and 16–24 residues, respectively (Figure 1C), indicating that *hsdM3* upregulation in these isolates is unlikely to have a functional consequence. Notably, the two isolates with mutations in *hgpB*, 60068 BAL Hi1 and 60295 BAL Hi1, both had a relatively high amount of DE, with 55 and 46 DE genes, respectively.

**DE Convergence and Functional Enrichment Analysis of NP-BAL Isolates**

We compared BAL with NP isolates from all patients to identify genes with convergent gene expression. However, no single convergent genetic mutations or DE genes were identified, suggesting more complex pathways involved in lung adaptation. Therefore, we conducted a functional enrichment analysis of the 134 non-redundant DE genes using COG assignments to identify enriched categories. Statistical analysis identified 4/21 COG categories that were significantly over-represented in the functional enrichment analysis (*Carbohydrate transport and metabolism, Energy production and conversion, Intracellular trafficking and secretion, and Cell motility, and secretion*), and 3/21 categories (*Translation, ribosomal structure and biogenesis, Coenzyme metabolism, and Function unknown*) that were under-represented (\(p \leq 0.05\), Figure 3). Of note, the *Function unknown* category encompasses 20.2% genes in the 86-028NP genome, yet DE was only observed in 3.5% of these genes. The basis for this under-representation may partly reflect an artificially high number of gene annotations in the NTHi genome. In support of this notion, 47/397 (11.8%) genes assigned to the *Function unknown* category exhibited zero reads across our 48 transcriptomes (Supplementary Dataset S2, gray shaded rows), representing 37.9% of all genes with zero reads.
reads. Approximately 51.8% of the other genes with zero reads corresponded with paralogous ribosomal and transfer RNA species (Supplementary Dataset S2).

The most significant over-represented COGs (p < 0.001) were Carbohydrate transport and metabolism, which is represented by 4.9% of 86-028NP genes, yet 17.3% of DE genes belonged to this category, and Cell motility and secretion, with only 0.6% genes in 86-028NP but with 5.8% of DE genes belonging to this COG. The most significant under-represented COGs were Translation, ribosomal structure and biogenesis, comprising 8.0% of 86-028NP genes but only 0.6% of the DE genes, and Function unknown, which comprises 20.2% of 86-028NP genes yet only 6.2% of DE genes belonged to this category (Figure 3 and Supplementary Dataset S1). Other significant COG categories (p < 0.01) were Energy production and conversion (5.0% of 86-028NP genes but 12.0% DE genes) and Coenzyme metabolism (4.4% 86-028NP genes but 0% DE genes).

**DISCUSSION**

Non-typeable *H. influenzae* is a pathogen of emerging public health importance (Erwin and Smith, 2007; Van Eldere et al., 2014). Conventionally considered a commensal of the upper airways, it is now recognized that NTHi is associated with increased severity and progression of multiple polymicrobial diseases of the lower airways such as COPD, bronchiectasis, and CSLD (Murphy, 2003; Purcell et al., 2014). In this study, we aimed to identify signals of NTHi pathoadaptation to the lower airways using 12 paired isogenic strains retrieved from the upper (NP) and the lower (BAL fluid) airways of 11 pediatric patients with CLSD or bronchiectasis. Previous studies have employed WGS to document the genome-wide changes leading to chronic adaptation and persistence of longitudinal NTHi isolates retrieved from COPD sputum in adults (Moleres et al., 2018; Pettigrew et al., 2018). The current work expands on these prior studies by examining both the genomic and transcriptomic profiles of NTHi adaptation in children’s airways. To ensure isolates were retrieved from the lower airways, we obtained NTHi isolates from the lower airways (BAL fluid) rather than sputum, the latter of which is fraught with upper respiratory tract contamination issues (Gronseth et al., 2017).

As part of our study criteria, only isogenic strains were selected to simplify the identification of convergent pathoadaptive mechanisms. As expected, comparative genomic analyses found minimal genetic differences between all isogenic NP-BAL pairs, with only nine total variants in our dataset, ranging from zero to three variants per pair. Five pairs were genetically identical, with the remaining seven encoding a total of two SNPs and seven indels. Notably, four of these indels occurred in tandem repeat regions, or SSR tracts, of phase-variable genes, which are abundant in the NTHi genome (Gilsdorf et al., 2004; Pettigrew et al., 2018). SSRs provide a rapid mechanism for gene expression modulation and adaptation (Vasu and Nagaraja, 2013; De Ste Croix et al., 2017). Two SSR indels occurred in the hypervariable Type I restriction-modification (RM) system methyltransferase gene, *hsdM3*. The remaining SSR indels occurred in the hemoglobin-haptoglobin binding protein B-encoding gene *hgpB*, which is involved in NP colonization and iron acquisition (Poole et al., 2013), and *lic3A*, which encodes a lipopolysaccharide (LOS) sialyltransferase that facilitates NTHi colonization in different anatomical sites (Hosking et al., 1999; Hood et al., 2001).

Our genetic findings concur with the recent longitudinal studies, which found that phase variation in *lic3A*, *hsdM3*, and *hgpB* are common in NTHi isolates retrieved from COPD airways (Moleres et al., 2018; Pettigrew et al., 2018). In addition to the *hgpB* SSR variant in 60068 BAL H1 variants that truncates the protein from 992 to 12 residues, a non-synonymous SNP (HgpB<sub>Arg407Lys</sub>) was seen in 60295 BAL H1. Likewise, the SSR region of *hsdM3* was deleteriously mutated in the 60051 and 60373_P1 BAL isolates, yielding truncated proteins of ≤ 24 residues (Figure 1C). The *hsdM3* gene is part of a Type I RM system that acts as a defense system against foreign DNA invasion by, e.g., bacteriophages (Murray, 2000), fine-tuning of competence through variable methylation patterns, and in the methylation of self-DNA that may alter gene expression (Vasu and Nagaraja, 2013). While the *hsdM3* mutations in the pediatric BAL isolates would render their corresponding Type I RM systems non-functional, the phase-variable nature of *hsdM3* enables rapid reverse switching back to an active state. Others have shown that phase-variable RM systems allow NTHi to rapidly generate genomic and phenotypic diversity, which may enhance survival and persistence, especially in new environments (Zaleski et al., 2005; Attack et al., 2015). Taken together, the commonality of SSR tract mutants affecting *hgpB*, *hsdM3*, and *lic3A* in airway-derived isolates demonstrates that phase variation of these loci is driven by strong diversifying selection, with this variation being key in the long-term persistence of NTHi within the hostile lung environment.

Outer membrane porin (OmpP1 and OmpP2) mutations are also common in COPD-derived NTHi isolates. Previous studies have identified null *ompP1* mutations in ~33% of COPD isolates, and in-frame *ompP2* mutations (i.e., missense SNPs and indels) in ~20% of COPD isolates (Moleres et al., 2018; Pettigrew et al., 2018). Similarly, we identified an in-frame *ompP2* mutation in the 65001 BAL H1 isolate that occurred in the extracellular loop six motif (Sikkema and Murphy, 1992) and increased OmpP2 length by three amino acids. However, we did not observe any genomic or transcriptomic *ompP1* variation between our 12 NP and BAL pairs. Although our study is cross-sectional in nature and our sample size is modest, this result suggests that *ompP1* variants may be much less common in pediatric NTHi lung-adapted strains than COPD lung-adapted strains. Biochemical profiles in COPD airways (e.g., fatty acids with bactericidal activity against NTHi wild-type strains) (Moleres et al., 2018) are potentially different to those present in pediatric bronchiectasis or CSLD airways, which may explain the greater selective pressure for OmpP1 inactivation in COPD airways. Further work is needed to document the prevalence of *ompP1* mutations in a larger pediatric lung isolate cohort to examine whether diseased pediatric airways do in fact impart different OmpP1 pressures on NTHi compared with isolates from COPD airways.
Across the 12 within-patient DE analyses, a total of 134 non-redundant genes (range: 0–58) were DE between NP-BAL pairs, with no correlation between the number of mutations and the number of DE genes. Indeed, NP-BAL pairs 60362, 60370 and 60373_P2 were devoid of mutations, yet DE was observed in 15, 2, and 4 genes, respectively. Likewise, 65001 BAL Hi1 encoded mutations in lic3A, ompP2, and a Type I RM system specificity gene, hsdS2, yet no DE was detected in this strain. This lack of correlation suggests that DE between isolate pairs is influenced by epigenetic factors [e.g., altered DNA methylation (Fox et al., 2007)], or by unidentified genetic mutations due to inherent limitations with short-read sequencing approaches (e.g., insertion sequence-mediated rearrangements or variants in paralogs). Future studies employing methylation signature detection and long-read sequencing will enable a deeper understanding of the basis for DE in these strains.

We did not identify any DE gene(s) common to all isolates, indicating no universal expression profile associated with lung adaptation (i.e., the BAL isolates). However, some degree of gene convergence was observed, with eight DE genes common to three isolate pairs, and 39 common to two pairs (Supplementary Dataset S1). Of these, the com and pil operons, required for the biogenesis and function of the type IV pilus, were DE in three pairs. The type IV pilus is involved in multiple biological roles including adherence to epithelial cells, competence, twitching motility, biofilm formation and long-term NP colonization (Bakaletz et al., 2005; Jurcisek et al., 2007). The pathway responsible for uptake, transport and incorporation of N-Acetylneuraminic acid ( Neu5Ac) into LOS (Apicella, 2012; Wong and Akerley, 2012), was DE in two pairs. Incorporation of Neu5Ac into the LOS enhances immune evasion and increases resistance to killing by human serum (Mandrell et al., 1992). Most DE genes (n = 87) occurred
in single pairs, suggesting multiple, parallel strategies toward lower airway adaptation. In addition, the directionality of DE was inconsistent, with gene upregulation in some pairs and downregulation in others (Figure 2). Lack of a convergent transcriptional signal across all strains may be due to sampling biases, for example, the inclusion of BAL isolates that have yet to adapt to the lower airways, the re-seeding of the NP environment with lung-adapted strains, or inherent differences in NP carriage time prior to lower airway infection. Greater breadth and depth of strains is needed to determine whether a convergent signal of lung adaptation can be identified in the NTHi transcriptome. Additionally, sampling of other disease sites (e.g., middle ear) may help to resolve NTHi adaptation strategies in the lung environment.

To identify parallel signatures of adaptation to the lung environment on a broader scale, we categorized the functions of within-patient DE genes based on COG categories. Overall, strains isolated from the lower airways showed significant enrichment of DE genes in 7/21 COG categories. Four categories were over-represented in the DE gene dataset when compared with their prevalence in the 86-028NP genome, with Carbohydrate transport and metabolism, Cell motility and secretion, Intracellular trafficking and secretion, and Energy production and conversion being significant (p < 0.05 or p < 0.001). In comparison, the COG categories Translation, ribosomal structure and biogenesis, Function unknown and Coenzyme metabolism were significantly under-represented (p < 0.01 or p < 0.001) in the DE gene dataset, suggesting that altered expression of these gene categories is negatively associated with lung adaptation. However, analysis of the 48 transcriptomes revealed that 11.8% genes in the Function unknown category had zero reads mapping to the 86-028NP genome (Supplementary Dataset S2, gray-shaded rows), suggesting that many of these so-called genes likely do not produce messenger RNA or proteins under any growth conditions, and may represent incorrect gene annotations. Further RNA-seq studies of NTHi cultures grown under different conditions will help to resolve these potential annotation issues. Another noteworthy observation was that Carbohydrate transport and metabolism and Energy production and conversion collectively contained ~35% of all DE genes and were primarily downregulated. This observation is consistent with Baddal et al. (2015), who observed the downregulation of genes involved in NTHi central metabolism in response to environmental stimuli. Intracellular trafficking and secretion and Cell motility and secretion were also over-represented in the BAL isolates and predominately downregulated (Figure 3 and Supplementary Dataset S1). Taken together, these results suggest that NTHi alters certain functional pathways in response to the lower airway environment, suggesting a degree of predictability in adapting to this new niche. Although further work is needed to consolidate these findings, our work sets the stage for identifying key functional pathways that may be exploited for targeted treatment and eradication of this pathogen.

Our study chose not to assess the wider polymicrobial community of the lower airways, as accurate transcriptional characterization of entire bacterial communities remains bioinformatically challenging (Lim et al., 2013). We instead chose to focus on an in-depth genomic and transcriptional analysis of NTHi, a species known to play a crucial role in the progression and severity of lung disease (Erwin and Smith, 2007; Van Eldere et al., 2014; Slack, 2017). Genetically diverse NTHi isolates from different patients were selected to enable identification of convergent adaptation strategies to the lung environment of pediatric patients. Due to the necessary passaging of strains to ensure purity and to obtain sufficient RNA material for sequencing, we cannot discount that these procedures may have influenced the NTHi expression profiles so that they no longer represent their in vivo counterparts. However, as previously demonstrated with other bacterial species, isolates appear to maintain their expression profiles even after microbiological handling and culturing in vitro (Viberg et al., 2017). For future studies, we recommend maximizing the number of strains isolated from individual patients together with longitudinal sampling to better understand pathogen adaptation over time.

In conclusion, our study provides new insights into NTHi adaptation to the lower airways in pediatric chronic lung diseases. Through genomic and transcriptomic characterization of 12 paired NTHi isolates from the NP and the lung, we found that NTHi employs several avenues of pathoadaptation that enables this pathogen to persist in the lower airways. Although we did not identify evidence of convergent pathoadaptive mechanisms at the single-gene level, our study identified parallel DE at the functional level, suggesting that NTHi adaptation to the pediatric airways is a complex but ultimately predictable process. The next steps require characterizing larger isolate panels using genomic and transcriptomic methods, particularly the elucidation of within-host NTHi population diversity from NP and lower airway sites across multiple patients, to better understand the pathoadaptive mechanisms that enable NTHi to persist and cause disease. Such findings will be crucial for the informed development of effective therapeutic interventions to prevent NTHi-driven chronic lung diseases.

DATA AVAILABILITY

The datasets generated for this study can be found in the NCBI database under the BioProject PRJNA484075. Genome assemblies for the 24 NTHi strains can be found in the NCBI GenBank (QWLW0000000-QWMT0000000), and Illumina WGS and RNA-seq data are available on the Sequence Read Database (SRR7719256-SRR7719303).

ETHICS STATEMENT

This study was carried out in accordance with the recommendations of the National Health and Medical Research Council’s (NHMRC) National Statement on Ethical Conduct
in Human Research (2007) with written informed consent from all subjects. All subjects gave written informed consent in accordance with the Declaration of Helsinki. The protocol was approved by the Human Research Ethics Committee of the Northern Territory Department of Health and Menzies School of Health Research, with the approval numbers HREC 07/63 and 07/85.

AUTHOR CONTRIBUTIONS

AC conducted the patient recruitment and specimen collection. EN and JB conducted the specimen processing, genotyping, and DNA extractions. AA, JB, HS-V, EP, and TH conducted the isolate selection. AA conducted the culturing, RNA extraction, and bioinformatics analysis with supervision and assistance from DS, EP, and TH. AA wrote the initial manuscript draft. DS, EP, and TH critically reviewed and edited the manuscript. DS, HS-V, AC, and EP conceived of the study. DS, HS-V, and EP obtained the funding. All authors reviewed and approved the final manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2019.01622/full#supplementary-material

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.