Combining β-Lactam Antibiotics Currently in Clinical Trials Are Efficacious in a DHP-I-Deficient Mouse Model of Tuberculosis Infection


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We report here a dehydropeptidase-deficient murine model of tuberculosis (TB) infection that is able to partially uncover the efficacy of marketed broad-spectrum β-lactam antibiotics alone and in combination. Reductions of up to 2 log CFU in the lungs of TB-infected mice after 8 days of treatment compared to untreated controls were obtained at blood drug concentrations and time above the MIC (T₅₀/MIC) below clinically achievable levels in humans. These findings provide evidence supporting the potential of β-lactams as safe and mycobacterial components of new combination regimens against TB with or without resistance to currently used drugs.

The widespread assumption that β-lactams, one of the safest Types of antibiotics, are not adequate drugs for the treatment of tuberculosis (TB) infection is grounded on (i) the constitutive production of BlaC, a broad-spectrum class A β-lactamase that protects Mycobacterium tuberculosis from β-lactam action (1), (ii) the impermeable nature of the mycobacterial cell wall, rendering most β-lactams weakly efficacious in vitro (2), (iii) their potentially limited penetration into macrophages, (iv) their failure to show significant levels of efficacy in commonly employed TB animal models of infection (3, 4), and (v) the unconvincing clinical proof of efficacy reported to date (5, 6). These arguments, together with further limitations associated with the lack of active and orally bioavailable β-lactams with long half-lives have profoundly affected the state of opinions in the TB research and development (R&D) community to the point where research in this area has virtually stalled for the past 3 decades.

β-Lactams have a long-standing record of clinical safety and a low potential for interaction with other antibactericidal and antituberculotic compounds. As a new class of antibacterial drugs, β-lactams would be the ideal complement to other novel agents to form a completely new antibacterial treatment regimen for treating multidrug-resistant (MDR) or extensively drug-resistant (XDR) TB. In summary, recently published data have produced the following findings. First, they have shown that the combination of some β-lactam carbapenems with clavulanate is efficacious against both drug-sensitive and MDR-TB strains. Activity was also shown against hypoxic M. tuberculosis cultures. Second, orally bioavailable BlaC-resistant faropenem matches the combination of meropenem-clavulanate in terms of potency and bactericidal behavior against M. tuberculosis (8). Third, a synthetic lethality screen has shown how mycobacteria can become hyper-susceptible to β-lactam action through a number of resistance mechanisms commonly induced by current antibacterial therapy (e.g., cell wall biosynthesis and efflux systems) (9). Last, MDR-TB strains are hypersensitive to the meropenem-clavulanate combination in vitro (10). Additionally, to date, it has not been possible to isolate spontaneous mutants resistant to the combination of meropenem and clavulanate or faropenem under laboratory conditions, hinting at a low propensity for the development of resistance. Finally, an increasing number of clinical reports have shown that the combination of meropenem and clavulanate contributes to sputum conversion and secession of symptoms in MDR/XDR therapeutically destitute TB patients (11-13).

Only recently, a European Development Clinical Trials Partnership (EDCTP)-funded early bactericidal activity (EBA) study on the use of different β-lactam combinations in drug-sensitive TB patients in South Africa was initiated (https://clinicaltrials.gov/ct2/show/NCT02349841) (14). In order to be able to support the previously mentioned meropenem- faropenem-amoxicillin-clavulanate clinical study (14), we chose to focus our efforts on the development of a mouse model of TB infection that could allow a closer reproduction of the human blood pharmacokinetic profiles of β-lactams in mice. The model should be able to address the ubiquitous and high-level expression of dehydropeptidase-I (DHP-I) (15), one of the main enzymes responsible for the cleavage of the lactam ring and hence the extremely quick clearance of β-lactams in vivo.

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β-lactam antibiotics in rodent tissues (including lungs) in comparison to that in humans (16).

The lack of significant efficacy encountered in different animal models of TB infection (4, 5, 17) has been a great source of frustration among TB researchers and is one of the reasons accounting for the very limited number of translational studies published to date in support of an eventual clinical evaluation. Such a model would need to address one of the main limitations associated with the use of rodent species for TB efficacy studies, i.e., the high activity and high level of expression of dehydropeptidase-I, an enzyme that acts as a major contributor to lactam ring cleavage in most tissue types of mice, rats, guinea pigs, and rabbits (16). The model should also be able to sustain both acute and chronic infections and generate significant and robust β-lactam efficacy responses to allow eventual drug-to-drug comparisons. While the model was specifically designed for the purpose of antibacterial drug discovery, such a model has been reported to be deficient in membrane-bound dipeptidase (MBD) (EC 3.4.13.19, also known as DHP-I), an enzyme hypothesized to be responsible for the conversion of leukothriene D4 (LTD4) to leukothriene E4 (LTE4) (18). This same enzyme was also shown to be responsible for the cleavage of cystinyl-bis-glycine (cys-bis-gly), and its null mutation led to the absence of β-lactamase activity in the lungs, kidneys, small intestines, and hearts of mice.

A preliminary evaluation of the model in terms of its capacity to sustain a viable M. tuberculosis H37Rv infection and improve the whole-blood exposure levels of meropenem showed how the progression of disease in terms of lung CFU counts was identical to that in the reference mouse strain C57BL/6 (see Fig. S1 in the supplemental material). Additionally, significant increases of meropenem both in terms of the maximum concentration of drug (Cmax) and area under the concentration-time curve (AUC) (Table 1) were measured. Most important, based on single-dose pharmacokinetic (PK) parameters, the time above the MIC (T>MIC) simulated in the DHP-I knockout (KO) mice increased significantly for the three considered meropenem MIC scenarios (0.23, 0.62, and 1.25 μg/ml) (Fig. 1). Once the principles of reduced β-lactam degradation and viability of the infection in vivo had been validated, a quick and preliminary evaluation of the effectiveness of β-lactam combination treatment to control TB infection in DHP-I-deficient mice was performed in a survival efficacy study (see Fig. S2 in the supplemental material). These combinations were selected while keeping in mind the clinical trial design that was implemented in the ongoing EDCTP-sponsored β-lactam phase IIa study (14) and previous knowledge regarding the in vitro antibacterial activities of their main components (MICs, 0.3, 1.25, 2.5, and 1.25 μ/ml for meropenem-clavulanate, faropenem, cefdinir, and amoxicillin-clavulanate, respectively) (9). The experiment resulted in an increased mean survival time (>39 versus 17.3 days) for β-lactam-treated groups of mice up to the time of final sacrifice (see Fig. S3 in the supplemental material). These experiments, while providing only qualitative evidence of the potential efficacy of β-lactams in a DHP-I-deficient murine model, were used as a basis for further evaluation of the model for the quantitative generation of efficacy signals. This second goal was addressed through the establishment of a new experimental design (see Fig. S4 in the supplemental material). This model showed how different β-lactam combination regimens gave rise to robust and significant CFU reductions in the lungs of infected DHP-I-deficient mice, with faropenem medoxomil containing the most significant reductions (−2 log CFU), followed by meropenem (−1.7 log CFU) and cefdinir (−1.3 log CFU), in comparison to the untreated control (Fig. 2). These results are in contrast to previously reported data for the meropenem-clavulanate combination (4, 5), albeit under different administration schedules. Most importantly, once-a-day treatment with the faropenem medoxomil-containing regimen was also able to produce a 1.3-log CFU reduction, further highlighting the benefits associated with DHP-I deficiency in terms of decreased pharmacokinetic clearance (Table 1). This observation also addresses one of the shortcomings thought to limit the potential of β-lactams for the treatment of human TB, i.e., the likely requirement for multiple daily administra-

![FIG 1 Whole-blood pharmacokinetic profiles simulated from a monocompartmental model after fitting of single-dose profiles of meropenem in TF3157 (a) and C57BL/6 (b). (c) A bicompartamental model was used for TF 3157 DHP-I KO mice. conc, concentration.](image-url)

FIG 2  The efficacy response was quantified by measuring the bacterial load in the lungs of infected mice for different combinations of β-lactams in a DHP-I KO acute TB model. Drugs administered subcutaneously were prepared in phosphate-buffered saline (PBS) buffer (300 mg/kg of body weight meropenem three times a day [t.i.d.]). Drugs administered orally were prepared in 1% methylcellulose (500 mg/kg of faropenem medoxomil t.i.d., 200 mg/kg of amoxicillin t.i.d., 50 mg/kg of clavulanate t.i.d., and moxifloxacin q.d.). All administrations (except for moxifloxacin group) were preceded by an oral dose of probenecid (P) (200 mg/kg, 1% methylcellulose). Only the moxifloxacin group was administered with P. Mice were treated with oral P alone, oral P and each drug component in clinical trials as potential combination part of novel TB regimens.

Drug combinations covering extensive periods of time (T > MIC) to drive efficacy. The benefits of the DHP-I-deficient model were further confirmed by the lack of observable β-lactam antibacterial effect in the reference C57BL/6 strain under the same experimental design (see Fig. S5 in the supplemental material).

In summary, based on the preliminary evidence presented here, we have been able to show how a DHP-I-deficient TF3157 murine model capable of sustaining significant β-lactam compound blood concentrations can be applicable to β-lactam drug discovery against TB. This model gives robust and significant efficacy signals measured as CFU reductions after treatment and can be used to discern between different β-lactam drugs. We believe that these data can modestly contribute to countering some of the past objections to the antibacterial potential of β-lactams. These results pave the way for the systematic in vivo characterization of both old and novel β-lactams as antibacterial leads and for the selection and eventual inclusion of safe and efficacious β-lactam drug components in clinical trials as potential combination partners in novel TB regimens.

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REFERENCES


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