Inactivation of the Thymidylate Synthase \(\text{thyA}\) in Non-typeable \textit{Haemophilus influenzae} Modulates Antibiotic Resistance and Has a Strong Impact on Its Interplay with the Host Airways

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Antibacterial treatment with cotrimoxazol (TxS), a combination of trimethoprim and sulfamethoxazole, generates resistance by, among others, acquisition of thymidine auxotrophy associated with mutations in the thymidylate synthase gene \(\text{thyA}\), which can modify the biology of infection. The opportunistic pathogen non-typeable \textit{Haemophilus influenzae} (NTHi) is frequently encountered in the lower airways of chronic obstructive pulmonary disease (COPD) patients, and associated with acute exacerbation of COPD symptoms. Increasing resistance of NTHi to TxS limits its suitability as initial antibacterial against COPD exacerbation, although its relationship with thymidine auxotrophy is unknown. In this study, the analysis of 2,542 NTHi isolates recovered at Bellvitge University Hospital (Spain) in the period 2010–2014 revealed 119 strains forming slow-growing colonies on the thymidine low concentration medium Mueller Hinton Fastidious, including one strain isolated from a COPD patient undergoing TxS therapy that was a reversible thymidine auxotroph. To assess the impact of thymidine auxotrophy in the NTHi-host interplay during respiratory infection, \(\text{thyA}\) mutants were generated in both the clinical isolate NTHi375 and the reference strain RdKW20. Inactivation of the \(\text{thyA}\) gene increased TxS resistance, but also promoted morphological changes consistent with elongation and impaired bacterial division, which altered \textit{H. influenzae} self-aggregation, phosphorylcholine level, C3b deposition, and airway epithelial infection patterns. Availability of external thymidine contributed to overcome such auxotrophy and TxS effect, potentially facilitated by the nucleoside transporter \(\text{nupC}\). Although, \(\text{thyA}\) inactivation resulted in bacterial attenuation in a lung infection mouse model, it
also rendered a lower clearance upon a TxS challenge in vivo. Thus, our results show that thymidine auxotrophy modulates both the NTHi host airway interplay and antibiotic resistance, which should be considered at the clinical setting for the consequences of TxS administration.

**Keywords:** *Haemophilus influenzae*, thymidylate synthase, thymidine auxotrophy, thymidine uptake, bacterial morphology, antibiotic resistance, airway infection

## INTRODUCTION

Non-typeable (non-capsulated) *Haemophilus influenzae* (NTHi) is a Gram negative coccobacillus that is a common commensal in the nasopharynx of healthy humans, and also an opportunistic pathogen causing respiratory infections such as acute otitis media, otitis media with effusion, community-acquired pneumonia, and exacerbations of chronic obstructive pulmonary disease (COPD; Agrawal and Murphy, 2011). COPD is characterized by a progressive and not fully reversible airflow limitation, accompanied by infiltration of the airways by neutrophils and mucus hypersecretion (Barnes, 2016). The chronic course and evolution of COPD is often characterized by periods of symptom exacerbation with a negative impact on the patient’s quality of life and evolution of the disease, and represent a significant cause of medical intervention and hospitalization.

Given that acute exacerbations of COPD (AECOPD) are mostly caused by bacterial and viral infections (Sethi, 2010), antibiotic therapy is routinely prescribed. Use of amoxicillin, tetracyclines, or cotrimoxazole (TxS), a combination of trimethoprim (TMP) and sulfamethoxazole (SMX), has long been regarded as standard therapy for patients with AECOPD, which has contributed to the increasing emergence of resistance to these antibiotics in common respiratory pathogens such as NTHi, and become a real challenge in the choice of adequate antibacterials (Nouira et al., 2010).

The treatment of respiratory infections has been a very important field of use of TxS, as a first-line agent against *H. influenzae* infected respiratory patients (Iyer Parameswaran and Murphy, 2009), and has also been considered as a prophylactic option in HIV-infected children therefore having an impact on *H. influenzae* carriage (Grant et al., 2009; Mwenya et al., 2010), altogether increasing TxS resistance levels in this pathogen. TxS interferes with the bacterial metabolism and replication by blocking the production of tetrahydrofolic acid (THF). THF is a co-factor for the thymidylate synthase, an essential protein encoded by the thyA gene, required for the conversion of thymidine from uracil (Stryer, 1995). During normal metabolism, dihydrofolate is reduced to THF by the dihydrofolate reductase (DHFR), encoded by the folH gene, also known as *drfA* and *folA*. TMP is a substrate analog of dihydrofolate and blocks its reduction to THF. Conversely, SMX is a substrate analog of para-aminobenzoic acid, and blocks the *folP* gene encoding the dihydropteroate synthetase DHPS, which is involved in production of dihydropteroate, a precursor of dihydrofolate. Both TMP and SMX have little toxicity to humans because humans do not synthesize folic acid but obtain it from dietary sources (Tristram et al., 2007). Inhibiting the production of THF prevents thymine synthesis and, hence, DNA replication, causing bacterial death and TxS susceptibility. However, if external thymidine is available, as shown in infected tissues (Besier et al., 2008), thymidine-dependent TxS resistance may emerge upon treatment. This aspect has been long reported (Maskell et al., 1978), and extensively analyzed in *Staphylococcus aureus*, where the underlying mechanism for thymidine dependency relies on mutations of the *thyA* gene. Thus, inactivation of the *thyA* gene generates thymidine auxotroph small colony variants (SCVs) with a strong impact in this pathogen’s physiology, virulence, and persistence, and uptake of external thymidine by the *S. aureus* NupC nucleoside transporter seems to bypass the effect of TxS (Kriegeskorte et al., 2014).

In NTHi, resistance to TxS is associated to polymorphisms and/or short insertions in the *folH* and *folP* genes, DHFR overproduction, or acquisition of the sulfonamide (SUL) genes *sul1* and *sul2* (de Groot et al., 1988, 1996; Enne et al., 2002). Existing evidence also relates *H. influenzae* TMP resistance to transient thymidine auxotrophy in isolates from sputum samples of chronic bronchitis patients receiving TMP (Platt et al., 1983). Despite this observation, little is known about thymidine-dependent antibiotic resistance and its underlying consequences for NTHi pathogenesis. Following this notion, we observed that among 2,542 NTHi strains isolated between 2010 and 2014 from clinical samples at Bellvitge University Hospital (Spain), 119 strains formed slow-growing colonies on Mueller Hinton Fastidious (MH-F) agar, a thymidine low concentration medium. This observation prompted us to hypothesize that such slow-growth could relate to thymidine auxotrophy, rendering TxS resistance due to the antibiotic administration. If so, such auxotrophy could also modify the dynamics of the NTHi-host interplay. To address these hypotheses, we screened the available NTHi isolates with a slow growth on MH-F agar and identified one thymidine auxotroph, easily reversible to the normal phenotype. Moreover, to question the relationship between thymidine auxotrophy and respiratory infection by NTHi, we employed two genome sequenced strains, NTHi strain 375, hereafter NTHi375, and *H. influenzae* (Hi) RdKW20 (Fleischmann et al., 1995; Mell et al., 2014), to generate thymidine auxotrophs by mutating the *thyA* gene, and systematically evaluated its effect on (i) bacterial resistance to TxS, morphology, growth, self-aggregation, and gene expression; (ii) NTHi-host interplay by assessing bacterial binding to a panel of components of the complement system, adhesion to- and invasion of cultured airway epithelia; (iii) NTHi respiratory infection in vivo by using a murine intranasal infection model upon TxS treatment. This work provides further evidence of the emergence of NTHi TxS resistance upon its administration, and shows for the first time...
that mutation of the thymidylate synthase encoding gene thyA in NTHi leads to a strong impact on its physiology and virulence, but also provides a survival advantage during TxS challenge. Our results provide a context for a better understanding of the potential effects of TxS treatment against NTHi respiratory infection.

MATERIALS AND METHODS

Bacterial Strains and Growth Conditions

Strains used in this study are described in Table 1. NTHi strains were grown at 37°C, 5% CO2 on chocolate agar (Biomérieux), MH-F agar (Biomerieux), or brain-heart infusion (BHI) agar supplemented with 10 µg/ml hemin and 10 µg/ml nicotinamide adenine dinucleotide (NAD), referred to as sBHI. NTHi liquid cultures were grown in sBHI (37°C, 5% CO2). Thymidine (Sigma-Aldrich) was dissolved in sterile distilled water (stock solution, 10 mg/ml). When necessary, media were supplemented with thymidine by (i) using sterile paper discs soaked on thymidine 300 µg/ml or 10 mg/ml, (ii) thymidine 300 µg/ml spreading on 20 ml chocolate agar or MH-F agar plates, (iii) thymidine 300 µg/ml addition into sBHI. Erythromycin 150 µg/ml (ErmR) or thymidine 300 µg/ml (SpecR) were grown at 37°C, supplemented with ampicillin 100 µg/ml (AmpR), erythromycin 150 µg/ml (ErmR) or SpecR, when necessary.

For thyA disruption, a DNA fragment containing the thyA gene and its respective adjacent regions (2,852 bp), was PCR amplified with Phusion polymerase (Thermofisher) using NTHi375 genomic DNA as template and primers thyA-F1 (5'-TGCCCTGAAATTCGCTCGGTATACCTTTA) and thyA-R1 (5'-CGATGATACTTAAAAAGTAATCGCGACCAGGCGG), generating pJET1.2-thyA. This cloned PCR product was disrupted by inverse PCR with Phusion polymerase, using primers thyA-F2 (5'-CGTTCTGTTGATGTCCGCTTGGA) and thyA-R2 (5'-ATATCATCAAAGCCCTCTACGTGC). An internal 226-bp fragment (nucleotides 312–537 in the thyA coding sequence) was replaced by a blunt-ended erythromycin resistance cassette excised by SmaI digestion from pBSLerm (Allen et al., 2005), generating pJET1.2-thyA. This plasmid was used as a template to amplify the thyA::ermC disruption cassette with primers thyA-F1 and thyA-R1, which was used to transform NTHi375 using the MIV method (Herriott et al., 1970). Transformants were screened by plating bacteria on sBHI agar with ErmR1 to obtain NTHi375ΔthyA. Same approach and disruption cassette were used to generate RdKW20ΔthyA. For mutant confirmation, NTHi375ΔthyA and RdKW20ΔthyA genomic DNA were used as template to be PCR amplified with four primer pairs: (i) thyA-F1 and thyA-R1, rendering a 3,814 bp product; (ii) thyA-F1 and pBSLerm-down (5'-GGTACACGAAAAACAGTTAAGGGR), rendering a 2,421 bp product; (iii) pBSLerm-up (5'-ATAAGAGGTTATATGAACGAGG) and thyA-R1, rendering a 2,190 bp product; (iv) pBSLerm-up and pBSLerm-down, rendering a 797 bp product (data not shown). Recombination events resulting from integration of the thyA disruption

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**TABLE 1** | Strains and plasmids used in this study.

| Description | Source |
|-------------|--------|
| **STRAINS** |        |
| Haemophilus influenzae | |
| NTHi375 | Wild-type, otitis media clinical isolate | Hood et al., 1999 |
| NTHi375ΔthyA | thyA::ermC, ErmR | This study |
| NTHi375ΔnupC | nupC::spec, SpecR | This study |
| RdKW20 | Laboratory strain, capsule-deficient serotype d | Fleischmann et al., 1995 |
| RdKW20ΔthyA | thyA::ermC, ErmR | This study |
| RdKW20ΔnupC | nupC::spec, SpecR | This study |
| NTHi233 | COPD isolate from a sputum sample, thymidine auxotroph | This study |
| **Escherichia coli** | |
| TOP 10 | Used for cloning assays | Thermofisher scientific |
| SW102 | Derived from DY380, it contains a defective λ prophage with the recombination proteins exo, bet, and gam being controlled by the temperature-sensitive repressor cI857 | Tracy et al., 2008 |
| **PLASMIDS** | |
| pJET1.2 | Cloning vector | Thermofisher scientific |
| pBSLerm | Source of an ErmR cassette | Allen et al., 2005 |
| pJET1.2-thyA | pJET1.2 with a 2,852 bp insert containing the thyANTHI375 gene (852 bp) and 1 kb flanking regions | This study |
| pJET1.2-thyA::ermC | pJET1.2 with a 3,814 bp insert containing a thyA::ermC disruption cassette | This study |
| pJET1.2-nupC | pJET1.2 with a 2,664 bp insert containing the nupC::spec disruption cassette | This study |
| pJET1.2-nupC::spec | pJET1.2 with a 3,501 bp insert containing a nupC::spec disruption cassette | This study |
| pRSM2832 | pKD13 derivative carrying a cassette containing a spectinomycin resistance gene flanked by FRT sites | Tracy et al., 2008 |
cassette in the NTHi375 and RdKW20 genomes were further verified by NTHi375ΔthyA and RdKW20ΔthyA genomic DNA PCR amplification with primers thyA-F1 and thyA-R1, and PCR product sequencing with primers thyA-F1, pBSLerm-up, pBSLerm-down and thyA-R1. When necessary, the thyA gene was PCR amplified with primers ThyA_Pro_F (5′-TGGCCGCTTGTAGCCGTTTG) and ThyA_R2 (5′-TCACCTAAGCTTCCGGTCTT) for DNA sequencing.

The nupC gene and its respective adjacent regions (2,664 bp) was amplified by PCR with Phusion polymerase using RdKW20 genomic DNA as template and primers nupC-F1 (5′-ATGAAACAGGTATGGAGCCAGTCCAT) and nupC-R1 (5′-GTTGAGTCAATTGGTGCAAGC). The gene-containing fragment was cloned into pET1.2, generating pET1.2-nupC. A SpecR cassette was PCR amplified from pRS2832 using gene-specific mutagenic primers nupC-F2 (5′-TGATAGAATAAGCCGAATTTTATTAACTAATCTAGGGGAAATCAATCGCTGGGATCGGACG) and nupC-R2 (5′-TACCGCAGCTTTTATTAGTTAAGATAATTAAAGTGCTGGATTTTTACTCATATCTTTAACCTTTTTTTCTAGGGGAAATCAATCGCTGGGATCGGACG), as described previously (Tracy et al., 2008). Primers were designed to delete sequences between the start codon and the last seven codons of nupC. E. coli SW102 cells were prepared for recombineering, co-electroporated with pET1.2-nupC (AmpR; 50 ng) and the nupC-specific mutagenic cassette (SpecR; 200 ng) as previously described (Sinha et al., 2012), and mutagenized clones containing pET1.2-nupC::spec were selected on LB agar with AmpR and SpecR. This plasmid was used as a template to amplify the nupC::spec disruption cassette with primers nupC-F1 and nupC-R1, which was used to transform NTHi375 and RdKW20 using the MIV method. Transformants were selected on sBHI agar with SpecR, to obtain NTHi375ΔnupC and RdKW20ΔnupC mutant strains.

Screening for NTHi Thymidine Auxotroph Clinical Isolates

Laboratory records of Microbiology Department at the Bellvitge University Hospital (Spain) regarding the difficulties of some NTHi isolates to grow on conventional antimicrobial susceptibility testing medium (Mueller-Hinton agar+5% defibrinated horse blood and 20 mg/l β-NAD, MH-F) were reviewed from 2010 to 2014. Screening for thymidine auxotrophs among NTHi clinical isolates was based on the interpretation of their growth characteristics. Isolates identified as having growth problems were further screened for thymidine auxotrophy by comparative growth on chocolate agar plates and low-thymidine MH-F agar plates. Thymidine auxotrophy was confirmed by testing strain growth on MH-F agar in the absence or presence of discs soaked with thymidine.

Susceptibility Testing under Non-standard Conditions

TxS susceptibility testing of NTHi thymidine auxotrophs failed to produce results by disc diffusion (Becton Dickinson), E-test (Biomérieux) or broth microdilution when inoculated on MH-F and incubated for 24 h, as specified by the European Committee on Antimicrobial Susceptibility Testing (EUCAST; http://www.eucast.org/clinical_breakpoints). Given that chocolate agar supported growth for all strains after 24 h of incubation, this medium was used for E-test- or TxS discs-based determination of minimal inhibitory concentrations (MIC). Strains were grown on chocolate agar, to generate bacterial suspensions normalized in phosphate-buffered saline (PBS) to OD600 = 1. Normalized suspensions were spread on chocolate agar or, when necessary, on MH-F agar, in the presence of E-test or TxS discs (23.75 mg/1.25 mg SMX:TMP) and incubated for 24 h before assessing the diameter of the growth inhibition zones.

Growth Curves

To monitor growth, NTHi strains grown on chocolate agar for 16 h were inoculated (2–5 colonies) in 20 ml sBHI, with or without thymidine, and incubated for 11 h with shaking. Cultures were diluted in 40 ml sBHI, with or without thymidine, to OD600 = 0.01 (RdKW20) or OD600 = 0.05 (NTHi375), incubated with agitation, and OD600 was recorded every hour for 8 h. Every 2 h, culture samples were serially diluted and plated on sBHI agar. Data are shown both as OD600 and c.f.u./ml. At the final time point, 10 μl of the bacterial cultures were placed on glass coverslips and fixed with 3.7% paraformaldehyde (PFA) in PBS pH 7.4 for 15 min at room temperature. Bacteria were labeled with a polyclonal rabbit anti-NTHi primary antibody diluted 1:600 and a donkey anti-rabbit conjugated to Cy2 (Jackson Immunological) secondary antibody diluted 1:100. Samples were analyzed with a Carl Zeiss Axioskop 2 plus fluorescence microscope and a Carl Zeiss Axio Cam MRC monochrome camera.

Confocal Microscopy

NTHi strains were grown on chocolate agar for 16 h, in the absence or presence of thymidine, and a colony was aseptically spread with an inoculation loop over a drop of distilled water on a microscopy slide. Samples were air-dried and stained for 15 min in the darkness with the cell-permeable fluorescent nucleic acid stain SYTO 9 (Life Technologies), following the manufacturer’s instructions. Samples were washed twice with distilled water and fluorescence was observed by confocal laser microscopy. Images were acquired using a Leica TCS-SL filter-free spectral confocal laser-scanning microscope (Leica Microsystems) equipped with a 488 nm argon laser, 543 nm and 633 nm He/Ne lasers (Centres Científics i Tecnològics-Campus de Bellvitge, Universitat de Barcelona, Spain) using a 63× magnification oil immersion objective (1.4 numerical aperture), and an image resolution of 1024 × 1024 pixels. Images were acquired randomly and analyzed using the Leica Confocal Software 2.5 (Leica Microsystems).

Transmission Electron Microscopy (TEM)

H. influenzae strains were examined by TEM after growth on chocolate agar following established procedures (Remuzgo-Martinez et al., 2015). Briefly, bacteria were applied to Formvar-coated grids, air dried, negatively stained with 1% phosphotungstic acid in distilled water for 10 s, and examined with a JEM-1011 transmission electron microscope (JEOL).
operating at 80 kV and equipped with an Orius SC1000 charge-coupled device (CCD) camera (Gatan).

**RNA Extraction and Real-Time Quantitative PCR (RT-qPCR) Analysis**

NTHi strains were grown for 16 h on chocolate agar. Bacteria (2–5 colonies) were inoculated into 20 ml sBHI, grown for 11 h, with or without thymidine, diluted into 40 ml fresh sBHI to OD<sub>600</sub> = 0.05, in the absence or presence of thymidine, and grown to OD<sub>600</sub> = 0.6. Bacterial total RNA was isolated using TRIzol reagent (Invitrogen). Total RNA quality was evaluated using RNA 6000 Nano LabChips (Agilent 2100 Bioanalyzer). All samples had intact 16S and 23S ribosomal RNA. Complementary DNA (cDNA) was synthesized from total RNA (1 µg) using SuperScript II Reverse Transcriptase reagents (Invitrogen). Real-time quantitative PCR was performed using Thermo Scientific Luminaris HiGreen qPCR Master Mix (Thermo Scientific) and fluorescence data was analyzed with BioRad CFX96 qPCR System (Bio-Rad). Relative quantities of mRNAs were calculated using the comparative threshold cycle (Ct) method and normalized using 16S ribosomal RNA (16SrRNA) as an endogenous control. Primer pairs were designed with Primer3 software: for nupC, nupC-RT-Fw (5′-ATTGTATCGACTCTAG), nupC-RT-Rv (5′-ATTGTCCTGCAATACTGCT); for tdk, tdk-RT-Fw (5′-TGTGTGTTTATGCTTGTTT) and tdk-RT-Rv (5′-CTTTAAATGACCTGCCTGTA); for 16S, 16S-Fw (5′-GGGCTTGTAGCCGGTGAAC) and 16S-Rv (5′-GCCAGTAATAATCGCCCTTCTTCTAG). Data are expressed as relative expression on mutant strains compared to their parental wild-type strains, considered to be 1. All measures were carried out in duplicate and at least three times (n ≥ 6).

**Bacterial Aggregation Assay**

Three to four colonies of NTHi grown on chocolate agar for 16 h were inoculated into 20 ml sBHI, grown for 11 h, diluted in sBHI to OD<sub>600</sub> = 1 and left standing at room temperature for 5 h (starting volume ~25 ml). The viability of each culture was tested by serial dilution and plating on sBHI agar at the beginning of each experiment (t = 0; data not shown). OD<sub>600</sub> readings were performed at regular time intervals on 700 µl aliquots collected from the top of each bacterial suspension. At least independent experiments (n ≥ 4) were performed for each strain.

**Phosphorylcholine (PCho) Quantification**

Three to four colonies of NTHi grown on chocolate agar for 16 h were inoculated into 20 ml sBHI, grown for 11 h, diluted into 40 ml sBHI to OD<sub>600</sub> = 0.05, grown to OD<sub>600</sub> = 0.6 in the absence or presence of thymidine, serially diluted, plated on sBHI agar for c.f.u. determination, and used to generate stocks stored at −80°C in sBHI with 20% glycerol as single use aliquots for further experiments. For PCho determination, ~1 × 10<sup>7</sup> c.f.u. were incubated for 1 h at 37°C with TEPC-15, a mouse monoclonal antibody specific for PCho (Sigma-Aldrich) diluted 1:25 in PBS-0.05% Tween 20. Samples were washed twice with PBS-0.05% Tween 20, and incubated with a fluorescein isothiocyanate (FITC)-conjugated rabbit anti-mouse (Serotec) diluted 1:300 in PBS-0.05% Tween 20 for 30 min at 4°C under dark conditions. Bacteria were washed with PBS-0.05% Tween 20, fixed in 3% paraformaldehyde (PFA) for 2–3 min at room temperature, and analyzed on a FACSCalibur flow cytometer (BD Biosciences) using forward and side scatter parameters to gate on at least 25,000 bacteria. Results are expressed as a relative percent fluorescence index (RFI), to measure both the proportion of fluorescent bacteria positive for PCho and the intensity of fluorescence (Ramos-Sevillano et al., 2015). Assays were performed in quadruplicate in at least three independent occasions (n ≥ 12).

**Binding of Complement Factors to NTHi**

Bacterial strains were grown as indicated for PCho determination. C3b deposition was analyzed as explained previously (Ramos-Sevillano et al., 2015). Briefly, a bacterial suspension containing ~1 × 10<sup>7</sup> c.f.u. was opsonized with human serum diluted 1:4 in PBS-0.05% Tween 20, and detected with a FITC-conjugated polyclonal goat anti-human C3b antibody (ICN-Cappel) diluted 1:300 in PBS-0.05% Tween 20 for 30 min at 4°C under dark conditions. CRP binding was measured as previously described (Ramos-Sevillano et al., 2015), by incubating ~1 × 10<sup>7</sup> c.f.u. with human serum diluted 1:4 in PBS-0.05% Tween 20 and detected with a polyclonal rabbit anti-human CRP antibody (Calbiochem) for 1 h at 37°C, followed by two washes with PBS-0.05% Tween 20. Bacterial suspensions were then incubated for 30 min with a FITC-conjugated polyclonal goat anti-rabbit antibody in PBS-0.05% Tween 20 for 1 h at 37°C. For both C3b and CRP binding, bacteria were finally washed with PBS-0.05% Tween 20, fixed in 3% PFA, and analyzed on a FACSCalibur flow cytometer as described above. Results are expressed as a RFI. Assays were performed in quadruplicate in at least three independent occasions (n ≥ 12).

**Cell Culture and Bacterial Infection**

A549 human alveolar basal epithelial cells (ATCC CCL-185) were maintained as described (Morey et al., 2011), seeded to 6 × 10<sup>4</sup> cells/well for 32 h, and serum-starved 16 h before infection. NCI-H-292 mucoepidermoid pulmonary human carcinoma epithelial cells (ATCC CRL-1848) were maintained as described (Euba et al., 2015c), and seeded to 4 × 10<sup>5</sup> cells/well 16 h before infection. Adhesion and invasion assays were performed and processed as described (Morey et al., 2011; Lopez-Gomez et al., 2012; Euba et al., 2015a,c). For infection, PBS-normalized bacterial suspensions (OD<sub>600</sub> = 1) were prepared by using NTHi strains grown on chocolate agar for 16 h, or in 20 ml sBHI for 11 h in the absence or presence of thymidine. A multiplicity of infection (MOI) of ~100:1 was used. To monitor adhesion, cells were infected for 30 min. Although this assay does not completely exclude a possible internalization of some bacteria, experimental conditions were previously set to mainly monitor adhesion (Morey et al., 2011; data not shown). For invasion assays, cells were incubated with bacteria for 2 h, washed three times with PBS, incubated for 1 h with RPMI 1640 medium containing 10% FCS, Heps 10 mM, and gentamicin 200 µg/ml to kill extracellular bacteria.
When necessary, A549 cells were infected for 30 min with bacterial suspensions (OD₆₀₀ = 1) generated with bacteria grown on chocolate agar, in the absence or presence of thymidine; infections were performed in RPMI 1640, and in the absence or presence of purified C3 (16 µg/ml), 2% C3-deficient human serum (HS), or C3-depleted HS reconstituted with purified C3. Human C3-deficient serum and human complement C3 were purchased from Sigma-Aldrich. Cells were then washed three times with PBS, lysed with 300 µl of PBS-saponin 0.025% for 10 min at room temperature, and serial dilutions were plated on sBHI agar. All infections were performed in triplicate at least three independent times (n ≥ 9). Results are expressed as c.f.u./ml.

Secretion of IL-8
For A549 cell stimulation by NTHi, bacteria grown on chocolate agar, in the absence or presence of thymidine, were collected with PBS, suspensions were normalized to OD₆₀₀ = 1, and used for a 2 h infection with a MOI of ~100:1. Cells were washed three times with PBS, and incubated for 6 h in RPMI 1640 medium containing 10% FCS, Hepes 10 mM and gentamicin 100 µg/ml. Supernatants were collected from the wells, cells debris removed by centrifugation and samples frozen at −80°C. IL-8 levels in the supernatants were measured by ELISA (Abnova KA0115) with sensitivity <2 pg/ml. Infections were performed in duplicate and at least twice (n ≥ 4). Results are expressed as IL-8 pg/ml.

Scanning Electron Microscopy (SEM)
A549 cells were seeded on glass coverslips and infected as described above, by using bacteria previously grown on chocolate agar. Coverslips were fixed in ice-cold 3% glutaraldehyde for 20 min at 4°C. Samples were dehydrated in a series of graded acetone, dried by the critical point method, coated with gold in a Fine coat ion sputter JFC-1100 226 (JEOL, Ltd) and observed with an Inspect S microscope (FEI Company) working at 15 or 20 kV (Lazaro-Diez et al., 2016).

NTHi Mouse Lung Infection
A CD1 mouse model of NTHi lung infection was used, as described previously (Morey et al., 2013; Euba et al., 2015a,b,c). CD1 female mice (18–20 g) aged 4–5 weeks were purchased from Charles River Laboratories (France), housed under pathogen-free conditions at the Institute of Agrobiotechnology facilities (registration number ES/31-2016-000002-CR-SU-US), and used at 22–25 g. Animal handling and procedures were in accordance with the current European (Directive 86/609/EEC) and National (Real Decreto 53/2013) legislations, following the FELASA and ARRIVE guidelines, and with the approval of the Universidad Pública de Navarra (UPNa) Animal Experimentation Committee (Comité de Ética, Experimentación Animal y Bioseguridad) and the local Government authorization. NTHi375 and NTHi375ΔthyA were used for lung infection, and mice were randomly divided into two groups (n = 5): (i) control, vehicle solution (0.1 ml PBS) administered by orosophageal gavage (Popper & Sons Inc.); (ii) one dose of TxS each 6 h administered by orosophageal gavage, starting at 6 h post-infection (hpi). TxS (Septrin, 8 mg/40 mg/ml TMP-SMX) treatment was performed at a dose of 960 mg/kg of body weight in 0.1 ml PBS. Infecting bacteria were previously grown on chocolate agar, in the absence or presence of thymidine. For NTHi intranasal infection, 40 µl of a NTHi suspension containing ~5×10⁸ c.f.u./ml (~2×10⁷ c.f.u./mouse) were placed at the entrance of the nostrils until complete inhalation, in mice previously anesthetized with ketamine-xylazine (3:1). At 12, 24, or 48 hpi, mice were euthanized using cervical dislocation. BALF samples were obtained by perfusion and collection of 0.7 ml of PBS, with help of a sterile 20G (1.1 mm diameter) Vialon™ intravenous catheter (Becton-Dickinson) inserted into the trachea. An aliquot of each recovered BALF was serially 10-fold diluted in PBS, and plated on sBHI agar to determine the number of viable bacteria. Results are expressed as mean ± SD of individual log₁₀ c.f.u./BALF. In parallel, lungs were removed; the left one was processed for viable bacterial counts (as described above), and the right lung was fixed in 10% neutral buffered formalin for histological purposes. Heads and necks containing upper airways, larynx, and tracheas were fixed in the same buffered formalin for histology. Uninfected mice receiving PBS or TxS were used as controls when necessary.

For cell counting, the remaining volume of each BALF sample was centrifuged at 5,000 r.p.m. for 3 min at 4°C. Each pellet was resuspended in 1 ml RPMI 1640 with 10% FCS and Hepes 10 mM, and total cell count determined using a hemocytometer. ~5×10⁴ cells in 200 µl RPMI 1640 with 10% FCS and Hepes 10 mM were used for cytokin preparation (1,500 r.p.m. for 10 min at room temperature, Thermo Shandon Cytospin). Giemsa stains were performed with an automated hematology slide preparation unit (SP-10, Sysmex Cosporation) according to the manufacturer’s instructions. Preparations were examined in a double-blinded manner with an optical microscope (BX, Olympus).

Histopathology and Lesion Score
Heads and necks were rinsed in running tap water for 1 h, immersed in 5% nitric acid for 24–36 h until complete decalcification, and 7–8 transaxial slices were made every 3–4 mm beginning at the nostrils and finishing in the caudal tracheas. Transaxial slices and lungs were embedded in paraffin, and 4–6 µm sections were stained with hematoxylin and eosin (H&E) by standard procedures, and examined by microscopy to determine the presence and extent of inflammatory lesions. Sections were examined blind as sets by a trained veterinary pathologist (Dr. M. Barberán). Parameters characterizing an acute inflammatory reaction in upper airways, larynx, trachea, and lung, including hemorrhages, hyperemia, polymorphonuclear cell infiltrates (PMNs), and alveolar macrophages, were subjectively scored on a scale of 0–3 (0: absent, 1: mild, 2: moderate, 3: severe). For tissue control, similar organs obtained from non-infected control and TxS treated mice were processed in an identical manner to the infected tissues.

Statistical Analysis
For self-aggregation, gene expression, diameter of growth or inhibition zones, PCho level, complement deposition, cell
infection, bacterial loads in lungs and BALF samples, and inflammatory cells in BALF samples, mean ± SD were calculated and statistical comparison of means performed using the two-tail t-test. For histopathology scoring, means ± SD were also calculated and statistical comparisons performed using one-way analysis of variance (ANOVA) followed by Fisher’s protected least significant difference (PLSD) multiple-comparison test. In all cases, a p < 0.05 value was considered statistically significant. Analyzes were performed using Prism software, version 6 for Mac (GraphPad Software) statistical package.

RESULTS
Isolation of a Thymidine Dependent NTHi Strain from the Sputum of a COPD Patient Receiving TxS

Among 2,542 clinical NTHi isolated between 2010 and 2014 at Bellvitge University Hospital (Spain), 119 (4.7%) isolates formed slow-growing colonies on MH-F agar. Thymidine auxotrophy was screened on the 119 isolates by comparative growth on chocolate agar and on low-thymidine MH-F agar. Thymidine auxotrophy was confirmed by absence of bacterial growth on MH-F agar, and further growth on MH-F agar in the presence of discs soaked in thymidine. One thymidine auxotroph NTHi isolate, named strain 8233, was identified in a sputum sample recovered from a COPD patient who had received TxS for 36 days to treat an acute exacerbation by Stenotrophomonas maltophilia. NTHi8233 produced small colonies on chocolate agar (data not shown) and did not grow on MH-F agar, except when discs soaked with thymidine 300 µg/ml or 10 µg/ml were added to the plate. Bacterial growth showed to be dependent on thymidine concentration (Figure 1A). The thymidylate synthase encoding gene thyA

Generation and Characterization of Thymidine Dependent H. influenzae strains

H. influenzae genome sequenced strains NTHi375 and RdKW20 were employed to generate thymidine dependent mutants by disruption of the thyA gene (thyA accession numbers NF38_0045 and HI0905, respectively). At the protein level, ThyANTHI375 and ThyARdKW20 displayed 98.2% identity (Figure S1E). NTHi mutants lacking the thyA gene were selected on sBHI agar and rendered normal size colonies (data not shown), did not grow on MH-F agar, and grew around discs soaked with thymidine on MH-F agar. Growth was dependent on the thymidine concentration present on the discs (Figure 1B and Table 2). Thus, the thyA mutants displayed a larger growth zone around discs soaked with thymidine 10 mg/ml than around discs soaked with thymidine 300 µg/ml (NTHi375ΔthyA, p < 0.005; RdKW20ΔthyA, p < 0.05). In contrast, wild-type (WT) bacteria presented a normal growth on MH-F agar. For NTHi375 WT and ΔthyA strains, the MICs of TxS were 0.25 and 0.38–0.5 µg/ml, respectively. For RdKW20 WT and thyA mutant strains, the MICs of TxS were 0.12 and 0.5–0.75 µg/ml, respectively. For purpose of illustration, larger growth inhibition zones were observed for WT than for thyA mutant strains on chocolate agar around TxS discs (Figure 1B). The morphology of NTHi375ΔthyA and RdKW20ΔthyA strains was assessed by confocal microscopy, showing non-septated filaments longer than their respective WT bacteria. Bacterial median length, measured by TEM in ~70 bacteria per strain, was (i) 1.42 ± 0.56 and 3.78 ± 2.86 μm for NTHi375 WT and ΔthyA strains, respectively (p < 0.0001); (ii) 1.43 ± 0.24 and 4.6 ± 4.98 μm for RdKW20 WT and ΔthyA strains, respectively (p < 0.0001). In agreement, SEM showed enlarged thymidine dependent bacteria on the surface of infected A549 human airway epithelial cells, which may modify the infectious process, compared to that shown by the WT strains (Figure 1B).

NTHi self-aggregates, which may promote microcolony formation on host cell surfaces (Meng et al., 2011; Mell et al., 2016). We asked whether the observed thymidine auxotrophy-driven increased bacterial size differs in its self-aggregation, by using tube-settling assays and monitoring the optical density of bacterial suspensions over time. Thymidine dependent mutants self-aggregated faster than their respective WT strains (Figure 1C).

Together, inactivation of the thyA gene in NTHi (i) impaired bacterial growth on MH-F agar, compensated by addition of the 6 nt duplication described above, even though its MIC of TxS remained unchanged.

Identification of NTHi8233 supports the notion that thymidine auxotrophy may be a bacterial response to TxS treatment that could be underestimated due to the lack of growth on MH-F agar and given that, as previously stated for other NTHi isolates (Platt et al., 1983), was shown to be reversible. This observation prompted us to generate mutant strains lacking the thyA gene on previously characterized NTHi genetic backgrounds for a detailed study of the impact of thymidine auxotrophy for this pathogen.
external thymidine, (ii) triggered enlarged bacilli which self-aggregate faster than their isogenic WT strains, (iii) increased resistance to TxS.

**Thymidine Auxotrophy Causes Growth Defects in *H. influenzae***

Growth in sBHI was analyzed for WT and *thyA* mutant strains. NTHi375Δ*thyA* had an extended lag phase and lower final OD$_{600}$ than the WT strain, which correlated with a reduced viability, measured as c.f.u./ml at the indicated time points (**Figure 2A**). Although not as pronounced, RdKW20Δ*thyA* also had a slightly extended lag phase and lower numbers of viable bacteria than those of the WT strain (**Figure 2B**). In both *thyA* mutants, the observed growth defects were partially restored by sBHI supplementation with thymidine 300 µg/ml, independently of its addition in the pre-culture used for further dilution in sBHI and
**TABLE 2 |** Bacterial growth in different media and conditions.

| H. influenzae strain | Chocolate agar | MH-F agar | Diameter of bacterial growth on MH-F agar around Thy\(^{a}\) disc (cm) |
|----------------------|----------------|-----------|--------------------------------------------------|
| NTHi375              | Yes            | Yes       | Thy 300 µg/ml                                   |
| NTHi375ΔthyA         | Yes            | No\(^{b}\) | 0.65 ± 0.05                                    |
| NTHi375ΔthyA+Thy\(^{c}\) | Yes     | No        | 1.25 ± 0.05                                    |
| RdKW20ΔthyA         | Yes            | Yes       | 1.83 ± 0.08                                    |
| RdKW20ΔthyA+Thy     | Yes            | No        | 0.9 ± 0.1                                      |
| RdKW20ΔthyA        | Yes            | Yes       | 1.4 ± 0.1                                     |

\(^{a}\)Thy, thymidine.

\(^{b}\)Total absence of bacterial growth.

\(^{c}\)NTHi375ΔthyA+Thy, bacteria previously grown on chocolate+Thy were used to inoculate chocolate agar or MH-F agar plates.

OD\(_{600}\) recording, or in the actual sBHI culture used for growth monitoring over time (Figures 2A,B). Immunofluorescence microscopy at the final time point of the growth curve showed non-septated long bacterial filaments for NTHi375ΔthyA grown in sBHI, but a mixture of non-septated filaments and bacteria with the average NTHi WT size for NTHi375ΔthyA pre-grown in sBHI supplemented with thymidine 300 µg/ml (Figure 2A, right). This could relate to the observed restoration of final OD\(_{600}\) but not of final bacterial counts for NTHi375ΔthyA when pre-cultured in sBHI supplemented with thymidine.

The thyA mutants, previously grown on chocolate agar with thymidine (chocolate+Thy), grew on MH-F agar around discs in a thymidine dependent manner, displaying a larger growth zone around discs soaked with thymidine 10 mg/ml than with thymidine 300 µg/ml (NTHi375ΔthyA, \(p < 0.005\); RdKW20ΔthyA, \(p < 0.005\)). Of note, thyA mutants previously grown on chocolate+Thy rendered a better growth on MH-F agar around discs soaked in thymidine than that of the same strains previously grown on chocolate agar (for thymidine 300 µg/ml, NTHi375ΔthyA, \(p < 0.05\); thyA, \(p < 0.05\); RdKW20ΔthyA, \(p < 0.005\)).

**Thymidine Auxotrophy Modifies the Expression of the Nucleoside Transporter Encoding nupC Gene in H. influenzae**

Both thyA mutants growth on MH-F agar around discs soaked in thymidine, and morphology/growth restoration in the presence of external thymidine, prompted us to speculate that these mutants may use external thymidine to overcome their nucleoside dependency. It has been previously shown that the nucleoside transporter encoding gene nupC is overexpressed in S. aureus thymidine dependent-SCVs, and that a ΔnupC mutant fails to use external thymidine for growth under TxS challenge (Chatterjee et al., 2008; Kriegeskorte et al., 2014). HI0519 and NF38_02480 are annotated as nupC in the RdKW20 and NTHi375 genomes, respectively (Fleischmann et al., 1995; Mell et al., 2014). Expression of the nupC gene in H. influenzae was assessed by RT-qPCR, showing a trend to be higher in the ΔthyA than in their isogenic WT strains grown in sBHI, with stronger evidence in NTHi375 (Figure S2A). As shown in Figure 2, sBHI supplementation with thymidine partially restored the growth defects shown by the thyA mutants. Following the notion that thymidine supplementation may restore mutant-related phenotypes, expression of the nupC gene was analyzed in ΔthyA bacteria when pre-cultured in sBHI with thymidine and then grown in sBHI, or when grown in sBHI supplemented with thymidine. Unexpectedly, when mutant bacteria were pre-cultured in sBHI supplemented with thymidine 300 µg/ml and then grown in sBHI, expression of the nupC gene was even higher than when pre-cultured and grown in sBHI (Figure S2A). Of note, a total absence of growth inhibition was observed for thyA mutant strains on chocolate agar around TxS discs, when pre-grown on chocolate+Thy (data not shown).

Next, NTHi375ΔnupC and RdKW20ΔnupC strains were generated, showing growth rates comparable to those of their respective WT strains in sBHI (Figure S2B), chocolate agar and MH-F agar (data not shown). To assess the biological function of nupC as a pyrimidine transporter responsible for the uptake of extracellular thymidine in H. influenzae, we tested the nupC mutants for their ability to use external thymidine for growth under TxS challenge in a TxS disc diffusion assay. Inhibition zones on chocolate agar and MH-F agar were comparable for WT and ΔnupC strains. Growth inhibition around TxS discs was also tested on chocolate agar and MH-F agar plates supplemented with thymidine. Unfortunately, no conclusive differences were observed because, as already described, bacterial growth on plates with high concentration of thymidine may produce haze or fine growth areas (Lorian, 1996).

Following this notion, we performed a search for pyrimidine transporters in all available genome sequenced H. influenzae strains, which did not reveal the presence of nucleoside transporters additional to NupC in this bacterial species. We also unsuccessfully attempted to generate a NTHi double mutant strain lacking both the thyA and nupC genes (data not shown), further supporting a role for NupC in the uptake of external thymidine.

Moreover, thymidylate is synthesized either by the thymidylate synthase ThyA or by the thymidine kinase TK, and inactivation of the former has been shown to result in upregulated expression of the later one in *Mycoplasma pneumoniae* (Wang et al., 2010). HI0529 and NF38_02430 are annotated as a tdk thymidine kinase in the RdKW20 and NTHi375 genomes, respectively (Fleischmann et al., 1995; Mell et al., 2014). Different to nupC, the tdk gene expression was shown to be similar in WT and ΔthyA strains grown in sBHI, independently of thymidine addition (Figure S2C).

Together, these results suggest that nupC may be a transporter for external thymidine in *H. influenzae*, whose increased expression could contribute to bypass the effects of TxS upon *de novo* thymidylate biosynthesis.
Thymidine Auxotrophy Reduces PCho Expression and C3b Deposition in H. influenzae

We have previously reported that bacteria lacking PCho self-aggregate slightly faster (Morey et al., 2013). Thymidine auxotroph mutants showed faster self-aggregation compared to that of their WT strains (Figure 1C), which could relate to changes in cellular morphology (i.e., elongation) or, alternatively, to a different amount of PCho residues on the bacterial surface. The level of PCho was measured on NTHi375 and RdKW20 WT and thyA mutant strains by flow cytometry using bacteria grown in sBHI. Thymidine auxotrophy was associated with a decreased detection of PCho in both NTHi375ΔthyA and RdKW20ΔthyA mutants, which was restored in NTHi375ΔthyA by bacterial growth in sBHI in the presence of thymidine. Indeed, PCho detection was higher in the thymidine dependent mutants grown in sBHI supplemented with thymidine than in their isogenic WT strains (Figure 3A and Figure S3A). Decreased levels of PCho in the thyA mutants may modify the bacterial ability to bind C-reactive protein (CRP; Weiser et al., 1998). However, CRP deposition on the bacterial surface was similar in WT and thyA mutant strains (Figure 3B and Figure S3B). Finally, to assess the possibility that cellular morphology may affect complement interaction, C3b deposition was analyzed. Thymidine auxotrophy was associated with decreased detection of C3b in NTHi375ΔthyA, which was restored by bacterial growth in sBHI supplemented with thymidine (Figure 3C). No differences were observed for C3b deposition between RdKW20 strains (Figure S3C).

Together, inactivation of the thyA gene caused a reduction in the amount of PCho residues on H. influenzae surface, leading to impaired C3b deposition in at least NTHi375, which suggests that...
Next, we assessed if auxotrophy-related invasion by RdKW20 WT and RdKW20 ΔthyA mutant strains was not assayed given the poor invasiveness of this strain, for both A549 cell adhesion and invasion (Figure 4A, right). Similar results were obtained for RdKW20 ΔthyA grown in sBHI, when compared to the WT strain, and to growth in sBHI with thymidine (Figure S4A). Given the impaired host cell interaction due to thymidine auxotrophy, we asked if inactivation of the thyA gene modifies the epithelial cell inflammatory response upon NTHi infection. Of note, the amount of secreted IL-8 was higher in A549 cells infected by NTHi375ΔthyA than by the WT strain, and such increase was reduced to WT levels upon infection by NTHi375ΔthyA previously grown on chocolate+Thy (Figure 4C). NTHi375ΔthyA lower epithelial infection rate was also observed during infection of NCI H-292 human bronchial epithelial cells (Euba et al., 2015c), for both adhesion and invasion (Figure 4B). NTHi375ΔthyA grown in sBHI showed lower adhesion to NCI H-292 cells than the WT strain, which was restored by thymidine supplementation, and adhesion by NTHi375ΔthyA grown in sBHI+Thy was higher than that shown by the WT strain (Figure 4B, right).

Complement opsonization, specifically C3, has been shown to enhance the bacterium-epithelial cell interaction for poorly encapsulated strains (de Astorza et al., 2004). Based on the observed differential C3b deposition by NTHi375 WT and NTHi375ΔthyA strains, we next tested the effect of C3 in NTHi infection of A549 cells. NTHi375 and NTHi375ΔthyA strains were grown on chocolate agar (ΔthyA) or chocolate+Thy (ΔthyA+Thy), and A549 cells were infected in the presence of human purified C3, C3-deficient serum, or C3-deficient serum reconstituted with human purified C3. The level of bacterial attachment did not change in the presence of C3, compared to that observed in control untreated cells (CON) or cells co-incubated with C3-deficient serum (Figure S4B).

In summary, thymidine auxotrophy modified NTHi ability to infect airway epithelial cells in terms of bacterial location and triggered inflammatory response. Changes in adhesion and IL-8 secretion could be restored by addition of external thymidine in the bacterial growth medium. The presence of C3 at the onset of infection did not modulate bacterial adhesion.

**Inactivation of the thyA Gene Attenuates H. influenzae Virulence and Confers Advantage under TxS Treatment In vivo**

Finally, we sought to determine the impact of thymidine auxotrophy in vivo, by using a mouse NTHi respiratory infection model system previously used for NTHi375 (Morey et al., 2013; Euba et al., 2015a,b). Mice were infected with NTHi375 WT and thyA mutant strains grown on chocolate agar, and bacterial loads were quantified in lungs and BALF samples at 12, 24, and 48 hpi. In lungs, NTHi375ΔthyA bacterial numbers were lower than those recovered for the WT strain at the three infection time points. The presence of C3 in the bacterial growth medium did not alter significantly bacterial challenge, as determined by thymidine dependency. In addition, the absence of thyA gene in our model system previously used for NTHi375 (Morey et al., 2013; Euba et al., 2015a,b).

**Thymidine Auxotrophy Modifies H. influenzae Interaction with Human Airway Epithelia**

The interplay of NTHi with the human respiratory epithelium plays a determinant role in the progression of infection (Clementi and Murphy, 2011). Next, we assessed if auxotrophy-related bacterial morphology changes could alter such interplay, by infecting A549 human type II pneumocytes with WT and thyA mutant strains (Morey et al., 2011; Lopez-Gomez et al., 2012; Euba et al., 2015c). Invasion by RdKW20 WT and thyA mutant strains was not assayed given the poor invasiveness of this strain, for both A549 cell adhesion and invasion (Figure 4A, right). Similar results were obtained for RdKW20 ΔthyA grown in sBHI, when compared to the WT strain, and to growth in sBHI with thymidine (Figure S4A). Given the impaired host cell interaction due to thymidine auxotrophy, we asked if inactivation of the thyA gene modifies the epithelial cell inflammatory response upon NTHi infection. Of note, the amount of secreted IL-8 was higher in A549 cells infected by NTHi375ΔthyA than by the WT strain, and such increase was reduced to WT levels upon infection by NTHi375ΔthyA previously grown on chocolate+Thy (Figure 4C). NTHi375ΔthyA lower epithelial infection rate was also observed during infection of NCI H-292 human bronchial epithelial cells (Euba et al., 2015c), for both adhesion and invasion (Figure 4B). NTHi375ΔthyA grown in sBHI showed lower adhesion to NCI H-292 cells than the WT strain, which was restored by thymidine supplementation, and adhesion by NTHi375ΔthyA grown in sBHI+Thy was higher than that shown by the WT strain (Figure 4B, right).

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FIGURE 4  

The thyA gene inactivation modulates *H. influenzae* interplay with cultured human airway epithelia. A549 type II pneumocytes (A) and NCI H-292 bronchial epithelial cells (B) were used to quantify adhesion and invasion by NTHi375 WT and ΔthyA strains. For adhesion, infecting inocula were prepared by using bacteria previously grown on chocolate agar following previously established methods (Morey et al., 2011; left). Alternatively, bacteria previously grown on sBHI, in the absence or presence of thymidine (sBHI±Thy), were used (right). NTHi375/ΔthyA showed significantly lower adhesion to A549 (chocolate agar, *p* < 0.005; sBHI, *p* < 0.005) and NCI H-292 (chocolate agar, *p* < 0.005; sBHI, *p* < 0.005) cells than the WT strain, which was restored by mutant pre-growth in sBHI with thymidine 300 µg/ml (A549, *p* < 0.05; NCI H-292, *p* < 0.005). NTHi375/ΔthyA showed significantly lower entry into A549 (*p* < 0.005) and NCI H-292 (*p* < 0.005) cells than the WT strain. (C) IL-8 release was higher in cells infected by NTHi375/ΔthyA than by the WT strain (*p* < 0.005); NTHi375/ΔthyA growth on chocolate+Thy reduced IL-8 secretion to WT levels (*p* < 0.005). *p* < 0.05, **p* < 0.005, ***p* < 0.005.

points tested. Thymidine dependency was also associated with significantly reduced bacterial counts in BALF at 12 and 24 hpi (Figures 5A,B).

To further investigate if thymidine auxotrophy, despite attenuation, confers an advantage to *H. influenzae* under TxS treatment, we used a therapeutic regimen consisting of one TxS dose (960 mg/kg, 1:5 TMP:SMX) every 6 h, starting at 6 hpi. Results indicate that TxS treatment was more efficient in reducing bacterial counts of the WT than of the thyA mutant strain. Thus, a significant reduction of the WT population in lungs at 12, 24, and 48 hpi, and BALF samples at 24 hpi, was observed under treated—compared to that of untreated conditions. In contrast, bacterial counts of NTHi375ΔthyA were unchanged at 12 and 24 hpi in both lungs and BALFs, despite TxS administration. The protective effect of thyA mutation against TxS was not observed at 48 hpi, maybe due to NTHi375ΔthyA severe clearance at this post-infection time point.

Based on the differences observed at 24 hpi, we attempted to restore the thyA mutant phenotypes by infecting with mutant bacteria previously grown in the presence of thymidine. However, intranasal inoculation of NTHi375ΔthyA grown on chocolate+Thy did not restore bacterial counts, and lung and BALF counts for NTHi375 WT were higher than those obtained for ΔthyA grown on chocolate+Thy. Unexpectedly, NTHi375ΔthyA previous growth on chocolate+Thy increased TxS efficiency, i.e., in TxS treated animals, lung, and BALF counts...
Inactivation of the thyA gene attenuates H. influenzae virulence and confers advantage under TxS treatment in vivo. CD1 mice were infected with $\sim 2 \times 10^7$ bacteria/mouse of NTHi375 WT (circle) or ΔthyA (square) strains. When necessary, a therapeutic regimen consisting of 960 mg/kg TxS (1:5 TMP:SMX) was administered orally each 6 h (dark symbols). Mice were euthanized at 12, 24, and 48 hpi, and bacterial loads were quantified in lungs (log_{10} c.f.u./lung) (A) and BALF (log_{10} c.f.u./BALF) (B). NTHi375ΔthyA showed significantly lower loads in lungs and BALFs at 12 (lung, $p < 0.001$; BALF, $p < 0.01$), 24 (lung, $p < 0.001$; BALF, $p < 0.001$), and 48 (lung, $p < 0.01$) hpi than those shown by the WT strain. TxS administration significantly reduced WT strain recovery from lungs and BALF samples at 12 (lung, $p < 0.05$), 24 (lung, $p < 0.001$; BALF, $p < 0.001$), and 48 (lung, $p < 0.001$) hpi, compared to WT infected untreated mice. Inverted triangles: CD1 mice were infected with $\sim 2 \times 10^7$ bacteria/mouse of ΔthyA strain previously grown on chocolate + Thy for 16 h; when necessary, a therapeutic regime consisting of 960 mg/kg TxS was administered orally each 6 h (dark inverted triangles). NTHi375ΔthyA previously grown on chocolate + Thy rendered significantly lower loads in lungs and BALFs than the WT strain ($p < 0.001$); NTHi375ΔthyA previously grown on chocolate + Thy showed significantly lower loads in lungs and BALFs of TxS treated—than in those of untreated mice ($p < 0.001$); in TxS treated mice, NTHi375ΔthyA previously grown on chocolate + Thy showed lower loads in lungs and BALFs than NTHi375ΔthyA previously grown on chocolate agar ($p < 0.01$). (C) BALF samples were analyzed for PMN (left) and alveolar macrophage (MΦ; right) counts at 12 hpi. Significantly higher numbers were observed in NTHi375 WT than in ΔthyA infected samples for both PMN ($p < 0.05$) and MΦs ($p < 0.05$). *$p < 0.05$, **$p < 0.01$, ***$p < 0.001$. **
were higher for NTHi375ΔthyA grown on chocolate agar than for NTHi375ΔthyA grown on chocolate+Thy. Also, lung and BALF counts for NTHi375ΔthyA grown on chocolate+Thy were higher in untreated—than in TxS treated animals (Figures 5A,B).

Microscopy score of the average histopathological lesion in samples of mice infected with WT or ΔthyA strains was also determined along the respiratory tract, and compared for untreated and TxS-treated mice (Table 3). Histopathological analysis of upper airways, larynxes, tracheas, and lungs from mice intranasally infected with WT or with ΔthyA rendered inflammatory lesions. Lungs and airways from control mice instilled with PBS did not show significant inflammation or pathological changes, independently of TxS administration. In both WT and ΔthyA-infected mice, the upper airways showed PMN infiltration of the lamina propria and lumens. PMN infiltration was found in larger numbers in the lumens of ΔthyA infected mice than in those infected with WT bacteria, at both 12 and 24 hpi. Conversely, PMN infiltration of the upper airway lamina propria was found to be larger in WT—than in ΔthyA infected mice at 24 hpi. Mild PMN infiltration of the lamina propria and lumens containing red blood cells and PMNs were the main findings in larynxes and tracheas. A tendency towards increased red blood cells containing lumens in WT-infected compared to ΔthyA infected mice was apparent at 24 hpi, although the differences did not reach statistical significance. As previously described (Euba et al., 2015b), lungs in infected mice showed areas of acute bronchopneumonia where alveolar septa were thickened with edema and hyperemic septal capillaries. Neutrophils, alveolar macrophages and scattered small hemorrhages were observed in alveolar spaces. Comparison of scored lesions at 12 hpi showed significantly more hemorrhages at the lower airway of mice infected with WT than at those infected with ΔthyA bacteria in control untreated mice. Hemorrhages were also found in larger proportions at the alveoli of mice infected with NTHi375 at 12 than at 24 hpi, but a lower proportion of alveolar macrophages was found at the lower airway of mice infected with NTHi375 at 12 than at 24 hpi. Significant differences were not found at the alveoli of ΔthyA infected mice between post-infection time points. Analysis of scored lesions in the airways of WT infected mice showed differences between control untreated- and TxS treated-mice. We observed a higher proportion of PMN infiltration in (i) the upper airway lumens at 24 hpi in TxS treated—than in control untreated mice, and (ii) the alveoli in control untreated—than in TxS treated mice at 12 hpi. Conversely, ΔthyA infection caused comparable lesions in both untreated- and TxS treated-mice (Table 3). Intranasal inoculation of NTHi375ΔthyA previously grown on chocolate+Thy rendered similar inflammatory lesions in both untreated- and TxS treated-animals; as a sole difference, a higher proportion of PMN infiltration was observed in the alveoli of TxS treated—than of control untreated mice (p < 0.05).

Last, we quantified accumulation of immune cells in the collected BALF samples at 12 hpi. NTHi375 WT pulmonary infection increased the accumulation of PMNs and alveolar macrophages, compared to that observed in the BALF of ΔthyA infected mice. TxS treatment showed a trend to reduce immune cell accumulation in BALF samples of WT infected mice. Unexpectedly, TxS treatment increased immune cell accumulation in BALF samples of ΔthyA infected mice (for both PMNs and alveolar macrophages; Figure 5C).

In summary, thymidine auxotrophy reduces NTHi virulence but, concomitantly, it confers an advantage under TxS treatment in vivo, suggesting that uptake of external thymidine from the infected tissue may contribute to bypass the bactericidal effect of TxS. Moreover, overall higher inflammatory traits were observed in WT—than in ΔthyA infected mice.

**DISCUSSION**

First-line antimicrobial agents must be effective, reliable, widely available and affordable in resource-poor settings (Grant et al., 2009). Following this notion, oral TxS has been recommended for years as initial antibacterial for acute otitis media, non-severe pneumonia or AECOPD because of its effectiveness and reasonable price, which, in turn, has resulted in increased resistance patterns by frequent respiratory pathogens such as *H. influenzae*. Thus, *H. influenzae* increased TxS resistance has been reported in clinical isolates from serotypes b, e, and f, and non-typeable strains from various pathological origins (Rowe et al., 2000; Leiberman et al., 2001; Campos et al., 2003a,b; Arguedas et al., 2005; Mohd-Zain et al., 2012; Puig et al., 2014; Greenhill et al., 2015). Besides *H. influenzae* TxS resistance due to changes in the sequence and/or expression of the folH and folP genes, or acquisition of the sul genes (de Groot et al., 1988, 1996; Enne et al., 2002), TMP resistance has been reported to arise in this pathogen as an indirect result of mutation to thymidine/thymine auxotrophy (Platt et al., 1983). Standard procedures for determination of *H. influenzae* antibiotic susceptibility at diagnostic laboratories involve the use of the low thymidine containing medium MH-F, which could underestimate the frequency of thymidine-dependent TMP/TxS resistance. Indeed, the routine evaluation of NTHi clinical isolates carried out in this study reported a proportion of strains designated as forming slow-growing colonies, which prompted us to screen their thymidine dependency, leading to identification of the thymidine auxotroph NTHi8233. Compared to ThyArΔkw230, ThyANTH8233 has a two-amino acid insertion at positions 90 and 91, which slightly modifies its predicted structure (Figure S1). Inactive thyA gene alleles have been reported for *S. aureus* due to in-frame deletions, deletions resulting in a frameshift, or point mutations resulting in amino acid transitions or non-sense mutations (Chatterjee et al., 2008; Kriegeskorte et al., 2014). In this study, we identified a NTHi thyA allele with an in-frame insertion compared to previously genome sequenced strains, which could alter its enzymatic activity. The ThyA canonical active site tryptophan (W83) proposed for bacterial sequences (Baugh et al., 2013) is conserved in ThyANTH8233, and future work will attempt to purify ThyANTH8233 and further assess its thymidylate synthase activity.

Moreover, NTHi8233 thymidine auxotrophy was found to be reversible, and such reversion was associated to disappearance
## TABLE 3 | Score of histopathological lesions found in the airways of control untreated or TxS treated mice, intranasally infected with NTHi375 WT or ΔthyA strains.

| Strain          | Treatment<sup>a</sup>/hpi | Upper airways | Larynx-trachea | Lung                  |
|-----------------|---------------------------|---------------|----------------|----------------------|
|                 |                           | PMNs<sup>c</sup> | PMNs lamina propria | Hyperemia | Red blood cells lumen | PMNs lamina propria | Hyperemia | Hemorrhage | Bronchial-alveolar PMNs | Alveolar macrophages |
|                 |                           | lumen         |                 |          |                      |                  |           |            |                          |                    |
| NTHi375         | Control/12 h              | 1.5 ± 0.9     | 1.2 ± 0.4       | 0.8 ± 0.2 | 1.1 ± 1.3             | 0.8 ± 0.4         | 1.3 ± 0.3 | 0.7 ± 0.3 | 2 ± 0.5                      | 0.7 ± 0.2           |
|                 | TxS/12 h                  | 1.1 ± 0.6     | 1               | 1.1 ± 0.4 | 0.5 ± 1               | 0.7 ± 0.3         | 1.5 ± 0.7 | 0.3 ± 0.5 | 1.3 ± 0.2                      | 0.5 ± 0.4           |
|                 | Control/24 h              | 0.6 ± 0.6     | 1.1 ± 0.3       | 1.1 ± 0.2 | 1.4 ± 0.9             | 0.7 ± 0.2         | 1.7 ± 0.6 | 0.2 ± 0.2 | 1.8 ± 0.4                      | 1.4 ± 0.3           |
|                 | TxS/24 h                  | 2.1 ± 0.9     | 1 ± 0.5         | 1.1 ± 0.5 | 1.7 ± 1.2             | 0.4 ± 0.2         | 1.7 ± 0.4 | 0.6 ± 0.6 | 1.8 ± 0.5                      | 1.7 ± 0.2           |
| NTHi375 ΔthyA   | Control/12 h              | 2.5 ± 0.3     | 0.9 ± 0.2       | 0.9 ± 0.2 | 1.5 ± 0.8             | 0.7 ± 0.3         | 1.1 ± 0.4 | 0.3 ± 0.2 | 1.7 ± 0.2                      | 0.4 ± 0.2           |
|                 | TxS/12 h                  | 2.1 ± 0.4     | 0.9 ± 0.2       | 0.6 ± 0.2 | 0.8 ± 1.1             | 0.8 ± 0.4         | 0.9 ± 0.4 | 0.1 ± 0.2 | 1.7 ± 0.5                      | 0.8 ± 0.4           |
|                 | Control/24 h              | 1.7 ± 1.1     | 0.6 ± 0.2       | 1 ± 0.3  | 0.5 ± 0.8             | 0.7 ± 0.2         | 1.6 ± 0.4 | 0.2 ± 0.2 | 2 ± 0.3                       | 1.1 ± 0.4           |
|                 | TxS/24 h                  | 1.8 ± 0.2     | 0.8 ± 0.2       | 0.8 ± 0.3 | 0.5 ± 1               | 0.6 ± 0.4         | 1 ± 0.4  | 0.1 ± 0.2 | 1.9 ± 0.4                      | 1.1 ± 0.2           |

<sup>a</sup>Control, animals administered vehicle solution; TxS, postinfection, one TxS dose was administered each 6 h.

<sup>b</sup>Statistical comparisons of mean values were performed using one-way ANOVA followed by Fisher’s PLSD multiple-comparison test.

<sup>c</sup>PMNs: infiltrates of polymorphonuclear cells.

<sup>d</sup>More recruitment of PMNs at the upper airway lumen of mice infected with NTHi375 ΔthyA than with WT at 12 and 24 hpi (both, p < 0.05).

<sup>e</sup>More recruitment of PMNs at the upper airway lamina propria of mice infected with WT than with ΔthyA bacteria at 24 hpi (p < 0.05).

<sup>f</sup>Higher alveolar hemorrhage in the lungs of mice infected with WT than with ΔthyA bacteria at 12 hpi (p < 0.05).

<sup>g</sup>Lower proportion of PMNs at the upper airway lumen (24 hpi) of WT bacteria infected mice untreated than TxS treated (p < 0.05).

<sup>h</sup>Larger proportion of PMNs at the alveoli (12 hpi) of WT bacteria infected mice untreated than TxS treated (p < 0.05).

<sup>i</sup>Higher alveolar hemorrhage in the lungs of mice infected with WT at 12 than at 24 hpi (p < 0.01).

<sup>j</sup>Lower numbers of alveolar macrophages at the alveoli of mice infected with WT bacteria at 12 than at 24 hpi (p < 0.01).
of the in-frame insertion in the thyA<sub>NTHi8233</sub> gene. This transient phenotype, in agreement with previous observations (Platt et al., 1983), prompts us to speculate that reversible thymidine auxotrophy may lead into the infrequent isolation of thymidine dependent NTHi strains from respiratory samples. In contrast, TxS resistance level remained unchanged in NTHi8233, independent of auxotrophy reversion. Note of note, increased TMP resistance has been previously found in H. influenzae thymidine prototrophs isolated soon after isogenic thymidine auxotrophs recovery from sputum samples exposed to subinhibitory TMP concentrations, suggesting that TMP resistance could be the result of a suppressor mutation (Platt et al., 1983). Following this observation, the MIC of TMP for NTHi8233 was >32 µg/ml, independent of auxotrophy reversion. The mechanism(s) underlying TMP/TxS resistance in NTHi8233 is currently unknown and will be subject of future study.

Reversible thymidine auxotrophy has been widely shown for S. aureus thymidine dependent-SCVs (Kahl, 2014), and made NTHi8233 inadequate for analysis of the impact of thymidine dependency on NTHi pathogenesis. For this purpose, the thyA gene was inactivated in strains NTHi375 and RdKW20. NTHi thyA mutants displayed alterations in morphology and growth, and dependence on external thymidine, comparable to those shown by S. aureus, Salmonella typhimurium, or S. maltophilia thymidine auxotrophs (Kok et al., 2001; Kahl et al., 2003, 2005; Anderson et al., 2007; Chatterjee et al., 2008). Moreover, NTHi thyA gene disruption caused a reduced PChO level and C3b deposition, impaired airway epithelial adhesion and invasion, and enhanced secretion of IL-8 by cultured epithelial cells. In some cases, minor phenotypic differences were observed between NTHi375 and RdKW20 thyA mutants, likely to be related to the known NTHi genomic heterogeneity (De Chiara et al., 2014). Altogether, the observed in vitro phenotypes, such as a deficient interaction with—and an increased inflammatory response by cultured airway epithelial cells upon infection by ΔthyA mutant strains, are likely to contribute to NTHi thymidine auxotroph attenuation upon murine lung infection. Modification of the biology of infection due to thymidine auxotrophy has been reported for other bacterial pathogens including S. aureus, S. typhimurium, Shigella flexneri, or Vibrio cholerae (Attridge, 1995; Cersini et al., 1998; Kok et al., 2001; Kriegeskorte et al., 2014). We acknowledge that a limitation of this study is the lack of genetic complementation for the thyA gene inactivation. Several approaches were unsuccessfully undertaken for plasmid-encoded thyA<sub>RdKW20</sub> gene complementation into RdKW20ΔthyA, and plasmid-encoded heterologous expression of the thyA<sub>NTHi8233</sub> allele into RdKW20ΔthyA (data not shown). Of note, we exclude a relationship between the thyA gene-related phenotypes shown in this study and overexpression of the immediately downstream gene, a putative homolog of the tada gene encoding a tRNA-specific adenosine deaminase, in the thyA mutants (data not shown). Overexpression of the tada gene has been shown to confer resistance to the bactericidal natural product xanthorrhizol in E. coli (Yogiara et al., 2015). Importantly, addition of external thymidine in the growth media restored, partial or totally, all analyzed in vitro phenotypes. Attenuation of NTHi375ΔthyA in vivo was not restored by infecting bacteria previously grown in the presence of external thymidine, which could be due to the observed partial restoration of thymidine prototrophy by available external thymidine, shown to be dependent on thymidine concentration on MH-F agar.

In addition to the impact of thymidine auxotrophy by inactivation of the thyA gene on NTHi virulence, TxS resistance was shown to increase in the ΔthyA mutant strains. Disruption of the thyA gene has been related to increased TMP resistance in other bacteria (Song et al., 2016). In agreement, the MIC of TMP was 1 and >32 µg/ml for NTHi375 WT and ΔthyA strains, respectively, and was 0.75 and >32 µg/ml for RdKW20 WT and ΔthyA strains, respectively. NTHi375ΔthyA pre-grown in chocolate+Thy was cleared faster in TxS-treated than in untreated mice, and the reason(s) for this observation is currently unknown. Although, originally intended to restore the observed increased expression of the nupC gene upon thyA inactivation, mutant pre-growth in sBHI with thymidine further amplified nupC expression, compared to sBHI. It should be noted that increased nupC gene expression by a thyA mutant in S. aureus was previously restored by complementing thyA gene disruption (Chatterjee et al., 2008). Our observation was unexpected and formally incomparable to that made for S. aureus. We speculate that it could be associated with the lack of growth inhibition around TxS discs observed for thyA mutant strains when pre-grown in chocolate+Thy, therefore suggesting that uptake of external thymidine by the NupC nucleoside transporter could contribute to circumvent the effects of TxS upon de novo thymidylate biosynthesis in NTHi. We suggest here for the first time that NupC could function as a primary thymidine transporter in NTHi if de novo thymidylate synthesis is blocked. Search in all available genome sequenced H. influenzae strains did not reveal the presence of additional nucleoside transporters in this bacterial species. As expected, we unsuccessfully attempted to generate a NTHi double mutant strain lacking both the thyA and nupC genes (data not shown), further supporting the proposed role for NupC.

Three additional aspects deserve further discussion. First, S. aureus thymidine dependent SCVs grow on Columbia blood agar (Kriegeskorte et al., 2014); S. maltophilia thymidine dependent SCVs grow on sheep blood-, brucella-, and chocolate-agar (Anderson et al., 2007), but neither grow on MacConkey agar and M9 minimal medium (Anderson et al., 2007), nor S. typhimurium thyA mutants grow on LB agar (Kok et al., 2001). Our results show that NTHiΔthyA thymidine auxotrophs grow rendering normal size colonies on chocolate—and sBHI-agar, but fail to grow on MH-F agar. Therefore, we may not consider the term thymidine dependent SCVs for H. influenzae. Second, S. aureus SCVs are recovered from several human and animal specimens, and are a highly dynamic subpopulation optimized for persistence, enabling the bacteria to hide inside the host cell without eliciting a strong host response (Kahl, 2014; Kahl et al., 2016). This may be unlikely for NTHi thymidine auxotrophs, given that we observed a significantly impaired airway epithelial cell invasion by the NTHi375ΔthyA mutant strain. Third, this study does not tackle the induction of NTHi thymidine auxotrophy by TxS challenge, and we therefore cannot speculate on the selection for thymine auxotrophs and TxS resistance due to prolonged antibiotic exposure. Unexpectedly, no statistically significant connection has been found between
regional TxS use and resistance among *H. influenzae* isolates (Karpanoja et al., 2008). We should also consider the potential emergence of mutations in the *thyA* gene as a consequence of treatment with other antibiotics with mutagenic activity such as ciprofloxacin, which concomitantly increases TMP resistance (Song et al., 2016). Hence, emergence of TxS resistance in response to antibiotics needs further analysis and continuous monitoring.

In conclusion, this study shows for the first time the impact of thymidine auxotrophy by disruption of the thymidylate synthase *thyA* on *H. influenzae* morphology and interplay with the host airway and, indirectly, on its resistance to TxS. We also show NupC as a potential facilitator of external thymidine uptake upon inhibition of thymidylate *de novo* synthesis. Thymidine auxotrophy lowers NTHi virulence, but also provides an advantage under TxS exposure. Thus, our results should be considered for the consequences of TxS administration in the clinical settings. Further studies will contribute to better assess the emergence of NTHI thymidine auxotrophs in clinical samples, likely to be currently underestimated, and to understand if NTHI thymidine dependency may be a response to antibiotic treatment with a survival advantage in specific environments.

**AUTHOR CONTRIBUTIONS**

IR, SM, BE, AF, JM, NL, MB, JR, FT, and CL have participated in the design and fulfillment of the experimental work. IR, SM, CA, JL, JY, and JG carried out in the conceptual design of the study.

IR, SM, JL, JY, and JG have written the manuscript (text, tables, and figures). All authors have participated in the correction of the manuscript to its final version.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: http://journal.frontiersin.org/article/10.3389/fcimb.2017.00266/full#supplementary-material

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