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New insight into structure-activity of furan-based salicylate synthase (MbtI) inhibitors as potential antitubercular agents

Laurent R. Chiarelli[a]* iD, Matteo Mori[b]* iD, Giangiacomo Beretta[c] iD, Arianna Gelain[b] iD, Elena Pini[b] iD, Josè Camilla Sammartino[a] iD, Giovanni Stelitano[a] iD, Daniela Barlocco[b] iD, Luca Costantino[d] iD, Margherita Lapillo[e], Giulio Poli[e] iD, Isabella Caligiuri[f] iD, Flavio Rizzolio[f,g] iD, Marco Bellinzoni[h] iD, Tiziano Tuccinardi[e] iD, Stefania Villa[b] iD and Fiorella Meneghetti[b] iD

[a]Department of Biology and Biotechnology "L Spallanzani", University of Pavia, Pavia, Italy; [b]Department of Pharmaceutical Sciences, University of Milano, Milano, Italy; [c]Department of Environmental Science and Policy, University of Milano, Milano, Italy; [d]Department of Life Sciences, University of Modena e Reggio Emilia, Modena, Italy; [e]Department of Pharmacy, University of Pisa, Pisa, Italy; [f]Pathology Unit, Department of Molecular Biology and Translational Research, National Cancer Institute and Center for Molecular Biomedicine, Aviano, Italy; [g]Department of Molecular Science and Nanosystems, Ca' Foscari University of Venezia, Venezia-Mestre, Italy; [h]Institut Pasteur, Paris, France

ABSTRACT
Starting from the analysis of the hypothetical binding mode of our previous furan-based hit (**I**), we successfully achieved our objective to replace the nitro moiety, leading to the disclosure of a new lead exhibiting a strong activity against MbtI. Our best candidate **1h** displayed a K_i of 8.8 μM and its antimycobacterial activity ($MIC^{99} = 250$ μM) is conceivably related to mycobactin biosynthesis inhibition. These results support the hypothesis that 5-phenylfuran-2-carboxylic derivatives are a promising class of MbtI inhibitors.

GRAPHICAL ABSTRACT
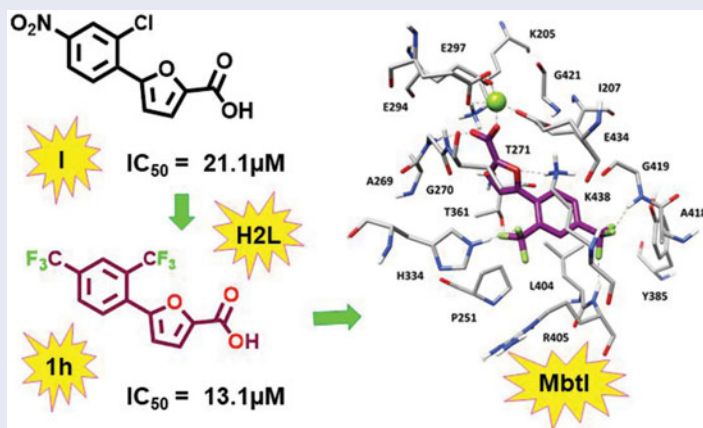
Introduction

Tuberculosis (TB) is an infectious disease caused by an obligate aerobic bacterium, known as *Mycobacterium tuberculosis* (Mtb). When the bacilli are inhaled, they reach the alveolar spaces of the lungs and are mainly ingested by macrophages. As a consequence, TB primarily affects the lungs, but at later stages it can also spread to other vital organs. Despite significant improvements with respect to diagnosis, treatment, and preventive measures have been successfully implemented in many healthcare systems around the world, this disease still remains the world's biggest threat to human health causing 54 million deaths between 2000 and 2017[1]. Standard therapeutic regimens have remained substantially unchanged over the past 60 years with outdated drugs and very long therapies that are still used for the treatment of new and relapse cases. In addition to the length of the cure, other hurdles related to the management of TB infections include interactions with drugs used in comorbid conditions, especially HIV, and severe side effects. All these issues contribute

Table 1. *In vitro* activity of compounds **1a–p**.

1a–i R$_1$=CF$_3$				1j–p R$_1$=NO$_2$			
Code	R$_2$	Residual activity at 100 μM (%)	Mbtl IC$_{50}$ (μM)*	Code	R$_2$	Residual activity at 100 μM (%)	Mbtl IC$_{50}$ (μM)*
1a	Cl	6.1 ± 2.5	28.5 ± 2.6	I**	Cl	11.3 ± 4.2	17.9 ± 3.2
1b	F	6.1 ± 1.1	27.6 ± 7.3	1j	F	32.1 ± 2.4	–
1c	Br	38.0 ± 3.7	–	1k	Br	26.4 ± 4.8	–
1d	OH	16.8 ± 2.7	35.9 ± 10.3	1l	OH	13.9 ± 4.0	29.8 ± 4.2
1e	CH$_3$	19.3 ± 2.2	31.5 ± 9.6	1m	CH$_3$	13.2 ± 5.8	28.5 ± 1.5
1f	NH$_2$	13.1 ± 2.4	34.6 ± 10.9	1n	NH$_2$	13.5 ± 3.8	24.2 ± 5.4
1g	CN	5.7 ± 1.5	18.5 ± 3.2	1o	CN	5.0 ± 1.9	24.4 ± 5.9
1h	CF$_3$	3.9 ± 1.7	13.1 ± 2.0	1p	CF$_3$	25.1 ± 3.7	41.8 ± 5.3
1i	*m*-CF$_3$	64.3 ± 4.5	–	–	–	–	–

*Only for compounds with residual activity ≤25%; **new fluorimetric assay value determined for the sake of comparison.

to determine a poor patient compliance that, together with the improper use of TB antibiotics, has led to the insurgence of multi-drug-resistant (MDR) and extensively drug-resistant (XDR) bacterial strains[2–4]. As confirmed by the latest WHO report, drug resistance is becoming a real emergency; therefore, there is a growing interest in the development of novel anti-TB compounds[5–8]. A few of them reached clinical trials and two drugs, delamanid and bedaquiline, have been recently approved; however, more information on their effectiveness, safety, and tolerability are still required because severe side effects have been reported[9,10]. In this scenario, the research of many more TB drug candidates to sustain an effective and productive drug pipeline is pivotal. Targeting Mtb iron uptake systems is now a validated strategy for the development of antimycobacterial compounds, because iron is essential for Mtb survival in the host and its acquisition is strongly correlated with virulence[11]. Among the four different iron acquisition pathways used by Mtb, the most thoroughly characterised one is based on the production of two types of siderophores: carboxymycobactins, which acquire iron extracellularly and transport it into the cytoplasm of the bacteria, and mycobactins, which facil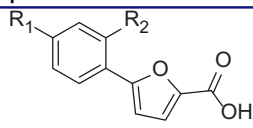itate the transport of iron through the cell wall into the cytoplasm. Notably, targeting this biosynthetic process is an attractive strategy, because its impairment lowers the pathogen virulence and survival without causing toxicity issues. Indeed, as this pathway is absent in humans, the risk of off-target effects is minimal; moreover, being an unexplored biological process for the development of drugs, there is no known resistance mechanism. The first step of the biosynthesis of these siderophores is catalysed by the Mg^{2+}-dependent enzyme salicylate synthase (Mbtl), a validated pharmacological target[12–15], whose crystal structure has been recently solved[16].

In this context, the aim of our project is the identification of new Mbtl inhibitors as potential antitubercular agents. Our previous computational studies generated a pharmacophore model, that allowed the identification of the interesting hit compound **I** (Table 1). Then, a structure-activity relationship study on monosubstituted derivatives underlined the importance of the nitro moiety for potency[14]. However, the nitro group is considered a structural alert for the development of a potential drug, since drugs containing nitro groups have been extensively associated with mutagenicity and genotoxicity[17]. On these bases, in the present work, we designed additional analogs (compounds **1a-p**, Table 1) exploring other hitherto unconsidered pharmacophoric features and evaluating the possibility of replacing the nitro group.

Materials and methods

Chemistry

Compounds **1a,b,d–i,l–n,p** were synthesised by a Suzuki-Miyaura reaction[18], followed by a base-catalysed hydrolysis of the ester function[19]. Compounds **1c,j,k,o** were synthesised by a Sandmayer reaction[20], starting from 2-furan carboxylic acid methyl ester and the diazonium salts of the appropriate amines; the ester function was then hydrolysed in basic conditions[21]. The general procedures, the synthetic pathways (Supplementary Schemes 1 and 2), the details concerning the specific synthetic steps and the analytical data are provided in the Supplementary Material.

Mbtl enzymatic assays

Recombinant *M. tuberculosis* Mbtl was produced and purified as previously reported[14]. Enzyme activity was determined measuring the production of salicylic acid by a fluorimetric assay slightly modified from Vasan et al.[12] Briefly, assays were performed at 37 °C in a final volume of 400 μL of 50 mM Hepes pH 7.5, 5 mM MgCl$_2$, using 1–2 μM Mbtl and the reactions were started by the addition of chorismic acid and monitored using a Perkin-Elmer LS3 fluorimeter (Ex. $\lambda = 305$ nm, Em. $\lambda = 420$ nm). Inhibition assays were performed in the presence of the compound at 100 μM (stock solution 20 mM in DMSO), at 50 μM chorismic acid. For significantly active compounds, IC$_{50}$ and K_i values were determined. To verify that the compounds were not PAINs, their ability to inhibit Mbtl activity was tested in the presence of 0.1 mg/mL of bovine serum albumin (BSA) or in the presence of 0.01% (v/v) Triton X-100 to confirm that they did not act as aggregate, and in the presence of 100 mM of 1,4-dithio-DL-threitol (DTT) to exclude an enzyme inhibition due to reaction with cysteines[22].

Minimal inhibitory concentration determinations and siderophore production assay

The MIC^{99} values of active compounds against *M. bovis* BCG were determined in low-iron Chelated Sauton's medium[23], using the resazurin reduction assay (REMA)[24]. Siderophore activity in the culture was tested in *M. bovis* BCG using the Universal CAS liquid assay. To this purpose, *M. bovis* was grown in 7H9 medium, subcultured in chelated Sauton's medium and then diluted to an
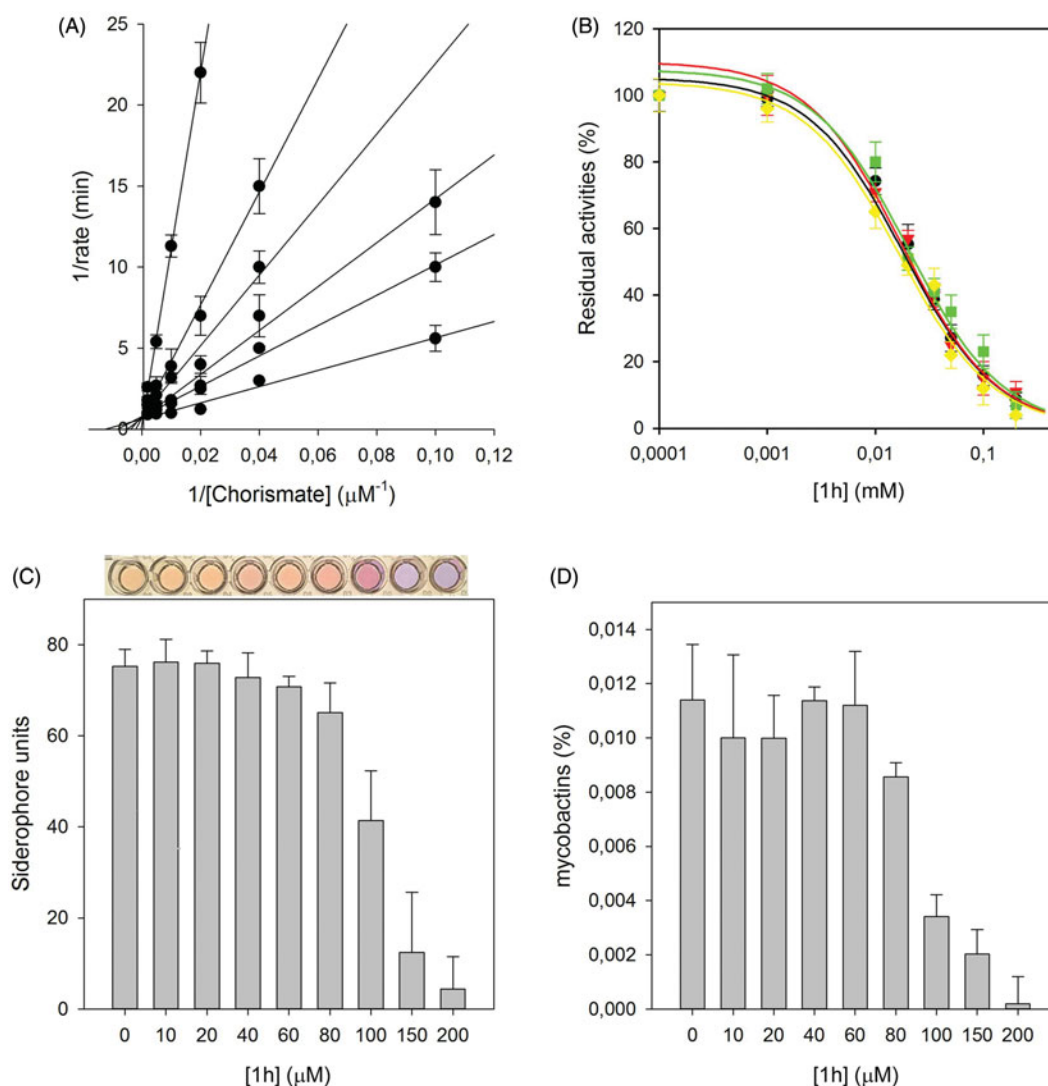
Figure 1. Biological characterisation of **1h**. The global reciprocal plot of data from MbtI steady-state kinetics analysis towards chorismic acid, in the presence of different concentrations of **1h** (A), highlights the competitive behaviour of the inhibitor. IC_{50} plots of **1h** in the presence of BSA (green squares), Triton X-100 (red triangles) and DTT (yellow diamonds) (B) confirm that the compound is not a PAIN. The Universal CAS assay performed on culture media of *M. bovis* BCG cells grown in the presence of **1h** (C), together with the determination of the mycobactins in the above-mentioned cells (D), confirm that the antimycobacterial activity is related to iron uptake inhibition. All data are mean ± SD of three replicates.

OD_{600} of 0.01 in chelated Sauton's containing different concentrations of the test compound. After 15 days of incubation at 37 °C, cells were harvested, supernatants were used to perform CAS assay and cell pellets were used for the determination of mycobactins. For CAS assay, an aliquot of 100 μL of supernatant was mixed with 100 μL of CAS assay liquid solution in a 96-well plate, incubated for 10 min at room temperature, and the absorbance was measured at 630 nm. For mycobactin determination, cell pellets were extracted in ethanol overnight, then 0.1 M $FeCl_3$ in ethanol was added until no color change was observed. The mixture was incubated at room temperature for 1 h, then mycobactins were extracted in chloroform, washed with water, evaporated and the residue was dissolved in methanol. The concentration of mycobactins was determined measuring the absorbance at 450 nm (1% solution of mycobactins gives an absorbance of 42.8).

Cell viability assay

MRC5 human fibroblast lung cells were maintained at 37 °C in a humidified atmosphere containing 5% CO_2, according to the supplier's indications. Cells (10^4) were plated in 96-well culture plates. The day after seeding, vehicle or compounds were added at different concentrations to the medium. Compounds were added to the cell culture at a concentration ranging from 200 to 0.01 μM. Cell viability was measured after 96 h according to the supplier's instructions (Promega, cat. n° G7571) with a Tecan M1000 PRO instrument. IC_{50} values were calculated from logistic dose-response curves.

Docking studies

All the compounds were docked into the minimised average structure of MbtI complexed with the lead compound **I**[14]. The software Gold with the four fitness functions implemented (i.e. GoldScore, ChemScore, Astex Statistical Potential, and ChemPLP) and plants were employed in this study as previously reported[14,25], since they have been already used showing good results in the previous virtual screening study on MbtI inhibitors performed in our laboratory[15]. The docking site was defined as the region comprising all residues that stayed within 10 Å from
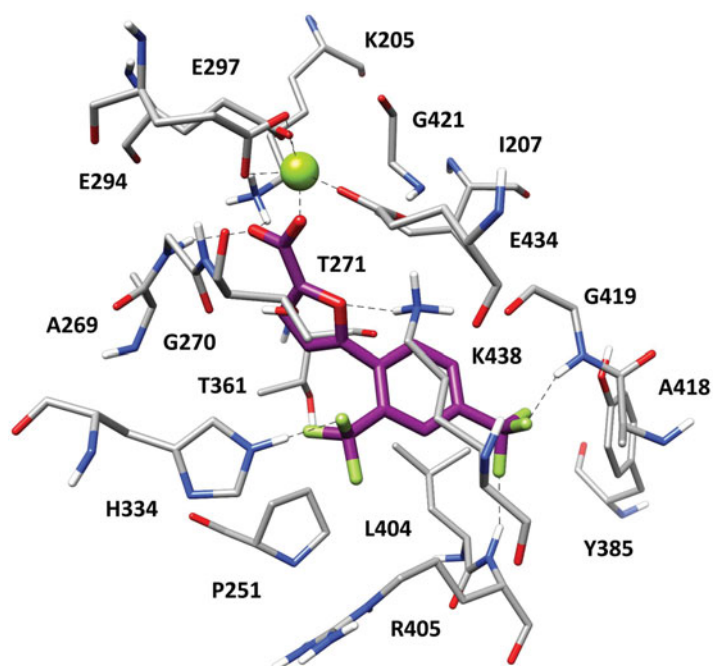
Figure 2. Minimised average structure of **1 h** within Mbtl binding site.

the reference compound **I**. By applying the five docking methods, five different binding dispositions (best-scored docking pose) resulted from the docking of each ligand into the protein binding site. The RMSD of these docking poses against the remaining four was evaluated by using the rms_analysis software of the Gold suite. On this basis, for each ligand docked into the protein binding site, a 5×5 matrix was generated reporting the RMSD results. By using an in-house programme, these results were clustered, so that among the five results, all of the similar docking poses were grouped together[26]. We selected an RMSD clustering threshold of 2.0 Å; therefore, the so-obtained clusters contained the group of poses that were less than 2.0 Å away from all others poses belonging to the same cluster. For each compound, the binding mode belonging to the most populated cluster identified by the consensus docking evaluation was taken into consideration and subjected to MD simulations.

MD simulations

All simulations were performed using AMBER 16[27]. General amber force field (GAFF) parameters were assigned to the ligands, whereas partial charges were determined using the AM1-BCC method as implemented in the Antechamber suite of AMBER16. MD simulations were carried out employing the ff14SB force field at 300 K. Magnesium ion was inserted considering its disposition and interaction into the 2FN1 PDB code. Ligand-protein complexes were placed in a rectangular parallelepiped water-box and solvated with a 20 Å water cap by using the TIP3P explicit solvent model. Sodium ions were added as counterions in order to neutralise the system. Before MD simulations, two steps of minimisation were performed; in the first stage, a position constraint of 500 kcal/(mol·Å²) was applied to keep the protein fixed thus minimising only water molecules. In the second stage, the whole system was energy minimised through 5000 steps of steepest descent followed by conjugate gradient (CG), until a convergence of 0.05 kcal/(mol· Å²) and imposing a harmonic potential of 10 kcal/(mol·Å²) to the protein α carbon. Particle mesh Ewald (PME) electrostatics and periodic boundary conditions were used

in the simulations. The time step of the simulations was 2 fs with a cutoff of 10 Å for the non-bonded interactions, while the SHAKE algorithm was applied to keep all bonds involving hydrogen atoms fixed. Constant-volume periodic boundary MD simulation was carried out for the first 0.5 ns, during which the temperature of the system was raised from 0 to 300 K. Then 19.5 ns of constant-pressure periodic boundary MD was performed at 300 K, using the Langevin thermostat in order to maintain constant the temperature of the system. A harmonic force constraint of 10 kcal/(mol· Å²) was applied to the protein α carbons during the first 3.5 ns, whereas in the last 16.5 ns no restraints were applied to the system. All the obtained MD trajectories were analyzed using the Cpptraj programme implemented in AMBER 16.

Result and discussion

In order to further develop our library of furan-based candidates, we examined the substitution pattern at the *para* and *ortho* positions of the phenyl ring. Herein, we disclose additional SAR data; firstly, we synthesised a phenyl di-substituted analog of **I**, characterised by the presence of the trifluoromethyl group (**1a**), because we knew that the more druggable *p*-CF₃ in place of the *p*-NO₂ group was still able to inhibit Mbtl[14]. Compound **1a** displayed encouraging inhibiting properties (residual enzyme activity at 100 μM, as % RA, of 6.1 ± 2.5, IC₅₀ = 28.5 ± 2.6 μM). Therefore, we took into account this compound as a new hit and we focussed on the effects of a variety of electron donating/withdrawing or hydrophilic groups in the *ortho* position of the phenyl ring, keeping the CF₃ group in *para* position (**1b–i**). Then, the effects of the previously analysed substituents in *ortho* position were evaluated in the second series of derivatives **1j–p**, bearing the original *p*-NO₂ group (Table 1).

The activity of all compounds (**1a–p**) was tested against the recombinant Mbtl, prepared and assayed as previously reported[14]. The substitution of the chlorine atom with the fluorine in **1 b** did not affect the activity, while the introduction of the bromine atom in **1c** diminished the inhibitory effect of the compound (% RA 38.0 ± 3.7), due to the low capability of bromine to act as

hydrogen bond acceptor together with its higher atomic radius determining a negative steric constraint. The presence of the electron donating groups in **1d**, **1e**, and **1f** did not lead to an improvement of the biological effects. When an electron withdrawing moiety was introduced in the *ortho* position, as in **1 g**, we detected an increased activity. This outcome was confirmed by the introduction of an *o*-CF$_3$ moiety that led to the disclosure of **1 h**, having an IC$_{50}$ value comparable to that of the most active inhibitor (**I**), previously identified. Derivative **1i** was then prepared and tested to assess the importance of having the electron withdrawing moiety in *ortho* position of the phenyl ring; interestingly, moving the CF$_3$ to the *meta* position significantly decreased the inhibitory activity of the compound. A parallel trend of biological results was obtained for the *p*-NO$_2$ derivatives **1j–p**. These compounds were prepared and tested to compare the impact of the CF$_3$ and NO$_2$ groups on the activity and to better understand their role in the interaction with the target. The outcomes reflected the similar biological behaviour observed for this series (**1j–p**) with respect to the previous one (**1a–i**), confirming the ability of CF$_3$ to act as a bioisostere of the nitro moiety, according to the bioisosterism between fluorine and carbonyl functionality[28]. Of note, the presence of two CF$_3$ substituents in **1 h** synergically contributed to enhance the activity, as shown by the comparison of the IC$_{50}$ value of **1 h** and the respective derivative **1p** belonging to the *para*-nitro series (13.1 vs. 41.8 μM).

For the most active compound, **1 h**, an accurate biological analysis was thus performed (Figure 1). Kinetic analysis allowed the determination of the K_i value 8.8 ± 0.7 μM, allowing the classification of this compound as competitive inhibitor of MbtI. Then, additional tests were performed to verify that **1 h** was not a PAIN compound. The addition of BSA and Triton X-100 did not influence its IC$_{50}$, proving that the compound did not form aggregates with the target. Similarly, the addition of DTT did not impact on **1 h** activity, showing that the ligand did not interact with the cysteine residues of the protein. To evaluate the antimycobacterial activity of **1 h** we used the nonpathogenic *M. bovis* BCG, whose siderophores closely resemble Mtb mycobactins[29,30]. The minimal inhibitory concentration (MIC^{99}) value, determined in iron-limiting conditions in chelated Sauton's medium, was similar to that of hit **I** (250 μM versus 500 μM). Moreover, to assess if the antitubercular activity was indeed related to iron uptake inhibition, the effects of the compound on siderophore production were evaluated by means of the Universal CAS liquid assay[30]. Firstly, the level of siderophore activity in the medium was measured and the colorimetric assay revealed that the removal of iron was inversely related to the concentration of the compound. Similarly, the quantification of mycobactins isolated from the cell pellet evidenced that the concentration of mycobactins decreased at higher **1 h** concentrations. All these outcomes suggested that the inhibitory effect towards Mtb growth was due to mycobactin biosynthesis inhibition. Finally, compounds **I**, **1 h** and **1p** were selected for evaluating their antiproliferative activity against human MRC-5 fibroblasts. The biological results showed that the three compounds did not affect the growth of these normal cells (MRC-5, IC$_{50}$ > 100 μM), thus indicating the potentially low level of toxicity of this class of compounds.

With the aim of elucidating the binding mode of this class of derivatives, molecular modelling studies were performed on the most active compound of the series. Compound **1 h** was subjected to a consensus docking protocol and the result was refined through 20 ns of MD simulation with explicit water molecules (see Supplementary Material for details). Figure 2 shows the minimised average structure of **1 h** bound to the catalytic site of MbtI. The two aromatic rings of the ligand lay on the hydrophobic wall constituted by I207, P251, T361, and L404, thus forming lipophilic interactions with these residues. On the other side of the binding site, the ligand core moiety shows multiple interactions with K438: a hydrogen bond formed by the furan oxygen and a cation-π interaction established through the phenyl ring, which also forms additional hydrophobic contacts with the side chain of the residue. The carboxylic group of **1 h** coordinates the Mg^{2+} ion and also forms H-bonds with the backbone nitrogen of G270 and the side chain of K205 that contribute to anchor the ligand to the MbtI binding site. According to the experimental results, the *p*-CF$_3$ group of the ligand replaces the *p*-NO$_2$ group of the parent compound **I** by forming two H-bonds with the backbone nitrogen of R405 and G419, while the *o*-CF$_3$ substituent of **1 h** shows an additional hydrogen bond with the side chain of H334. The same consensus docking and MD protocols were then applied on compounds **1a** and **1q** to gain a better interpretation of the obtained SAR. In agreement with the experimental data, the binding mode predicted for **1a** confirmed the importance of the *o*-CF$_3$ substituent of **1 h**. In fact, the *o*-chlorine atom of **1a** does not show any hydrogen bond with H334. Moreover, although the ligand perfectly chelates the Mg^{2+} ion, the presence of the chlorine atom determines a small shift in the binding orientation that hampers the formation of H-bonds with G270 and K205 predicted for **1 h** (Figure S1). Nevertheless, the presence of an *o*-CF$_3$ moiety was found to be detrimental when associated with the *p*-NO$_2$ group originally present in **I**, as observed for compound **1p**. Interestingly, MD simulations suggested that **1p** is not able to form the H-bond and cation-π interactions with K438 (Figure S2). In order to establish the H-bond network among its *p*-NO$_2$ group, the backbone nitrogen of R405 and the side chain of Y385, **1p** moves toward these residues and slightly rotates toward K438, which thus moves away from the ligand to avoid steric clashes. Due to its orientation within MbtI binding site, **1p** cannot even interact with H334 through the *o*-CF$_3$ as predicted for **1h**.

To conclude, starting from the analysis of the hypothetical binding mode of our previous furan-based hit **I**, we successfully performed the bioisosteric replacement of the nitro group. Our preliminary hit optimisation study led to the disclosure of a new compound (**1 h**) exhibiting a strong activity against MbtI, comparable to the most potent competitive inhibitor reported so far[14]. Our best candidate **1 h**, characterised by the presence of two CF$_3$ substituents in the *ortho* and *para* positions of its phenyl ring, displayed an IC$_{50}$ value of 13.1 ± 2.0 μM (K_i = 8.8 ± 0.7 μM) and the antimycobacterial activity showed by this compound (MIC^{99} = 250 μM) is conceivably related to mycobactin biosynthesis inhibition. Moreover, preliminary assays against noncancerous human fibroblast lung cells did not reveal cytotoxicity issues. These results support the hypothesis that 5-phenylfuran-2-carboxylic derivatives are a promising class of MbtI inhibitors and allowed us to gather new information for a further optimisation of this class of compounds.

Disclosure statement

Funding

ORCID

Laurent R. Chiarelli http://orcid.org/0000-0003-0348-9764
Matteo Mori http://orcid.org/0000-0002-7491-1494
Giangiacomo Beretta http://orcid.org/0000-0003-0987-0857
Arianna Gelain http://orcid.org/0000-0001-9104-732X
Elena Pini http://orcid.org/0000-0003-0554-566X
Josè Camilla Sammartino http://orcid.org/0000-0003-3707-3118
Giovanni Stelitano http://orcid.org/0000-0002-5219-4770
Daniela Barlocco http://orcid.org/0000-0002-8593-1853
Luca Costantino http://orcid.org/0000-0001-5334-8084
Giulio Poli http://orcid.org/0000-0002-8061-5632
Isabella Caligiuri http://orcid.org/0000-0002-4355-5597
Flavio Rizzolio http://orcid.org/0000-0002-3400-4363
Marco Bellinzoni http://orcid.org/0000-0002-8887-6917
Tiziano Tuccinardi http://orcid.org/0000-0002-6205-4069
Stefania Villa http://orcid.org/0000-0001-6616-1163
Fiorella Meneghetti http://orcid.org/0000-0002-6511-7360

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