In Vivo Biosynthesis of Terpene Nucleosides Provides Unique Chemical Markers of \textit{Mycobacterium tuberculosis} Infection

Graphical Abstract

Highlights

- \(N^6\)-tuberculosinyladenosine (formed from 1-tuberculosinyladenosine) is described

- These lipid-linked nucleosides are detected in \textit{M. tuberculosis} infected mouse lung

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In Brief

To make better diagnostic tests for tuberculosis, we identified molecules that are abundantly produced by \textit{M. tuberculosis} within infected mammalian tissues. Two molecules, 1-TbAd and \(N^6\)-TbAd are specifically produced by \textit{M. tuberculosis} and can be sensitively detected using mass spectrometry, making them attractive targets for clinical test development.
In Vivo Biosynthesis of Terpene Nucleosides Provides Unique Chemical Markers of Mycobacterium tuberculosis Infection

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SUMMARY

Although small molecules shed from pathogens are widely used to diagnose infection, such tests have not been widely implemented for tuberculosis. Here we show that the recently identified compound, 1-tuberculosinyladenosine (1-TbAd), accumulates to comprise >1% of all Mycobacterium tuberculosis lipid. In vitro and in vivo, two isomers of TbAd were detected that might serve as infection markers. Using mass spectrometry and nuclear magnetic resonance, we established the structure of the previously unknown molecule, N6-tuberculosinyladenosine (N6-TbAd). Its biosynthesis involves enzymatic production of 1-TbAd by Rv3378c followed by conversion to N6-TbAd via the Dimroth rearrangement. Intact biosynthetic genes are observed only within M. tuberculosis complex bacteria, and TbAd was not detected among other medically important pathogens, environmental bacteria, and vaccine strains. With no substantially similar known molecules in nature, the discovery and in vivo detection of two abundant terpene nucleosides support their development as specific diagnostic markers of tuberculosis.

INTRODUCTION

Tuberculosis (TB) remains a leading cause of death worldwide, resulting in 1.5 million deaths annually (World Health Organization, 2014), yet no rapid, sensitive, and specific diagnostic test exists. Diagnosis based on detection of Mycobacterium tuberculosis in patient samples mainly relies on sputum microscopy, which is insensitive, or on in vitro culture, which is slow, insensitive and infeasible in many clinics. T-cell antigen recall tests, such as intradermal injection of purified protein derivative (PPD) or interferon-γ release assays (IGRA) are in widespread use (Lalvani and Pareek, 2010) but give delayed results or are expensive and have suboptimal test characteristics related to sensitivity and specificity. Vaccination with live Bacille Calmette-Guérin (BCG) in most parts of the world leads to antigen-specific T-cell responses, which create false-positive results, rendering the PPD test unusable in many populations. Accordingly, there is now strong consensus that developing better diagnostic tests for M. tuberculosis infection is the key issue for improved disease control through rapid initiation of antibiotics and categorization of patients for vaccine trials (Hankom et al., 2008).

Detection of pathogen-specific shed molecules or antigens provides rapid and specific diagnosis of many infectious diseases. Such antigen tests have long been a mainstay of diagnosis for infection by cryptococci, legionella, and other pathogens (Shelhamer et al., 1996). The strengths of antigen test technology are high diagnostic specificity and rapid detection of molecules using a simple ELISA of urine or serum (Couturier et al., 2014). Therefore, the key criterion for discovery of chemical targets for antigen tests is specific expression of the target by the disease-causing pathogen, combined with lack of expression among other microbes, especially those that are abundant in the environment or cause diseases that mimic the disease of interest. Other desirable criteria related to test sensitivity involve
identifying targets with broad expression among most infecting strains in clinical settings, expression of the antigen at high concentrations, expression in vivo under conditions of infection, and lack of host degradation or metabolism to unrecognizable chemical forms.

In addition to antigen capture, pathogen-specific molecules can be coated onto plastic and used to detect target-specific host antibodies, functioning as a serological test. Such tests have not yet moved into widespread clinical use for TB due in part to specificity concerns that may be related to immune responses to environmental mycobacteria, mildly pathogenic mycobacteria, or live vaccine strains (Lawn et al., 2012) against TB (Baumann et al., 2014).

Despite the widespread use of antigen and serological tests in other infectious diseases, neither type of test is widely used for TB. Although the urine lipoarabinomannan (LAM) ELISA has usefulness for TB-HIV coinfection (Lawn et al., 2012), no antigen or serological test has emerged as having widespread clinical usefulness for tuberculosis. The current chemical targets for testing were chosen based on their ready availability and represent only a small fraction of the candidate small molecules that could be developed. For example, among 169 subclasses of mycobacterial lipids in the MycoMass and Lipid DB databases, more than 90% are expressed only by mycobacteria (Layre et al., 2011; Sartain et al., 2011). Thus, the potential range of specific mycobacterial targets for diagnostics development is vast and largely unexplored.

We initiated a comprehensive effort to discover specific chemical targets for diagnostic testing using a newly developed high-performance liquid chromatography (HPLC)-mass spectrometry (MS)-based lipidomics platform (Layre et al., 2011). A recent comparative lipidomics screen of mycobacteria sought to identify those molecules that are present in *M. tuberculosis* but are lacking in a BCG vaccine strain (Layre et al., 2014). These two species are evolutionarily related and share more than 99% sequence identity but only *M. tuberculosis* causes widespread disease. BCG has been administered to more than 3 billion people and it induces immune responses that can cause false-positive immunological tests for TB (World Health Organization, 2014). By identifying cell wall lipids absent in BCG, we reasoned that such molecules would also be lacking in BCG vaccines and in other less related bacteria that confound the diagnosis of TB. Also, given the lower infectious potential of BCG, molecules selectively expressed in *M. tuberculosis* might be virulence factors. This screen identified a previously unknown diterpene nucleoside, 1-tuberculosinyladenosine (1-TbAd) (Layre et al., 2014).

Having established the lack of 1-TbAd in BCG, here we sought to determine if 1-TbAd has the key test characteristics needed for use as a diagnostic target, including high abundance, specific expression, and shedding from intact bacteria in ways that lead to detection in vivo. Unexpectedly, we identified an abundant but previously unknown diterpene nucleoside derived from *M. tuberculosis* present in vivo. The discovery of two terpene nucleosides, which are specifically and abundantly expressed by *M. tuberculosis*, provides two highly attractive targets for development for new diagnostic tests for TB.

**RESULTS**

1-TbAd Is a Major Lipid in *M. tuberculosis*

1-TbAd was identified in *M. tuberculosis* using an electrospray ionization (ESI)-MS-based lipidomics platform. Among 7852 ions, 1-TbAd (C_{30}H_{46}N_{5}O_{4}^+; m/z 540.3545) was the second most intense ion in the lipidome (Layre et al., 2014). The high intensity might have resulted from 1-TbAd’s accumulation to high concentration. Yet it was difficult to imagine that any abundant class of molecule would have escaped detection over decades of study of a pathogen of worldwide importance. Alternatively, 1-TbAd might have been present only in trace amounts, yet its intrinsically charged nature and amphipathic character might have promoted particularly efficient ionization in ESI-MS. To measure the mass of 1-TbAd as a percentage of all lipid, we analyzed extracts from *M. tuberculosis* strain H37Rv using the method of standard additions. We compared the area under the curve of retention time versus intensity measured at the mass (m/z 540) of 1-TbAd (A_{540}) and phosphatidylethanolamine (A_{220}), which controls for the efficiency of ESI-MS detection. As with other polar lipids (Layre et al., 2011), we observed a nearly linear relationship between mass input and 1-TbAd signal intensity (Figure 1A). In three experiments, we determined that 1-TbAd comprises 1.1%, 1.4%, and 1.5% of mycobacterial lipids. Thus, contrary to expectations that terpene nucleosides previously escaped detection due to scarcity, 1-TbAd is a highly abundant molecule that comprises a major class of lipid in *M. tuberculosis*. The basis for lack of prior detection remains unknown, but the near co-elution of 1-TbAd and abundant membrane phospholipids in normal-phase thin-layer chromatography and HPLC methods might have obscured 1-TbAd detection in the past (Figure 1B).

**Constitutive Biosynthesis of 1-TbAd**

1-TbAd was initially isolated from bacteria grown in a formulated rich medium (7H9) in late logarithmic phase (Layre et al., 2014). Because rv3377c and rv3378c protect bacteria against low pH, we reasoned that TbAd production might be pH responsive. However, we detected equivalent production with high absolute intensity (>10^6 counts) when growing at neutral (7.4) or lysosomal pH (4.5) (Figure 1C). Furthermore, 1-TbAd production was not substantially inhibited during the stationary phase, with laboratory strains of different origins (Layre et al., 2014), or when growing in minimal (Sauton) or rich (7H9) medium (Figure 1D). Thus, 1-TbAd is constitutively produced at high concentrations among diverse in vitro conditions and we could not identify signals for its inhibition.

**1-TbAd Biosynthetic Gene Expression**

To function as a sensitive marker of infection, molecules must be expressed in most strains of *M. tuberculosis* that infect patients and among the seven recognized *M. tuberculosis* lineages that exist worldwide (Comas et al., 2013). 1-TbAd derives from geranylgeranyl pyrophosphate (GGPP) and adenosine, which are present in nearly all organisms. Lipid cyclization and coupling to adenosine are performed by specialized enzymes, Rv3377c (prenyl cyclase) and Rv3378c (tuberculosinyl transferase), which together with polyprenyl synthases comprise a functional gene island (Mann and Peters, 2012) that is necessary and sufficient.
for 1-TbAd biosynthesis (Layre et al., 2014). Knowledge of the essential TbAd biosynthetic genes allowed a survey of the genomes of 432 phylogeographically diverse clinical isolates of the M. tuberculosis complex (MTB complex) for an intact biosynthetic locus. TbAd genes were detected among all clinical strains examined.

We identified 25 unique single nucleotide polymorphisms (SNPs) in rv3377c and 22 unique SNPs in rv3378c affecting a total of 150 M. tuberculosis strains (Table S1). While the proportion of non-synonymous SNPs (nSNP) (18/25 in rv3377c vs 14/22 in rv3378c) in both genes was similar, more strains contained nSNPs in rv3377c (101 strains with nSNP in rv3377c versus 52 strains with nSNP in rv3378c). To understand the selection forces driving rv3377c and rv3378c sequence diversity, we calculated the ratio of the rates of non-synonymous to synonymous single nucleotide changes (dN/dS) of both genes with respect to the inferred MTB complex ancestor. The whole-gene dN/dS ranged from 0.97 to 0.47 for rv3377c and rv3378c, respectively. The dN/dS for rv3378c was similar to the global dN/dS of the M. tuberculosis genome (0.45–0.67) in MTB complex (Comas et al., 2010), whereas the dN/dS of rv3377c was significantly higher.

Bioinformatic algorithms that seek to identify which mutations are likely to alter protein function suggested that 60% (15/25) of nSNPs identified in rv3377c are not likely to affect protein structure (Table S1). However, ten SNPs are predicted to have notable effects on structure. The replacement of a glycine at position 31 with a valine is predicted to affect the protein structure in the entire lineage 6, also known as M. africanum. In summary, we found that most clinical isolates have an intact rv3377c-rv3378c locus and no clear evidence for selection driving the mutations identified, with the exception that lineage 6 might have an inactivating mutation.

Pathogen-Specific Gene Expression
To determine whether 1-TbAd biosynthesis exists in species other than M. tuberculosis, we sought orthologs of rv3377c and rv3378c across the biological kingdoms. Confirming a prior analysis (Mann and Peters, 2012), few candidate orthologs were found among non-mycobacterial species. Low stringency searches did return candidate rv3378c orthologs in Zea mays and Dictyostelium discoideum. However, considering all non-mycobacterial species, we did not identify candidate orthologs organized into a locus with both essential genes present. Among mycobacteria, orthologs of rv3377c and rv3378c could not be identified in most species, including disease-causing members of the M. avium complex, M. kansasii, and M. marinum (Figure 2A). Instead, orthologs were present only within the MTB complex: in M. bovis, M. kansasii, and M. africanum. However, coding mutations were found in the locus in all species other than M. tuberculosis as detailed in Figure 2A. Two mutations were found in the Pasteur strain of BCG (Pasteur), including a frameshift mutation that is likely inactivating (Figure 2A). Thus, we could not identify an intact orthologous locus outside the MTB complex and we found a wild-type sequence only in M. tuberculosis.

However, gaps in the analysis remained. Some but not all of the rv3378c mutations, including those present in M. africanum, have been proven to inactivate lipid biosynthesis (Chan et al., 2014; Layre et al., 2014). Second, BCG vaccine strains are of particular interest because they cause false-positive immunological tests. Although the locus in BCG strain Pasteur was inactivated through frameshift, other BCG or M. bovis strains might never have acquired these mutations, or they might have done so and then reverted to wild-type sequences. The 12 widely used BCG vaccine strains show variations in efficacy (Ritz et al., 2008), so differential expression of TbAd might account for this (Pethe et al., 2004). However, further analysis of all common vaccine strains used worldwide (Pasteur, Copenhagen, Japan, Mexican, Australian, Russia, Glaxo, Prague, Phipps, Connaught,
Denmark, Tice) confirmed the presence of the frameshift mutation in all cases, so 1-TbAd is likely absent from all major vaccines used worldwide.

1-TbAd Expression in Pathogens

A general limitation of genetic analysis is that genes might exist that are functionally equivalent to rv3377c and rv3378c but lack the sequence identity needed for identification as orthologs. Therefore, we undertook biochemical analysis for 1-TbAd production, focusing on non-tuberculous mycobacteria and microbes, whose infection can mimic tuberculosis disease. By adapting the existing HPLC-MS lipidomics protocol (Layre et al., 2011), we validated a rapid method to specifically detect 1-TbAd within complex lipid mixtures. Analyzing total lipids, molecular events (linked intensity, mz, and retention time values) corresponding to all lipids are recorded and the events corresponding to the mass (m/z 540.35) and retention time (23 min) of authentic 1-TbAd are reported (Figure 2B). Monitoring of unFractionated lipids avoids time-consuming methods needed to purify 1-TbAd, as well as artifacts introduced through sample handling. Datasets can be subjected to further data mining for events corresponding to other lipid products that are upstream and downstream in biosynthetic pathways, isomers of 1-TbAd, as well as TbAd variants with defined changes in mass due to chain length variation or oxidation. Initial validation showed that no other mycobacterial lipid obscured the event corresponding to the mass and retention time (tR) of 1-TbAd. Furthermore, total lipid extracts of the human THP-1 macrophage-like cell line did not generate detectable signals at this mass and tR coordinate, even when amplified 1000-fold (Figure 2B, inset). Thus, no host lipid interfered with TbAd detection in this system, which is a key consideration for diagnostic applications.

Lipid extracts from non-actinomicete bacteria (Escherichia coli, Staphylococcus aureus) and fungi (Aspergillus fumigatus, Candida albicans) did not produce a 1-TbAd signal. Turning to mycobacteria, we could not detect 1-TbAd among reference strains of environmental (M. fallax) or non-pathogenic laboratory strains of mycobacteria (M. smegmatis, M. phlei) that lack rv3378c orthologs (Figure 2C). In agreement with the genetic results, we did not detect 1-TbAd among disease-causing bacteria that are related to M. tuberculosis but lack identifiable orthologs of rv3377c-3378c (M. avium, M. marinum) or those with orthogonal loci containing a known frameshift mutation (M. bovis). Among all organisms tested to date, only M. tuberculosis produces 1-TbAd.

Detection of 1-TbAd In Vivo

To determine if 1-TbAd is detectable in vivo, we infected BALB/c mice via inhalation for 21 days followed by collection of whole-lung homogenates. Target validation in vivo does not require scanning of the complete lipidome, but instead focuses on sensitive and specific detection of predetermined targets, which are likely to be highly diluted within host tissues. We developed a reversed-phase method to increase chromatographic acceleration. Table S1 lists polymorphisms for these genes in the M. tuberculosis complex.

Figure 2. Genetic and Biochemical Analysis of TbAd Biosynthetic Genes, rv3378c, rv3377c, and 1-TbAd in Lung Pathogens, Environmental Mycobacteria, and Vaccine Strains

(A) The TbAd locus encodes a geranylgeranyl pyrophosphate cyclase (Rv3377c) and tuberculosyl adenosine transferase (Rv3378c). Orthologous genes are shown by arrows and SNPs are indicated by vertical lines. Table S1 lists polymorphisms for these genes in the M. tuberculosis complex. (B and C) Lipid extracts from human macrophage-like cells (THP-1) containing 1-TbAd, M. tuberculosis H37Rv, or the indicated species were subjected to HPLC-MS.

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resolution, while taking advantage of the specificity and accuracy of triple quadrupole mass detection. This method detected a 1-TbAd standard at 11.8 min based on its known transition for 1-TbAd (m/z 540 → 408 → 136). Direct analysis of unfractionated lipids from whole-lung homogenates did not detect signal corresponding to 1-TbAd in uninfected lung but did detect a signal corresponding to 1-TbAd that was well above background signals in six of six infected mice tested, with representative data shown in Figure 3A. Thus, 1-TbAd is produced in vivo and is readily detected ex vivo in a one-step HPLC-MS method.

**Unknown Terpene Nucleoside**

Ion chromatograms of the ion 540 → 408 also showed a second, later-eluting peak (14.6 min) with transitions that were distinct from 1-TbAd in all six mice tested (Figure 3A). Unlike 1-TbAd, the unknown lipid contained a 540 → 148 ion and a relatively bright 540 → 408 transition that was equivalent in intensity to the 540 → 136 ion. Returning to normal-phase HPLC-quadrupole time-of-flight (Q-TOF)-MS experiments using in vitro grown *M. tuberculosis*, ion chromatograms measured at m/z 540.4 also showed two peaks at ~25 and ~7 min, which corresponded to late-eluting 1-TbAd and an early-eluting unknown lipid detected in mice (Figure 3B). Collision-induced dissociation (CID)-MS analysis of the unknown showed fragment ions of m/z 148.062 and 408.313, matching the pattern of the late-eluting unknown from reversed-phase chromatography, suggesting that they were the same molecule (Figures 3A and 3C). Thus, the unknown later-eluting compound detected in mice was likely derived from *M. tuberculosis* rather than the host. Also, separate tracking of the unknown and 1-TbAd in the bacterial pellet and conditioned supernatant demonstrated a much higher ratio of signal for the unknown in supernatants (Figure 3B). Thus, the early-eluting unknown was likely generated during or after transit from the cytosol to the extracellular space.

**1-TbAd Isomers**

The unknown (m/z 540.355) and 1-TbAd (m/z 540.3545, calculated) had identical molecular ion masses (±2 ppm) and similar fragmentation patterns. The unknown had a much lower retention time in normal-phase chromatography. Therefore, the unknown was likely a much less polar isomer of 1-TbAd. CID-MS ions were assigned unequivocally as C_{30}H_{45}N_{5}O_{4} (m/z 136.0618) and C_{25}H_{38}N_{5}O_{4}^{+} (m/z 268.1040), matching the mass of adenine and adenosine, respectively. The neutral loss leading to m/z 268.1040 suggested the loss of C_{20}H_{32}O_{2}, likely a diterpene moiety (Figure 3B). A fragment ion with the formula C_{25}H_{38}N_{5}O_{4}^{+} (m/z 268.3125) matches the mass of geranylgeranyl (GG) pyrophosphate and tuberculinosyl pyrophosphate is hydrolyzed to isotuberculinosin in vitro (Nakano et al., 2011). Therefore, CID-MS of the unknown detected ions consistent with a diterpene adenosine structure, but bright ions at m/z 408.31, 148.06 were seen only in the early-eluting isomer (unknown).
1-GGAd and 1-isoTbAd were candidate structures (Figure 4A). However, they would be expected to have largely the same ionic properties as 1-TbAd. Alternatively, a tuberculosinyl linkage at the N6 adenosine position would mimic common N6-linked adenine compounds including zeatin (Heyl et al., 2012) and would be expected to alter the ionic properties observed in the unknown. Therefore, we considered N6-tuberculosinyladenosine as a candidate structure (Figure 4A) and we synthesized N6-(E,E,E)- and N6-(E,E,Z)-geranylgeranyl-adenosine as a model N6-linked compound to evaluate this hypothesis (Figure 4B; Data S2).

Identification of N6-TbAd
Detection of the isomer in cultured M. tuberculosis provided a route to purifying it in greater quantity. After growing M. tuberculosis cultures in roller bottles, ~50 mg of lipid was fractionated by normal-phase open-column chromatography followed by reversed-phase HPLC. Adding 0.2% trifluoroacetic acid (TFA) to the mobile phase increased the retention of 1-TbAd in reversed-phase chromatography but did not affect the unknown isomer, confirming that the two compounds differed in their ionic properties. This purification sequence produced ~300 μg of each isomer so that the combined yield (~1.2%) confirmed ESI-MS studies (Figure 1A).

Analysis of the MS data and 1H correlated spectroscopy (COSY), nuclear Overhauser effect spectroscopy (NOESY), and heteronuclear multiple-quantum correlation (HMQC) nuclear magnetic resonance (NMR) spectra (800 MHz) (Table 1) established the structure of the unknown as N6-tuberculosinyladenosine. The resonances from the tuberculosinyl side chain positions 14 and 15 (labeled with a and b in the corresponding N6-(3-methyl-2-butenyl)adenosine spectrum) adjacent to the heterocyclic adenine moiety are shifted similarly, relative to 1-TbAd, due to the positive to neutral charge shift.

The presence of the m/z 148 fragment in N6-TbAd results from charge-directed fragmentation after protonation of a double bond that can occur with the N6-TbAd but not the analogous 1-TbAd.
The combination of 1-D and 2-D NMR data lead to the determination of the side group structure as a tuberculosinyl group in $N^6$-linkage, defining $N^6$-TbAd. Detailed spectra are shown in Data S1. The expected COSY and NOESY cross peaks are seen within the terpene, base and sugar fragments. The only NOESY cross peak between the three fragments is between H1 and H4. The allylic methylene peak from the two H15 protons is broad rather than the expected doublet due to slow rotation about the C6-N bond. *, not observed.

**Table 1. NMR Analysis**

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<td>29.3</td>
<td>1.02</td>
<td>2', 3', 10', 18'</td>
<td></td>
</tr>
<tr>
<td>20α</td>
<td>16.4</td>
<td>0.65</td>
<td>1', 7', 11', 17'</td>
<td></td>
</tr>
</tbody>
</table>

The combination of 1-D and 2-D NMR data lead to the determination of the side group structure as a tuberculosinyl group in $N^6$-linkage, defining $N^6$-TbAd. Detailed spectra are shown in Data S1. The expected COSY and NOESY cross peaks are seen within the terpene, base and sugar fragments. The only NOESY cross peak between the three fragments is between H1 and H4. The allylic methylene peak from the two H15 protons is broad rather than the expected doublet due to slow rotation about the C6-N bond. *, not observed.

*Ottria et al., 2010*. The adenine protons at δ 8.24 (H-8) and 8.26 (H-8) are characteristic of an $N^6$-substituted adenosine and are different from those of 1-TbAd at δ 8.53 and 8.66. All COSY and NOESY correlations are consistent with the assignment of the unknown as $N^6$-TbAd. The absorption of the allylic methylene group is a broad peak at 4.22–4.16 ppm due to slow rotation at room temperature around the C-N bond (Uzawa and Anzai, 1988). The diagnostic ion at m/z 148 in the $N^6$-TbAd is explained by an alternate protonation of the double bond at C-14” and subsequent charge-directed fragmentation, which would be suppressed in 1-TbAd, having its positive charge stabilized in the adenine ring (Figure 4D).

**Dimroth Rearrangement of 1-TbAd to $N^6$-TbAd**

A plausible mechanism to account for the biological origin of $N^6$-TbAd is that 1-TbAd converts to $N^6$-TbAd by a Dimroth...
Figure 5. A Two-Step Mechanism for 6-TbAd Biosynthesis
(A) The Dimroth rearrangement represents a specific and plausible mechanism that accounts for the rearrangement from cationic 1-linked adenosine to neutral 6-linked adenosine compounds. The pKa values derived from analogous compounds and illustrate that only the 1-linked form is predominantly charged.

(B) Incubation of recombinant Rv3378c, tuberculosinyl pyrophosphate, and adenosine yielded a strong signal for the 1-TbAd isomer (~24 min) but no detectable signal at the retention time for 6-TbAd (~8 min).

(C) 1-TbAd was treated with dimethylamine in methanol to effect the conversion (Ottria et al., 2010). MS3 analysis of the reaction products detected the diagnostic fragment ion for 6-TbAd at m/z 148.2.

(D) Lung lipids were spiked with 1-TbAd, extracted for lipid and analyzed using normal-phase HPLC-Q-TOF-MS under conditions that detect both isomers.

(E) These data support a two-step mechanism in which 1-TbAd is the predominant or sole product of Rv3378c followed by conversion to 6-TbAd, which can occur during export of 1-TbAd to the extrabacterial space.

DISCUSSION

These studies show that 1-TbAd and the newly identified in vivo biotransformation product, 6-TbAd, possess key features of high-quality targets for development in antigen capture diagnostic or serological tests. These features include abundant expression and efficient shedding, as well as absence of expression in common sources of false-positive test results including vaccine strains, non-mycobacterial lung pathogens, and environmental bacteria. Furthermore, these studies provide evidence for intact biosynthetic genes among clinical strains and expression by reference strains under many different conditions surveyed, suggesting that it could be a sensitive biochemical marker of infection. Taking advantage of the broad lipidomics detection methods, the data demonstrate sensitive detection of both TbAd isomers without interference from any other host.
bacterial, or fungal lipids using a standardized and simple HPLC-
MS test.

Another criterion for use of shed molecules as diagnostic tools is that they remain intact during in vivo infection or, if they are metabo-
lized by the host, generate recognizable transformation products that are distinct from all host molecules. These data identify the complete struc-
ture of N⁶-TbAd as a previously unknown in vivo biotransformation product of M. tuberculosis. Here we report a small proof-of-principle study in which both TbAd isomers are readily detected in the lung. Starting from a systematic screen of all M. tuberculosis-specific compounds, these two terpene nucleosides meet all the criteria needed for entry into human population studies.

Furthermore, these data determined the origin and stepwise appearance of the TbAd isomers in ways that add to a basic un-
derstanding of tuberculosis pathogenesis. Prior studies by Pethe et al. (2004) identified that Rv3378c acts by some unknown mechanism to inhibit phagosomal acidification and promote mycobacterial survival in macrophages. Rv3378c is localized in the bacterial cytosol and yet might act in some way on events in the phagosomal space. Our results identify the product of Rv3378c as an amphipathic diterpene nucleoside (Layre et al., 2014) and show that the product of Rv3378c is efficiently ex-
ported to the extracellular space, so that this product does escape to the phagosomal compartment that is involved in pH regulation. Unexpectedly, 1-TbAd undergoes biotransformation to N⁶-TbAd, a previously known in vivo metabolite within the infected host.

1-TbAd and N⁶-TbAd show only one difference in structure but the site of tuberculosinyl linkage determines their pKₐ and charge state. Quantitative consideration of the ionic properties of 1-TbAd (pKₐ = 8.5) and N⁶-TbAd (pKₐ = 3.8) (Kapinos et al., 2011) predict that N⁶-TbAd is neutral at pH 7.4, whereas 1-TbAd exists as a mixture of [TbAd] and [TbAdH⁺]. During active infection, M. tuberculosis typically grows in moderately acidic phagosomal compartments or extracellular caseous material. Therefore, 1-TbAd would change in charge states as it leaves the neutral pH environment of the mycobacterial cell wall, traverses the acidic phagosome, and enters the neutral environment of infected tissues and finally the peripheral circu-
l nation. These effects would alter its water solubility and adher-
ence to anionic membranes in the three compartments. Furthermore, the Dimroth rearrangement is favored under basic conditions. At lower pH, the concentration of nucleophiles decreases so 1-TbAd is less prone to rearrangement in the acidic environment of localized acute infection. Thus, 1-TbAd has an intrinsic chemical feature that would stabilize its structure while in the phagosome and the N⁶-form might be more efficiently generated in non-acidic compartments that are distant from the site of infection. In this scenario, 1-TbAd is more important in control of the export to the phagosome and N⁶-TbAd might represent an altered, possibly inactivated, form of 1-TbAd whose importance derives its role as a recognizable infection marker.

These studies set the stage for human population studies to detect both TbAd isomers as targets for diagnosis in clinically assayable fluids such as sputum, serum, and urine. Both isomers are readily detected ex vivo using a routine liquid chromatography-MS method. For serological applica-
tions, the difference in charge state strongly predicts differing host antibody responses to 1-TbAd and N⁶-TbAd, and know-
ledge of these differing ionic properties may guide separate development of these terpene nucleosides as targets for ELISAs in the future. Although LAM ELISAs have emerged as a clinical test with some utility, one limiting factor for this and other mycobacterial lipid-specific ELISAs is thought to be false-positive results from subclinical exposures to non-tuber-
culosus mycobacteria or BCG vaccination (Baumann et al., 2014; Lawn, 2012). Here we demonstrated the lack of both TbAd isomers and the rv3377c-3378c genes in common sources of false-positive immune response. These discovery-
based chemical studies open two new paths to the develop-
ment of simple and specific antigen or serological tests, which are urgently needed for better care of patients and vaccine research.

**SIGNIFICANCE**

The discovery and detection of 1-tuberculolinsyladenosine and its rearrangement product N⁶-tuberculolinsyladenosine, produced in vivo during the course of mycobacterial infec-
tion in a murine model, provides unique chemical markers to diagnose tuberculosis disease.

**EXPERIMENTAL PROCEDURES**

**Bacteria Cultures**

M. tuberculosis was grown in the absence of detergent in 7H9 medium supplemented with 10% albumin-dextrose-catalase or oleic acid-albumin-dextrose-catalase or in Sauton’s medium until mid-log phase with or without phosphate-citrate buffers adjusted at the given pH. Supernatant was sepa-
rated from bacteria by centrifugation at 2000 x g and filtration of supernatant cultures (0.2 μm). Cell-associated or secreted lipids were extracted using suc-
cessive contact in chloroform and methanol mixes or ethyl acetate, respec-
tively, as previously described (Layre et al., 2011).

**Purification of 1-TbAd and the Unknown TbAd Isomer**

Lipids were prepared by serially extracting desiccated or wet pellets of Myco-
bacterium tuberculosis H37Rv and other named species with 1:2 (v/v), 1:1, and then 2:1 chloroform/methanol. The combined extracts were dried at room tem-
perature and stored in 1:1 chloroform/methanol solution (~10 mg/ml). Fifty mil-
ligrams was concentrated under nitrogen resulting in a slurry that was loaded on an open silica gel column (2 cm × 1.6 cm). Lipids were eluted with 10 ml of solvent in the following sequence: chloroform, chloroform/isopropanol (95:5, v/v), chloroform/isopropanol (90:10, v/v), and chloroform/methanol (60:50, v/v). Fractions were monitored for ions with a nominal m/z of 540 and the m/z 540 → m/z 408 → m/z 148 transitions in the nanospray MS² and MS³, with the m/z 148 ions seen in the unknown (N⁶-TbAd, not 1-TbAd. N⁶-TbAd eluted using isopropanol/chloroform while 1-TbAd required meth-
anol/chloroform for elution. After evaporation, 1-TbAd and its isomer were further purified using reversed-phase HPLC using an octadecyl-modified silica (5 μm) semi-preparative column (250 × 10 mm) run under isocratic conditions at 3.0 ml/min. 1-TbAd was purified using 99.5:0.5 (v/v) methanol/water/ trifluoroacetic acid monitoring by UV at 260 nm. Collected fractions were evaporated to dryness in 0.5-ml portions with 2.0 ml of acetonitrile added to help minimize exposure of the collected 1-TbAd to TFA. N⁶-TbAd was purified using 5:90:5 chloroform/methanol/water (v/v/v) as the mobile phase, moni-
tored by UV at 260 nm, and eluted at 14.5 min.

**Ion Trap MS**

Multistage CID-MS experiments were completed on a linear ion trap (Thermo Scientific LXQ) using a nanoelectrospray ionization source and borosilicate glass pipettes pulled to a 2-μm tip.
Basic Dimroth Conversion
1-TbAd was converted to N6-TbAd by dissolving a small portion of 1-TbAd in 2 M dimethylamine in methanol at room temperature for 3 days.

NMR Analysis
A Bruker Avance 800 NMR instrument was used to analyze samples dissolved in deuterated methanol (0.5 mg/ml) that were compared with a 1-TbAd standard (Layre et al., 2014).

HPLC-ESI-Q-TOF Analyses
Extracted cell-associated or secreted lipids were concentrated under reduced pressure at room temperature. One hundred micrograms were resuspended at 0.5 mg/ml in 70:30 hexane/isopropyl alcohol and 20 µl were injected for analysis by HPLC-ESI-MS (Agilent 6520 QToF) as described (Layre et al., 2014).

Reversed-Phase HPLC-MS
Mouse (BALB/c) lungs were harvested after 3 weeks and analyzed using an injection volume of 50 µl using a reversed-phase, linear gradient elution method developed using a Zorbax Eclipse Plus-C18 column (2.1 mm × 50 mm, 3.5-µm particle size; Agilent Technologies). The solvents were (A) 0.1% formic acid in milliQ water and (B) 0.1% formic acid in acetonitrile. After equilibration for 8 min with 100% solvent (B), a two-part linear solvent gradient was used as follows: 0–10 min, 0% to 100% solvent B; 10–22 min, 100% to 0% solvent B. Mouse experiments were conducted under oversight of the Animal Ethics Subcommittee of the University of KwaZulu-Natal (protocol 107/13).

N6-TbAd Structural Analysis
Synthesis of N6-linked adenosine analogs and detailed descriptions of N6 products are described in the Supplemental Experimental Procedures.

Genetic Analysis of rv3377c and rv3378c
Detection of SNPs in rv3377c and rv3378c was done from multiple alignments of the orthologous sequences of both genes retrieved from the genomes of 431 strains representative of the seven main phylogenetic lineages of the MTB complex, comprising 271 published genomes (Comas et al., 2010), 161 whole genomes that are currently being analyzed, and 12 published BCG strains (Copin et al., 2014). Nucleotides differing between the query strains and the reference strain (Comas et al., 2010) were recorded as SNPs. The genetic diversity of rv3377c and rv3378c in the MTB complex was done by multiple alignments of the orthologous sequences of both genes retrieved from the genome of 432 strains. CODEML from PAML 3.14b (Yang, 1997) was used to estimate dN/dS rates. The codon frequencies were derived from the genome of 432 strains. CODEML from PAML 3.14b (Yang, 1997) was used to estimate dN/dS rates. The codon frequencies were derived from the genome of 432 strains. CODEML from PAML 3.14b (Yang, 1997) was used to estimate dN/dS rates. The codon frequencies were derived from the genome of 432 strains. CODEML from PAML 3.14b (Yang, 1997) was used to estimate dN/dS rates. The codon frequencies were derived from the genome of 432 strains. CODEML from PAML 3.14b (Yang, 1997) was used to estimate dN/dS rates.

Lipid Extracts of Microbial Species
The chloroform/methanol extraction and sources of mycobacterial, non-mycobacterial, and fungal species used has been described (Ly et al., 2013). In addition, M. marinum, C. albicans, and A. fumigatus were obtained from the American Type Culture Collection.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures. Data S1 and S2, and one table and can be found with this article online at http://dx.doi.org/10.1016/j.chembiol.2015.03.015.

AUTHOR CONTRIBUTIONS
D.C.Y and E.L. performed the normal-phase LC-MS experiments; B.B.S. performed and interpreted NMR on N6-TbAd; Z.W., J.B., and A.J.M. synthesized chemical analogs; S.P., A.T., and J.A. performed analysis of mouse samples; W.B., J.D.E., and D.B.M conceived the overall collaboration and experiments; and D.C.Y and D.B.M wrote the paper.

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