A High-Throughput Screen against Pantothenate Synthetase (PanC) Identifies 3-Biphenyl-4-Cyanopyrrole-2-Carboxylic Acids as a New Class of Inhibitor with Activity against Mycobacterium tuberculosis

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Abstract

The enzyme pantothenate synthetase, PanC, is an attractive drug target in Mycobacterium tuberculosis. It is essential for the in vitro growth of M. tuberculosis and for survival of the bacteria in the mouse model of infection. PanC is absent from mammals. We developed an enzyme-based assay to identify inhibitors of PanC, optimized it for high-throughput screening, and tested a large and diverse library of compounds for activity. Two compounds belonging to the same chemical class of 3-biphenyl-4-cyanopyrrole-2-carboxylic acids had activity against the purified recombinant protein, and also inhibited growth of live M. tuberculosis in manner consistent with PanC inhibition. Thus we have identified a new class of PanC inhibitors with whole cell activity that can be further developed.

Introduction

Nearly one third of the human population is infected with Mycobacterium tuberculosis, the causative agent of tuberculosis (TB) [1]. Despite the existence of approved drug regimens against TB, it continues to claim approximately 1.4 million lives every year [1], and the emergence of increasingly drug resistant strains has made the need for improved therapies more urgent.

A hallmark of M. tuberculosis is its lipid-rich cell wall, which is an essential element of intracellular survival and pathogenicity, and is also thought to contribute to the difficulty of effectively delivering antimicrobial agents into the cell. The significance of this lipid-rich cell wall is underscored by the large number of genes (~250) encoding enzymes in fatty acid metabolism present in the M. tuberculosis genome [2], making this pathway a promising target for new antibacterial drug discovery. Indeed, several anti-tubercular agents are known to inhibit cell wall biosynthesis.

The panC gene encodes the enzyme pantothenate synthetase (PS or PanC), necessary for the production of pantothenate (vitamin B5) in bacteria. Pantothenate is a key precursor for the biosynthesis of coenzyme A (CoA) and acyl carrier protein (ACP), critical components of fatty acid synthesis. The gene encoding PanC is essential for optimal growth in vitro [3], and when genetically disrupted in M. tuberculosis, the resulting strain is auxotrophic, requiring pantothenate supplementation for growth [4,5]. Additionally, pathogenicity is severely attenuated in the pantothenate auxotroph [6]. PanC is absent in mammals, who scavenge pantothenate from their diet using pantothenate permease [7,8], of which there is no homolog in M. tuberculosis. This suggests the potential for developing drugs that do not have cross-reactive toxicity to homologs in the host, and makes PanC an attractive drug target in M. tuberculosis.

PanC catalyzes the ATP-dependent condensation of pantoate and 6-alanine to form pantothenate, simultaneously releasing AMP and pyrophosphate [9]. Previously described assays of PanC activity coupled AMP production to an enzyme cascade that results in the oxidation of NADH to NAD+ in reactions catalyzed by myokinase, pyruvate kinase and lactate dehydrogenase [9]. An enzyme-based screen of M. tuberculosis PanC (PanCMyc) against a library of 4080 compounds identified a weak in vitro inhibitor with...
no observable whole cell activity [10]. However, because this assay relied on a kinetic measurement, and because absorbance measurements of NADH at 340 nM are often complicated by auto-fluorescence in a compound library, we chose to adapt this assay for high throughput screening. The kinetic assay was modified to generate a fluorescent signal that can be measured as a single time-point (end point assay). We used this assay to conduct a high-throughput screen against a large and diverse compound library, and identified several novel inhibitors of PanCMTB, some of which are active against live M. tuberculosis.

**Materials and Methods**

**Chemicals**

Chemicals were obtained from Sigma Aldrich, unless otherwise noted. Pantoate was synthesized as previously described [9]. Myokinase (M3003), pyruvate kinase (P1506), and L-lactic dehydrogenase (L2500) all isolated from rabbit muscle, were purchased from Sigma Aldrich.

**Isolation and purification of recombinant PanCMTB (Rv3602c)**

An Escherichia coli expression vector (pET28b+) encoding PanCMTB with an amino-terminal 6X-Histidine tag (Dr. Courtney Aldrich, University of Minnesota) was transformed into E. coli BL21(DE3), grown to mid-log phase and induced with 0.2 mM IPTG at 18°C for 16 hours. A cell lysate was prepared by treatment with lysozyme and sonication; the cleared lysate was applied first to a nickel column to isolate His-tagged proteins, followed by an additional step of purification by size exclusion chromatography (HiTrap and Sepharose 200, Amersham).

PanCMTB, purified to apparent homogeneity by SDS-PAGE, was concentrated to 2–5 mg/mL in 50 mM HEPES, 50 mM NaCl, 5 mM MgCl₂, and 5% glycerol, flash frozen and stored at −80°C until use.

**Low-throughput assay for PanCMTB activity- kinetic NADH depletion**

The activity of recombinant PanCMTB was measured, as previously described [9]. Briefly, compounds or carrier DMSO alone were incubated with PanCMTB, the coupling enzymes, and their reagents for five minutes. The reaction was then initiated by addition of the PanCMTB substrates, pantoate and β-alanine. The reaction was conducted in wells of a black clear-bottomed microplate containing a final volume of 40 μL per well. The final concentrations were 0.4 mM NADH, 10 mM ATP, 1 mM pantoate, 5 mM β-alanine, 1 mM phosphoenol pyruvate (PEP-K), 10 mM MgCl₂, 2% DMSO, 1 μg/mL of PanC, and 18 U/mL each of myokinase, pyruvate kinase and lactate dehydrogenase in 100 mM HEPES pH 7.8. The plate was immediately transferred to a SpectraMax micro-plate reader and the rate of NADH depletion was monitored by measuring the absorbance at 340 nm every 20 s for a total of 20 min.

**High-throughput assay for PanCMTB activity- single-timepoint fluorescence**

The assay was run using a Beckman Coulter Core robotic system. Key components included an ORCA arm, Multimek, Nanoscreen, Victor 2, custom deck chiller and custom plate shuttles. The automated components were controlled and scheduled using SAMI software. Assay results were determined using custom software and managed with Collaborative Drug...
Discovery’s (CDD) Laboratory Information Management System (LIMS).

The assay is shown in Figure 1A. The kinetic NADH depletion assay was initiated as described above. After 30 minutes 10 μL of solution containing fluorescent reagents was added to each well. The final 50 μL reaction contained 12.5 U/mL of diaphorase and 5 mM resazurin. After thorough mixing, the plate was transferred to a micro-plate reader and the level of resulting NADH-dependent resorufin determined by measuring fluorescence (excitation and emission filters set at 560 nM and 590 nM, respectively).

Preparation of compound plates for HTS and for CRC

HTS library sets of individual compounds were plated at 1 mM - diluted first into HEPES and finally into the assay for a final assay concentration (FAC) of 20 μM. For concentration response curves (CRCs), the compounds were plated in single wells at 10 mM and diluted 3-fold across 9 adjacent wells in a row. Serial dilutions were made in 100% DMSO to ensure accurate concentrations of compound across the series. This entire plate was then treated in the same format as the HTS library plates, for FACs ranging from 200 μM-0.01 μM in 2% DMSO (details provided in Text S1 and shown in Figure S3).

Analog Retrieval

Analog retrieval from the Lilly collection was performed by chemical fingerprint similarity search (Tanimoto similarity threshold of 0.8). The molecular composite fingerprints used for this study consisted of four components. First, 2048 bits were derived from linear paths similar to Daylight fingerprints, then 160 bits that had their origins in the MACCS keys. A further 160 bits were derived from the second bit vector, but where bits were set only when repeated features were found. In addition, 8 molecular properties were added (atom count, size of largest ring, number of rings, number of ring atoms, number of aromatic atoms, number of fused ring atoms, number of heteroatoms) [11,12].

Strains

*M. tuberculosis* H37Rv (ATCC 27294) was used as the parent strain in all experiments (also referred to by the abbreviation RvS). PanC-TetON, a strain in which panC (Rv3602c) is under the conditional expression of a tetracycline-inducible promoter [5] was used in this study. The strain was maintained and cultured in the presence of 50 μg/mL of hygromycin, 20 μg/mL kanamycin and 50 μg/mL pantothenate supplement (vitamin B5), unless otherwise specified.

*M. tuberculosis* viability assays

*M. tuberculosis* strains RvS and PanC-TetON were grown to log phase (OD600~0.3), diluted to a final theoretical OD600 of 0.002 and dispensed into a 96-well round-bottom plate (Corning, Acton, MA) in a final volume of 180 μL. To these cells 20 μL of compound diluted in 7H9 and 10% DMSO was added to yield 200 μL (final concentration of 1% DMSO). For each strain, control wells containing no compound were used as a measure of 100% growth, while wells containing a 1:100 dilution of the starting culture were used as a measure of 99% inhibition. RvS was plated in 7H9 containing 0.2% w/v glycerol, 10% v/v OADC supplement (oleic acid, albumin, D-glucose, catalase; Becton Dickinson) and 0.05% w/v Tween 80 (7H9-GAT) with 0.5 μg/mL anhydrotetracycline (ATc). PanC-TetON was plated in 7H9-GAT with 12.5 μg/mL of hygromycin, 6.25 μg/mL kanamycin and 0.5 μg/mL ATc, either in the presence or absence of 50 μg/mL pantothenate. Plates were incubated at 37°C for 6 days, cells were resuspended by pipetting and 20 μL of the total cell mixture was used in a Bac-Titer Glo™ (Promega, Madison, WI) assay of cell viability, as per the manufacturer’s instructions. Luminescence readings were conducted on a FluosStar Omega plate reader (BMG Lab Tech, Cary, NC). Data from dose-response experiments was represented as the percent inhibition compared with the no-drug controls and analyzed with Graphpad Prism™ (San Diego, CA). The MIC<sub>50</sub> for each growth condition was calculated by fitting the data to a non-linear least-squares curve.

Cytotoxicity assay

Cytotoxicity was measured against the African green monkey adult kidney cell line (Vero). Vero cells were plated at 25,000 cells/mL in black 96-well assay plates pre-populated with compound dilutions and controls. Cells were incubated in a humidified 5% CO<sub>2</sub> environment at 37°C for 48 h; intracellular ATP levels were measured using CellTiter-Glo<sup>®</sup> Reagent (Promega, Madison, WI) [13]. Luminescence was measured using a Victor 2 plate reader and percentage inhibition of growth calculated. Results were expressed as toxicity concentration (TC<sub>50</sub>) = concentration of compound required to inhibit growth by 50%.

Results

Assay description

Recombinant PanC<sub>MTB</sub> was expressed and purified in *E. coli*. We confirmed the enzymatic activity of the purified enzyme using the kinetic assay [9] (Figure 1B) and then adapted it to be more amenable to high throughput screening. Rather than a kinetic measurement of NADH depletion the level of NADH was determined by the terminal addition of the enzyme diaphorase and its substrate, resazurin, which is converted into the fluorescent dye, resorufin, in an NADH-dependent reaction. In this end-point fluorescence assay the PanC<sub>MTB</sub> reaction was initiated and then allowed to proceed, depleting the NADH present in solution. After a defined incubation time, while the PanC<sub>MTB</sub> reaction was still progressing linearly, the diaphorase and resazurin were added, initiating the reduction of the dye with the remaining NADH. We titrated the concentrations of diaphorase and resazurin so that the fluorescent signal developed rapidly and demonstrated linear proportionality to the amount of NADH present in solution (Figure 1D). Therefore, the final fluorescence was inversely proportional to the activity of PanC<sub>MTB</sub> (Figure 1C).

In a set of representative experiments, the reaction conducted with varying levels of PanC<sub>MTB</sub> resulted in varied rates of NADH depletion (Figure 1B). The reactions were then coupled to the fluorescence reagents, and levels of the resulting NADH-dependent resorufin measured (Figure 1C). When no PanC<sub>MTB</sub> was present the fluorescence signal was high. When increasing amounts of PanC<sub>MTB</sub> were present, the rate of AMP production rose, causing progressive depletion of NADH in the initial reaction, ultimately resulting in a drop in the final fluorescence (Figure 1C). In addition to determining assay conditions under which the fluorescence readout was proportional to levels of active PanC<sub>MTB</sub>, we confirmed that the assay was also proportionately sensitive to varied levels of the substrate, pantoate, by holding the concentration of recombinant PanC<sub>MTB</sub> constant, but varying the concentrations of pantoate (data not shown). After successfully coupling the kinetic NADH depletion assay to an end-point fluorescent reaction, we optimized it for translation to an HTS format and tested the robustness of this system to confirm that it was HTS-compatible.
Development and validation for HTS compatibility

All validation steps and the final HTS were performed in 384-well plates using a 320 array of compounds (Figure 2A) with the two outside columns on each side of the plate (columns 1, 2, 23 and 24) reserved for appropriate controls, and the central 20 columns for 320 wells of test compounds. As a negative control reaction, to mimic 100% inhibition, we compared reactions conducted without substrate or without enzyme; there was no significant difference between the two. The substrate-free negative control reaction was simpler to incorporate into the HTS flow deck layout/geometry and was chosen as the “maximum inhibition” control. Two different concentrations of control inhibitor (nafonyl oxalate) that consistently displayed ~50% and ~80% inhibition of the reaction were placed in the two outer most columns of the plate (1 and 24). The positive controls (full reactions with no inhibition) and negative controls were contained in columns 2 and 23, respectively, in order to minimize any chance of edge-effects. Each of the four controls was performed in 16 wells per plate, a number of replicates that provided powerful statistical evaluation. We confirmed that the reagents were stable in the assay conditions (Text S1). In order to minimize the auto-oxidation of NADH we used a custom-chilling unit maintained at 4°C as a reservoir for NADH-containing solutions. All solutions were freshly prepared every four hours.

Various parameters in the assay were developed and optimized, summarized below and detailed in the Text S1. We created individual programs for each of the separate liquid additions in this assay. This included optimization of the liquid handling steps for each of the robots; adjusting the heights, aspiration and dispensation speeds as well as other protocol details including mixing reagents (Figure S4); and washing of tips. Synchronizing the NADH depletion initiation and the fluorescence generating reaction as tightly as possible reduced variation across a plate. Finally we used the appropriate optic filters in the Victor2 Wallac plate-reader that allowed us to obtain optimal excitation and emission of resorufin with minimal spectral overlap from the light source, and minimal auto fluorescence (details in Text S1).

Validation of HTS compatibility was performed based on NCGC guidelines [14] using the 320-array format described earlier (Figure S1). After confirming low variation of signal across

![Figure 2](https://example.com/figure2.png)

Figure 2. Design and implementation of HTS campaign. A) Plate layout for 320-array with four sets of controls (n = 16 for each) in columns 1, 2, 23, and 24. B) Results from Blank Plate Validation using DMSO alone in all 320 assay wells.
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the 320 test wells with uninhibited reactions (Figure 2B, % CV = 2.27) we tested three different concentrations of the control inhibitor, nafronyl oxalate. (2.5 μM, 250 μM and 750 μM) that resulted in a minimum (min), medium (mid) and maximum (max) level of inhibition. Two replicate plates of each were run on a single day, and repeated on three separate days (for a total of six plates). The resulting % CV across each of the plates ranged from 3.7%–12.9% (Figure S1), acceptable for continuing with the HTS.

We performed a final validation of HTS-compatibility by testing a small subset of the LISSP4 library. Three compound plates, each containing 320 compounds, were randomly selected and each was assayed at 5 μM, 10 μM and 20 μM (Figure S2). In addition a 10th plate with no compounds in the 320-array was run alongside these plates. We found that 20 μM showed good levels of inhibition with low background signal. We also saw dose-dependent inhibition from one well, indicating that the pilot screen was successful (Figure S2). The Z’ factor [15] for the controls of each of the three plates tested at 20 μM were 0.803, 0.804 and 0.824 confirming that the assay performance was robust at this concentration.

Screening LISSP4 Library (27.5 K compounds) and Diversity Library (62.6 K compounds)

Having determined that the fluorescently coupled end-point assay was compatible with high-throughput screening, we proceeded to screen two large compound libraries derived from the Eli Lilly screening collection of >800 K physical samples. The library can be subdivided into a set representing compounds previously identified as actives against human targets (LISSP4 – Lilly Strategic Screening Paradigm 4th iteration) and a set of generally diverse compounds (Diversity 4th iteration) not identified as actives at the time of library construction. LISSP4 contains a representative set of 27582 molecules from active (typically <1 μM) classes of druggable human proteins: proteases, kinases, phosphodiesterases (PDE), GPCRs, nuclear hormone receptors, transporters, etc. Diversity4 contains 62651 molecules representative of compounds from the remainder of the library. The sets were created in November 2008.

The LISSP4 library (27,582 compounds) was screened at a final assay concentration of 20 μM (Figure 3A). Two compounds which showed maximum inhibition were identified as promiscuous inhibitors and were removed from further studies. One compound inhibited 41.1% and eleven compounds inhibited >20%. Many compounds showed modest inhibition (10–20%). Although these had relatively low inhibition in the primary assay a subset of these were considered significant because they inhibited ≥3 standard deviations away from the mean inhibition across that particular plate.

Although the results from the LISSP4 library confirmed that the assay could identify inhibitors of PanC activity, this library yielded a relatively low hit rate. Therefore, we decided to screen a more diverse compound set, the Diversity library (62,651 compounds) at 20 μM (Figure 3A). Hits were selected as those with inhibition >15% and at least 3 standard deviations higher than the background within that plate. There were more than 180 hit compounds including 6 at >40% and 3 at >60% inhibition. In total there were 222 primary hits from both libraries.

Confirmation of PanC<sub>MTB</sub> inhibitory activity

We confirmed our primary hits using two assays. The first was a counter-screen to identify compounds whose activity is due to inhibition of one of the coupling enzymes in the reaction. The second was to run concentration response curves (CRCs).

In addition to the 222 primary hits, the seven most potent primary hits were used as seeds for analog retrieval from the Lilly collection. These seven primary hits comprised 5 distinct classes of compounds; three compounds belonged to the same class. Analog retrieval was performed by chemical fingerprint similarity search and supplemented by substructure searches of identified MedChem Studio scaffold (C2) (Simulation Plus Inc.). This resulted in an additional 78 compounds that were assayed. Of the 300 compounds run against the secondary assays 27 compounds were positive in counter-screen (>10% inhibition), indicating that they targeted one or more of the coupling enzymes in the assay. They were eliminated from further analyses.

The remaining 273 compounds were tested in CRCs against PanC<sub>MTB</sub>. Compounds which were active in CRCs (below 200 μM) were grouped into classes using MedChem Studio scaffold clustering; one class of particular interest was identified. This class contained three active compounds from the primary screen and two analogs retrieved by similarity searches. The two analogs (compounds 1 and 2) had EC<sub>50</sub> of 1.8 +/- 1.1 μM and 4.0 +/- 1.1 μM respectively (Figure 3B and C). Compounds of this class did not inhibit the counter screen, and showed dose-dependent inhibition of the PanC<sub>MTB</sub>. The most potent of the four primary hits from this class inhibited PanC in the HTS assay by 71.6% at 20 μM. Tanimoto similarity between the two exemplified hits (1 vs. 2) in the fingerprint used in this work space was 0.81.

Characterization of biochemical enzyme inhibition

We used the low-throughput continuous PanC<sub>MTB</sub> enzyme assay to determine the mode of inhibition, holding all reagents constant but varying a single substrate. Reaction rates were measured as the rate of NADH oxidation and the data were fitted to nonlinear regressions with GraphPad Prism to generate Michaelis-Menten plots (Figures 4A and B). In addition, inhibition constants were determined by nonlinear regression analyses using the general equation of mixed inhibition in GraphPad Prism™, where the resulting parameter “a” determines the mode of inhibition [16]. Both compounds showed non-linear fits closer to competitive inhibition with respect to the substrate pantoate, generating α = 128 and α = 78 for 1 and 2, respectively. Ki inhibition constants with respect to pantoate were 174 +/- 20 nM for 1 and 297 +/- 37 nM for 2.

Whole-cell activity against M. tuberculosis

The compounds were tested for growth inhibitory activity against wild-type M. tuberculosis by monitoring ATP levels in cultures exposed to compounds. In each growth condition, bacteria grown in the absence of drug were used as a control reflecting 100% growth. We found that both compounds 1 and 2 were active against live M. tuberculosis, with MIC<sub>50</sub> values of 115 μM and 54 μM respectively (Figure 5). Further, we wanted to ascertain the degree to which this activity could be attributed to the specific inhibition of PanC<sub>MTB</sub> in whole cells. To do this, we tested the activity of compounds against a conditional PanC-knockdown strain (PanC Tet-ON<sub>TM</sub>) in which PanC expression is held under the control of a tetracycline-inducible promoter; in the presence of low concentrations of tetracycline <i>pantC</i> expression is dramatically reduced [5] and growth is partially attenuated. We used these conditions to maintain growth, while still having artificially lowered PanC levels, making the cell line more sensitive to PanC-mediated inhibition. We found that under lowered <i>pantC</i> expression the MIC<sub>50</sub> of both 1 and 2 dropped about two-fold (Figure 5). The MIC<sub>50</sub> of 1 was reduced from 115 μM in wild-type to 69 μM in the PanC under-expressor strain. Similarly 2 was more potent against the PanC under-expressor strain with an MIC<sub>50</sub> of 24 μM compared to 54 μM in the wild-type. This
suggested that the growth defect is linked to PanC-mediated inhibition. In the presence of higher tetracycline levels panC is induced in the PanC Tet-ON\textsubscript{M} strain but this only partially alleviates the growth defect achieving \textasciitilde 80% of wild-type growth. However, with supplemental pantothenate the pressure caused by the PanC deficiency is relieved and growth is restored to wild-type levels [5]. Therefore to confirm that the hypersensitivity to the PanC inhibitors was not due simply to the attenuated growth, the compounds were also tested against PanC Tet-ON\textsubscript{M} grown in the presence of pantothenate. As expected, the increased sensitivity was abolished when pantothenate was included in the growth medium, with 1 and 2 displaying MIC\textsubscript{50} values similar to those in wild type RvS (111 and 53 \textmu M, respectively; Figure 5).

The two compounds were tested for cytotoxicity against the Vero cell line; both showed some toxicity with TC\textsubscript{50} of 70 \textmu M for compound 1 and 96 \textmu M for compound 2. We also tested for...
Detailed information on tip handling during HTS, compound plating for concentration response curves, reagent stability testing, robotics optimization, and optics for resorufin measurement.

**Figure S1** Validation of HTS-compatibility assaying plates with max, mid, and min concentrations of a control inhibitor.

(PDF)

**Figure S2** Data generated in pilot study from one compound plate assayed at three different concentrations.

(PDF)

**Figure S3** Plate layout for concentration response curves (CRCs).

(PDF)
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Conceived and designed the experiments: AK AC JO EAK GA MV TM PAH DRS TP. Performed the experiments: AK AC JO EAK. Analyzed the data: AK AC JO EAK MV TM PAH DRS TP. Contributed reagents/materials/analysis tools: AK AC JO EAK GA MV TM PAH DRS TP. Wrote the paper: AK AC MV DRS TP.

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