Thioridazine: A Non-Antibiotic Drug Highly Effective, in Combination with First Line Anti-Tuberculosis Drugs, against Any Form of Antibiotic Resistance of Mycobacterium tuberculosis Due to Its Multi-Mechanisms of Action

Leonard Amaral 1,2,* and Miguel Viveiros 3

1 Institute of Hygiene and Tropical Medicine, Universidade Nova de Lisboa, Lisbon 1349-008, Portugal
2 Institute of Medical Microbiology and Immunobiology, University of Szeged, Szeged 6720, Hungary
3 Unidade de Microbiologia Médica, Global Health and Tropical Medicine, Instituto de Higiene e Medicina Tropical, Universidade Nova de Lisboa, Lisbon 1349-008, Portugal; MViveiros@ihmt.unl.pt
* Correspondence: lamaral@ihmt.unl.pt; Tel.: +351-21-3652600

Academic Editor: William R. Schwan
Received: 8 October 2016; Accepted: 5 January 2017; Published: 14 January 2017

Abstract: This review presents the evidence that supports the use of thioridazine (TZ) for the therapy of a pulmonary tuberculosis infection regardless of its antibiotic resistance status. The evidence consists of in vitro and ex vivo assays that demonstrate the activity of TZ against all encountered Mycobacterium tuberculosis (Mtb) regardless of its antibiotic resistance phenotype, as well as in vivo as a therapy for mice infected with multi-drug resistant strains of Mtb, or for human subjects infected with extensively drug resistant (XDR) Mtb. The mechanisms of action by which TZ brings about successful therapeutic outcomes are presented in detail.

Keywords: antibiotic resistant pulmonary tuberculosis; thioridazine; in vitro activity; ex vivo activity; in vivo activity (murine model and human trails); the non-killing macrophage; enhancement of intracellular killing of Mtb by thioridazine; mechanism of actions of thioridazine

1. Introduction

The resurgence of tuberculosis during the 1980s was followed in the 1990s in New York City by a dramatic increase in the rate of pulmonary tuberculosis (TB) infections accompanied with emerging levels of resistance to the first line anti-tuberculosis (anti-TB) drugs isoniazid and rifampicin—termed multi-drug resistant tuberculosis (MDR-TB) [1]. Resistance to these drugs plus resistance to the second line drugs, amikacin, kanamycin, capreomycin and fluoroquinolones—termed extensively drug resistant tuberculosis (XDR-TB)—was soon noted in various parts of the globe [2]. More recently, resistance to all anti-tuberculosis drugs—termed totally drug resistant tuberculosis (TDR-TB) was first noted in Italy in 2006 and later reported in Iran, India and South Africa [3]. Although the global acquisition of tuberculosis infections has decreased by 1.5% per year [4,5], the progression of increased resistance of Mycobacterium tuberculosis (Mtb), as a consequence of prolonged and problematic therapeutic regimens, threatens the progress that has been made since 1990 in the control and prevention of TB [5]. Although the efficacy of the repurposed and newly recommended antibiotic for resistant forms of Mtb—Linezolid—has been recently evaluated from data provided by 23 studies in 14 different countries, involving more than 500 patients, suggests an overall success rate of 77% [6], the drug is notorious for producing a plethora of serious side effects such as neuropathy and hematological disorders [7]. Other newly approved drugs for MDR and XDR-TB therapeutics—bedaquiline and delamalid—are following the same path with their recognized efficacy against resistant forms of TB.
being threatened by their market cost and cumulative reports of side effects and lack of safety [8]. Despite the occurrence of these side effects, in the absence of better forms of effective therapeutic regimens, the World Health Organization (WHO) continues to recommend the use of these extremely costly drugs for therapy of MDR and XDR-TB [4,6,8]. But is there an inexpensive drug, in comparison to the cost of these drugs, that has been extensively studied and has been safely used for the therapy of psychosis for over 50 years producing no serious side effects if the patient are monitored properly? The answer resides in thioridazine!

It is the purpose of this review to present all of the evidence, much of it confirmed by many groups around the world, which strongly supports that use of thioridazine (TZ) in combination with antibiotics to which the Mtb isolate was previously resistant for therapy of MDR, XDR and most probably, TDR.

2. Mycobacterium tuberculosis and Phenothiazines: Chlorpromazine, Thioridazine, in Vitro Activities

Phenothiazines are heterocyclic compounds. The first such compound was synthesized by Bernthsen in 1883 via the reaction of diphenylamine with sulfur. Methylene blue (MB) is a phenothiazine that was synthesized from a phenothiazine derivative by Heinrich August Bernthsen in 1883. Soon thereafter, the chemist Paul Erhlich used the MB dye for staining live cells and found that it could reduce movement of microorganisms [9]. This observation spurred experiments with humans that showed that the dye could render the subject sedated and was effective in the treatment of schizophrenia [10]. These discoveries led to the synthesis of chlorpromazine (CPZ) by Paul Charpentier in 1950, introduced by Rhone Poulenc as the first true neuroleptic in 1957 [10] (Figure 1).

![Thioridazine and Chlorpromazine](image)

**Figure 1.** Structures of the phenothiazines thioridazine and chlorpromazine [9].

Because of its worldwide use, anecdotal observations suggested that it had antimycobacterial properties [11]. By 1977, the in vitro antimycobacterial properties of CPZ were clearly shown [12] and confirmed 10 years later [13]. As a consequence of the emergence of a pulmonary tuberculosis epidemic in New York City during the late 1980s, and later the large percentage of MDR-TB coupled with the absence of new and effective anti-TB drugs at that time, the search for new anti-TB drugs began. The observations that CPZ had potential anti-TB activity spurred the study demonstrating that the in vitro concentrations of CPZ needed to inhibit the replication of Mtb could be exceeded and safely achieved in the CPZ treated patient, and clinically relevant concentrations ex vivo could effectively promote the killing of phagocytosed Mtb [14]. Regrettably, CPZ is also a drug that causes very serious side effects [15]. Thioridazine (TZ) is an equally effective neuroleptic phenothiazine. It produces significantly fewer side effects when used with moderation and maintaining an evaluation of the patient for underlying cardiopathy. TZ is therapeutically safe, as proven by the 60 plus years it has been in use, and is still widely used today in many countries to control psychosis. Consequently,
TZ was examined for in vitro activity against antibiotic susceptible and antibiotic resistant isolates of Mtb and compared to the activity of CPZ against the same strains [16]. The MICs in vitro for CPZ and TZ were calculated as ranging between 4 and 32 µg/mL, depending on the system and the antibiotic resistance status of the tested strain, and they were equally effective [17]. For the M. tuberculosis H37Rv fully antibiotic susceptible reference strain this range was determined, by many authors, to be 8–15 µg/mL depending on the system (Table 1).

Table 1. Minimum inhibitory concentrations of thioridazine for M. tuberculosis H37Rv reference strain according to the literature.

| Culture Systems          | TZ             | References |
|--------------------------|----------------|------------|
| MICs                     |                |            |
| MB7H10                   | 8–10 µg/mL     | [18–20]    |
| Broth microdilution      | 8–15 µg/mL     | [21–23]    |
| Bactec TB-460            | 15 µg/mL       | [24,25]    |
| Bactec MGIT 960          | 15 µg/mL       | [26]       |
| Cytotoxicity *           | IC50 13.78 µM  | [23]       |

* Evaluated against human monocyte-derived macrophages after 72 h of incubation at 37°C, 5% CO₂.

Both CPZ and TZ had similar activity against strains susceptible to isoniazid (INH) and rifampicin (RIF) as well as to strains resistant to these antibiotics and as many as five other antibiotics. The in vitro effects of TZ [20,27,28] as well its derivatives, have since been repeatedly confirmed [23,29]. However, the minimal inhibitory concentrations that completely inhibited the replication of Mtb in vitro employed in all of the cited studies exceeded many fold that which can be safely achieved clinically (ca. 0.5 mg/L of plasma in a patient chronically treated with TZ). Because CPZ had been shown to reduce the resistance of a number of pathogenic bacterial species to antibiotics [30], presumably by interacting with the cell wall of the bacterium [31], the effect of TZ on the resistance of isolates of Mtb to antibiotics was also evaluated (Figure 2). Briefly, although all of the phenothiazines tested were able to reduce the resistance to first line anti-TB drugs, the very mild neuroleptic TZ demonstrated great effectiveness at concentrations that were clinically achievable and similar to those employed for the initial therapy of psychosis [32]. Although the mechanism by which TZ reduced antibiotic resistance of Mtb was not readily understood, studies in other groups of bacteria demonstrated that TZ reversed the resistance of Escherichia coli to tetracycline by inhibiting the over-expressed efflux pump of the bacterium that was responsible for its multi-drug resistant phenotype [33]. Consequently, a large number of clinical isolates of Mtb that were susceptible to isoniazid (INH) were induced to extremely high level resistance to INH; this resistance could be totally reversed with a small and clinically relevant concentration of TZ [34]. Further studies showed that TZ reversed resistance of Mtb that had been induced to high level resistance to INH via the interference with the over-expressed efflux pumps genes of the organism mmpL7, p55, efpA, mnr, Rv1258c and Rv2459 [35] thereby confirming the previous observations on the role of efflux pumps in the multidrug resistance of the organism [34,36].

The effect of TZ on Mtb is not limited to the inhibition of efflux pumps. Studies by Dutta et al. show that, besides efflux pumps, the genes that code for essential proteins of the cell envelope are affected by TZ, albeit at concentrations that exceed the minimum inhibitory concentration of the drug [37]. Among the genes affected were those that encode efflux pumps that extrude antibiotics, oxido-reductases, enzymes involved in fatty acid metabolism and aerobic respiration, and genes that are co-expressed with the global SigmaB regulon, which are involved in the response to stress [38]. Other studies have confirmed these observations and have extended the understanding that TZ affects a large number of essential genes that code for proteins of the plasma membrane, many of which are involved in controlling essential energy production, active transport and permeability processes in response to antibiotic and oxidative stress stimuli [39]. In particular, several studies confirmed that TZ acts in mycobacterial respiratory chain components involved in ATP oxidative phosphorylation,
namely, the type-II NADH-menadione oxidoreductase (NDH-2)—a key component of respiratory chain of Mtb—thus raising the hypothesis that this is the main molecular target of TZ and making it also effective against latent TB [40–42]. NDH-2 catalyzes the first reaction of the electron transfer chain of Mtb that leads to ATP oxidative phosphorylation. During this reaction, NDH-2 transfers two electrons from NADH to menadione, which is reduced to menadionol form. Yano et al. have shown that the respiratory functions leading to de novo ATP synthesis and NADH regeneration might be the Achilles’ heel of hypoxic nonreplicating mycobacteria, making TZ an attractive drug with activity both against replicative and dormant mycobacteria [40–42]. This hypothesis has been confirmed by Sohaskey et al. who demonstrated that concentrations of TZ exceeding the MIC for actively replicating Mtb also inhibit/kill dormant Mtb, becoming a promising drug to control latent tuberculosis and shorten anti-TB drug regimens if used directly on the human macrophage [43,44]. However, the question of whether TZ can be clinically useful for inhibiting the replication of Mtb and simultaneously killing dormant Mtb remains doubtful unless science demonstrates that these effective in vitro concentrations can be achieved at the site of the pulmonary system where the infective organism normally resides, namely, the pulmonary macrophage. Because TZ is concentrated by cells such as macrophages that are rich in their lysosome content [45,46] to levels that theoretically are assumed to greatly exceed the concentration present in the medium (in fact never measured inside the macrophage, only measured in TZ-loaded culture lysates) [14,47], the noted effects of TZ on essential genes may take place in vivo.

![Figure 2](image-url)

**Figure 2.** Killing effect of thioridazine over three days of infection. Effect of thioridazine on the intracellular survival of *M. tuberculosis* (Mtb) H37Rv and an XDR Mtb strain within human monocyte-derived macrophages at day three post infection. Isoniazid (INH) was tested at 0.1 µg/mL, and thioridazine (TZ) at 2.5 µg/mL. Data are presented as a mean of the percentage of the survival (adapted from [26]).

### 3. Thioridazine and Its Effect on Intracellular *Mycobacterium tuberculosis*

To test the hypothesis above, and based upon the evidence that TZ was equal to CPZ with respect to its antimycobacterial properties in vitro (and the fact that CPZ was also very effective ex vivo), the rationale and the experiments performed by Crowle et al. [14] were repeated by Ordway et al. with CPZ and TZ against clinical strains of MDR Mtb [24], and later by Machado et al. against XDR Mtb, where TZ showed an excellent synergistic effect with first line drugs [26] (See Figure 1 as an example). In these works TZ was shown to enhance the killing of intracellular antibiotic susceptible and MDR/XDR Mtb by monocyte-derived human macrophages that have little killing action of their own at concentrations in the medium which are equivalent or lower than those present in the plasma of a thioridazine-treated psychotic patient (0.5 mg/L of plasma). These TZ ex vivo studies were extended to a large number of TZ derivatives, some of which revealed to be more effective than TZ and all of which expressed no toxicity at their effective ex vivo concentrations [29]. The same rationale and technical approach was further expanded from TZ and CPZ to other ion channel blockers.
such as verapamil, flupenthixol and haloperidol, with very successful results in enhancing the killing activity of the infected macrophage, regardless of the drug resistance profile of the infectious Mtb, with moderate and acceptable toxicities and excellent synergistic effects with first and second line anti-TB drugs [26].

The mechanism by which TZ promotes the killing of intracellular Mtb was at first opined to be the result of TZ being concentrated within the phagolysosome as predicted by Daniel and Wojcikowski [45,46] to a level compatible to its minimum bactericidal concentration of 60 mg/L. Although the concentrated effect in phagocytosed mycobacteria may take place, up to now this has never truly been demonstrated to occur in vivo. Controversial and disputed studies of Segal and associates [48,49] have also suggested another hypothesis which, if correct, could alter the form of therapy used against the MDR and XDR Mtb [24,50].

This hypothesis involves a series of stages that begin with: binding of the infecting Mtb organism to the receptor of the plasma membrane of the macrophage [51–53], followed with the invagination of the receptor-mycobacterium forming the phagosome which travels through the cytoplasm of the macrophage [54,55] and eventually fuses with a lysosome [56] to form the phagolysosome unit. The activation of dormant hydrolases (zymogen granules) requires a low pH [56] which is created by vesicular ATPases of the phagolysosome unit which are dependent upon the retention of ions [57]. Because the macrophage plasma membrane have their Ca^{2+} channels (L-type) inhibited in presence of TZ, this inhibition will result in a significant increase of Ca^{2+} from intracellular stores within macrophages. Accumulation of these ions in the cytoplasm of the macrophage causes an indirect acidification of the phagolysosome (schematic overview in Figure 3). Consequently, we considered the possibility that since TZ inhibits efflux pumps of bacteria [34–36] and also acts against efflux pumps of human cells [58–60], and phenothiazines in general inhibit Ca^{2+}/ion channel transport [25,61], TZ may also inhibit the efflux of ions from the phagolysosomal unit leading to the indirect acidification of the compartment and the activation of hydrolytic enzymes. This possibility is supported by recent studies of Machado et al. [26,35] demonstrating that TZ promotes the acidification of the phagolysosomal unit by indirect inhibition of macrophage ion channels. The inhibition of these channels therefore activates the hydrolytic enzymes via the coupling of the vesicular ATPases and consequent killing of the entrapped Mtb organism. This hypothesis is further supported by separate research using another ion channel blocker, verapamil [62,63]. These studies not only provide support for the use of TZ for therapy of Mtb drug resistant infections by a non-antibiotic compound [64–67], but also introduce an alternative therapeutic strategy that targets the killing machinery of the pulmonary macrophage infected with Mtb [62,63,67–70]. Drugs that target mycobacteria will eventually cause the organism to become resistant via the development of mutations at the gene coding level of the antibiotic target, and the alternative form of therapy with TZ evades this mutagenic response and assists the still effective antibiotics against drug resistant forms of Mtb.

In Figure 3 a model of the putative mechanism of action of thioridazine inside the macrophage is depicted, combining all the contributions made so far to elucidate its remarkable enhancing activity of the macrophage killing activity [26,48–50].
4. Mono and Combinational Therapy with TZ: The Mouse and the Human

The question of whether the in vitro and ex vivo effects of TZ are reflected successfully in the murine model needed answering, and to this end, mono-TZ therapy of the Mtb infected mouse [72] as well as combination therapy with first line antibiotics in this model have both proven to be effective [73–75]. Nevertheless, because the results in the mouse model not always are reproduced in humans, the effectiveness of TZ-combination therapy needed to be investigated. To this end, TZ in combination with antibiotics to which the infective organism was initially resistant produced complete cures in 17 out of 18 XDR-TB patients in Argentina [76]. Mono-therapy of five terminal XDR-TB patients with TZ significantly improved their quality of life (elimination of night sweats, improved appetite, weight gain, reduction of disease-associated stress) and did contribute to a longer life span [71], but because TZ does not restore lost pulmonary tissue, the patients succumbed to the disease. Studies by Abbate et al. [70] and Udzwadia et al. [77] showed that the use of TZ was safe with no significant effects on QT intervals or any other cardiac property as per the rigorous monitoring carried out in these trials.
5. Important Considerations for Therapy of MDR/XDR Mtb Patients with TZ in Combination with Antibiotics to Which the Infecting Organism Is Resistant

The initial response of bacteria to an antibiotic or noxious agent is to over-express its efflux pumps [26,33–35,78–88]. When the concentration of the agent is progressively increased, the genes that control and code for the efflux pumps of the responding organisms are progressively increased [33–35,85,87–90]. However, when the initial concentration of the antibiotic is maintained below its MIC during repeated passages, the bacterium responds with progressive increases in its efflux activity of the pre-existing pumps [26,35,90]. Eventually, the appearance of resistance to a large variety of unrelated antibiotics begins to occur with a progressive concomitant increase of transport activity ultimately leading to the MDR phenotype with basal (normal) levels of efflux activity [35,90]. These observations tend to explain why antibiotic resistance of an infecting bacterium continues to increase although the dosing of the patient remains unchanged [27,79,84]. Moreover, they also suggest that in order to define the antibiotic status of a clinical isolate from a patient who may be a suitable candidate for adjunct therapy with TZ, the antibiotic profile as well as the activity of the efflux pump system of the infecting organism should be determined. Whereas the determination of an antibiotic resistance panel is routine for a laboratory that performs diagnostic studies for a suspected pulmonary tuberculosis infection, there are at this time few laboratories that perform the needed assays that define the efflux pump status of the infecting Mtb isolate. Fortunately, there are methodologies that have been developed which are not difficult to perform by a routine tuberculosis laboratory that do not require expensive instrumentation. When this cost is compared to the huge cost associated with therapy of an MDR Mtb infection due to mutations or to an over-expressed efflux pump system, where therapy is expected to be highly problematic [91], the cost is indeed minor. Based upon the above, any patient who is considered to be a candidate for adjunct TZ therapy with antibiotics must first have a clinical isolate evaluated for susceptibility to first line antibiotics. The status of the efflux pump system and the ability of TZ to reverse its in vitro resistance to specific antibiotics of the panel must also be investigated before treatment [91]. In addition to these assays, it would be of great interest to determine the effect of TZ on the survival of the infecting isolate by the patient’s own macrophage. Given positive answers from the above assays (antibiotic susceptibility panel; defined efflux pump system; ability of TZ to reverse resistance to the antibiotic(s) for which the isolate was initially resistant; and, effective enhanced killing activity of the macrophage-trapped isolate), the patient may well be a good candidate for therapy with TZ as an adjunct to antibiotics whose initial resistance was due to an over-expressed efflux pump system [91]. During the time the clinical isolate is accordingly being investigated as suggested, the patient must be evaluated for any cardiopathy. It must be noted that the use of TZ is safe and the suggested protocol for dosing the patient is one that begins with a low level of 25 mg/day that is increased weekly to 50, 100 and 200 mg/day. This protocol has been shown not to reduce the QT interval (increased time between contractions of left and right ventricles) [92–94], a side effect repeatedly noted in MDR-TB patients treated with fluoroquinolones, bedaquiline or delamanid and a limitation factor for the use of new regimens including synthetic drugs [95]. However, approximately 6% of the Eastern European population has a mutation in the p450 cytochrome which reduces the metabolism of TZ, and consequently, the build-up of plasma TZ levels will result [93]. This build-up may be rapid and reach levels which are known to reduce the QT interval [94]. Consequently, the patient should be monitored for cardiac function prior to therapy with TZ in order to rule out any cardiopathy that may worsen with TZ dose, and monitoring should continue for the first week of therapy with TZ and periodically thereafter. It is important to note that TZ is safe to use for up to 1000 mg/day when introduced to the patient gradually [76,77,93,94], coupled to knowing the patient’s clinical history and performing cardiac monitoring as recommended. At this time, the time required for therapy leading to a negative TB culture and radiological evaluation consistent with cure is not known although as per Abbate et al. complete cures were achieved with XDR-TB patients within a few months of TZ adjunct therapy [76,77]. It may well be that full recovery of XDR TB patients takes place within a period of time commensurate with that routinely producing
complete cures of the patient infected with antibiotic susceptible Mtb with daily doses that are far below those used for the therapy of a psychotic patient.

6. Costs Associated with the Care of an MDR-TB Patient

The average cost for hospitalization during the period from 2005 through 2007 for an MDR-TB patient in the USA was $81,000 per year and for the XDR-TB patient $285,000 (3.5 times than that for the MDR-TB) [96]. Regardless of this huge expenditure per patient, the mortality rate for MDR-TB in the USA is still significantly higher than for antibiotic susceptible TB infections, and for XDR-TB significantly higher than for MDR infections, especially if the patient is co-infected with HIV or presents with AIDS [97]. Nevertheless, due to the development of a variety of clinical diagnostic programs, therapeutic monitoring such DOTS, and the wide introduction of rapid laboratory methods for the identification, isolation and susceptibility test to first line TB drugs, the frequency of TB infections susceptible and resistant to first line TB drugs has fallen dramatically [4,97]. In countries that are poor, the situation is totally reversed and the incidence of all forms of TB infections continues to rise rapidly [5,98]. Although it is not possible at this time to advance the status of TB control, therapy, etc. in these global regions, given the severity of increasing antibiotic resistance, it is reasonable that therapy of a clinical presentation of tuberculosis can be pursued without the luxury of what is present in wealthier countries. However, therapy with first line drugs is costly and if not properly administered leads to MDR, and progressively more resistant forms of TB. WHO has recommended linezolid to be included in the empirical protocols for the therapy of MDR/XDR-TB infections [4,5]. However, this drug has a very narrow therapeutic window and because the optimal dosing strategy that minimizes the substantial toxicity associated with prolonged use has not been determined [99,100], blind use of this drug is extremely expensive and problematic. Consequently, if far less expensive drugs such as those that make up the line of defense are available, and because TZ is safe when used as prescribed, and because the effective daily dose of TZ used by Abbate et al. was, via increments, limited to 200 mg/day [76], TZ in combination with first-line drugs may prove to be significantly effective for therapy of any form of tuberculosis. Given that TZ when concentrated by the phagolysosome will be effective against the efflux pumps that are responsible for MDR phenotype of the bacterium, and given that TZ may also reach a level in the phagolysosome compatible with that which is bactericidal in vitro, and coupled to the enhancement of killing by the macrophage housing the infective organism, the potential that TZ has for therapy of any TB pulmonary infection is significant and should continue to be further supported.

7. Conclusions

The body of results and evidences gathered so far, coming from many different contributions from different teams around the world, enable us to propose the following mechanism of action for thioridazine and other ion channel blockers in the bacteria: after entering the cell, the compounds will generate a cascade of events which starts with the inhibition of the respiratory chain complexes, though we cannot say at the present moment if the respiratory chain is a direct target. The inhibition of the bacterial respiratory chain will lead to dissipation of the membrane potential, reduction of ATP levels, efflux inhibition, oxidative stress, and increase in intracellular ion levels. On the host cell, treatment with these compounds results in phagosome acidification that synergizes with several components of the host immune response, such as lysosomal hydrolases, leading to bacterial growth restriction. Both effects cooperate and result in an enhanced killing activity that can be highly efficient when combined with antituberculosis drugs.

Promising examples of the future use of thioridazine in new short term therapeutic regimens against any form of antibiotic resistance of Mtb come from the recent studies that demonstrated the possibility of effectively using nanoparticles containing thioridazine and rifampicin for rapid tuberculosis treatment in vitro and in a zebrafish model [101,102]. The use of TZ as a therapeutic adjuvant for anti-TB therapy is currently being expanded in Argentina and India. Even the World
Health Organization, who has not shown great interest in the repurposing of this narcoleptic drug for TB has recently considered thioridazine as a World Health Organization group 5 drug for multidrug-resistant tuberculosis treatment due to its efficacy and safety [103].

Acknowledgments: The authors are thankful to our colleagues who contributed to this work for over 20 years, namely, Joseph Molnar, Jetta Kristiansen, Diane Ordway, Marta Martins, Diana Machado, Isabel Couto, Eduardo Abatte, Zarin Udwadia, Martin Boeree, Notten Dutta and Dick van Soolingen. Miguel Viveiros is thankful to FCT Fundação para a Ciência e a Tecnologia (FCT) for funding the Global Health and Tropical Medicine (GHTM) Research Center (Grant UID/Multi/04413/2013) and for the many grants provided to the authors from the FCT and Gulbenkian Foundation during the 20 year period of this study. A special acknowledgement is due to Diana Machado for her kind assistance with the figure design and revision of the paper.

Conflicts of Interest: The authors declare no conflict of interest.

References
1. Munsiff, S.S.; Nivin, B.; Sacaju, G.; Mathema, B.; Bifani, P.; Kreiswirth, B.N. Persistence of a highly resistant strain of tuberculosis in New York City during 1990–1999. J. Infect. Dis. 2003, 188, 356–363. [CrossRef] [PubMed]
2. Matteelli, A.; Roggi, A.; Carvalho, A.C. Extensively drug-resistant tuberculosis: Epidemiology and management. Clin. Epidemiol. 2014, 6, 111–118. [CrossRef] [PubMed]
3. Parida, S.K.; Axelsson-Robertson, R.; Rao, M.V.; Singh, N.; Master, I.; Lutckii, A.; Keshavjee, S.; Andersson, J.; Zumla, A.; Mæurer, M. Totally drug-resistant tuberculosis and adjunct therapies. J. Intern. Med. 2015, 277, 388–405. [CrossRef] [PubMed]
4. World Health Organization. Global Tuberculosis Report 2015; World Health Organization: Geneva, Switzerland, 2015. Available online: http://www.who.int/tb/publications/global_report/en/ (accessed on 3 October 2016).
5. World Health Organization. Key Bottlenecks in M/XDR-TB Control and Patient Care; World Health Organization: Geneva, Switzerland, 2009; Available online: http://www.who.int/tb/challenges/mdr/bottlenecks/en/ (accessed on 3 October 2016).
6. Agyeman, A.A.; Ofori-Asenso, R. Efficacy and safety profile of linezolid in the treatment of multidrug-resistant (MDR) and extensively drug-resistant (XDR) tuberculosis: A systematic review and meta-analysis. Ann. Clin. Microbiol. Antimicrob. 2016, 15. [CrossRef] [PubMed]
7. Zhang, X.; Falagas, M.E.; Vardakas, K.Z.; Wang, R.; Qin, R.; Wang, J.; Liu, Y. Systematic review and meta-analysis of the efficacy and safety of therapy with linezolid containing regimens in the treatment of multidrug-resistant and extensively drug-resistant tuberculosis. J. Thorac. Dis. 2015, 7, 603–615. [PubMed]
8. Olaru, I.D.; von Groote-Bidlingmaier, F.; Heyckendorf, J.; Yew, W.W.; Lange, C.; Chang, K.C. Novel drugs against tuberculosis: A clinician’s perspective. Eur. Respir. J. 2015, 45, 1119–1131. [CrossRef] [PubMed]
9. Ohlow, M.J.; Moosmann, B. Phenothiazine: The seven lives of pharmacology’s first lead structure. Drug Discov. Today 2011, 16, 119–131. [CrossRef] [PubMed]
10. Healy, D. Explorations in a New World. In The Creation of Psychopharmacology; Harvard University Press: Cambridge, CA, USA, 2004.
11. Gonzalez-Gonzalez, E. Use of largactil in pulmonary tuberculosis. Rev. Esp. Tuberc. 1958, 27, 134–136. [PubMed]
12. Mólnár, J.; Béládi, I.; Földes, I. Studies on antitubercular action of some phenothiazine derivatives in vitro. Zentralbl. Bakteriol. Orig. A 1977, 239, 521–526. [PubMed]
13. Kristiansen, J.E.; Vergmann, B. The antibacterial effect of selected phenothiazines and thioxanthenes on slow-growing mycobacteria. Acta Pathol. Microbiol. Immunol. Scand. B 1986, 94, 393–398. [CrossRef] [PubMed]
14. Crowle, A.J.; Douvas, G.S.; May, M.H. Chlorpromazine: A drug potentially useful for treating mycobacterial infections. Chemotherapy 1992, 38, 410–419. [CrossRef] [PubMed]
15. Gardos, G.; Cole, J.O. Maintenance antipsychotic therapy: Is the cure worse than the disease? Am. J. Psychiatry 1976, 133, 32–36. [PubMed]
16. Amaral, L.; Kristiansen, J.E.; Abebe, L.S.; Millett, W. Inhibition of the respiration of multi-drug resistant clinical isolates of Mycobacterium tuberculosis by thioridazine: Potential use for initial therapy of freshly diagnosed tuberculosis. J. Antimicrob. Chemother. 1996, 38, 1049–1053. [CrossRef] [PubMed]

17. Amaral, L.; Kristiansen, J.E.; Viveiros, M.; Atouguia, J. Activity of phenothiazines against antibiotic-resistant Mycobacterium tuberculosis: A review supporting further studies that may elucidate the potential use of thioridazine as anti-tuberculosis therapy. J. Antimicrob. Chemother. 2001, 47, 505–511. [CrossRef] [PubMed]

18. Ångeby, K.; Juréen, P.; Chryssanthou, E.; Schön, T. Tentative susceptibility testing breakpoint for the neuroleptic drug thioridazine, a treatment option for multi-and extensively drug resistant tuberculosis. Int. J. Mycobacteriol. 2012, 1, 177–179. [CrossRef] [PubMed]

19. Musuka, S.; Srivastava, S.; Dona, C.W.S.; Meek, C.; Leff, R.; Pasipanodya, J.; Gumbo, T. Thioridazine pharmacokinetic-pharmacodynamic parameters “Wobble” during treatment of tuberculosis: A theoretical basis for shorter-duration curative monotherapy with congeners. Antimicrob. Agents Chemother. 2013, 57, 5870–5877. [CrossRef] [PubMed]

20. Vesenbeckh, S.; Krieger, D.; Bettermann, G.; Schönfeld, N.; Bauer, T.T.; Rüssmann, H.; Mauch, H. Neuroleptic drugs in the treatment of tuberculosis: Minimal inhibitory concentrations of different phenothiazines against Mycobacterium tuberculosis. Tuberculosis 2016, 98, 27–29. [CrossRef] [PubMed]

21. Li, G.; Zhang, J.; Li, C.; Guo, Q.; Jiang, Y.; Wei, J.; Qiu, Y.; Zhao, X.; Zhao, L.L.; Lu, J.; et al. Antimycobacterial activity of five efflux pump inhibitors against Mycobacterium tuberculosis clinical isolates. J. Antibi. 2016, 69, 173–175. [CrossRef] [PubMed]

22. Coelho, T.; Machado, D.; Couto, I.; Maschmann, R.; Ramos, D.; von Groll, A.; Rossetti, M.L.; Silva, P.A.; Viveiros, M. Enhancement of antibiotic activity by efflux inhibitors against multidrug resistant Mycobacterium tuberculosis clinical isolates from Brazil. Front. Microbiol. 2015, 6. [CrossRef] [PubMed]

23. Pieron, M.; Machado, D.; Azzali, E.; Santos Costa, S.; Couto, I.; Costantino, G.; Viveiros, M. Rational design and synthesis of thioridazine analogues as enhancers of the antituberculosis therapy. J. Med. Chem. 2015, 58, 5842–5853. [CrossRef] [PubMed]

24. Ordway, D.; Viveiros, M.; Leandro, C.; Bettencourt, R.; Almeida, J.; Martins, M.; Kristiansen, J.E.; Molnar, J.; Amaral, L. Clinical concentrations of thioridazine kill intracellular multidrug-resistant Mycobacterium tuberculosis. Antimicrob. Agents Chemother. 2003, 47, 917–922. [CrossRef] [PubMed]

25. Martins, M.; Viveiros, M.; Amaral, L. Inhibitors of Ca2+ and K+ transport enhance intracellular killing of M. tuberculosis by non-killing macrophages. In Vitro 2008, 22, 69–75. [PubMed]

26. Machado, D.; Pires, D.; Perdigão, J.; Couto, I.; Portugal, I.; Martins, M.; Amaral, L.; Anes, E.; Viveiros, M. Ion channel blockers as antimicrobial agents, efflux inhibitors, and enhancers of macrophage killing activity against drug resistant Mycobacterium tuberculosis. PLoS ONE 2016, 11, e0149326. [CrossRef] [PubMed]

27. De Knecht, G.J.; ten Kate, M.T.; van Soolingen, D.; Aarnoutse, R.; Boeree, M.J.; Bakker-Woudenberg, I.A.; de Steenwinkel, J.E. Enhancement of in vitro activity of tuberculosis drugs by addition of thioridazine is not reflected by improved in vivo efficacy. Tuberculosis 2014, 94, 701–707. [CrossRef] [PubMed]

28. Van Ingen, J. The broad-spectrum antimycobacterial activities of phenothiazines, in vitro: Somewhere in all of this there may be patentable potentials. Recent Pat. Antinfect. Drug Discov. 2011, 6, 104–109. [CrossRef] [PubMed]

29. Martins, M.; Schelz, Z.; Martins, A.; Molnar, J.; Hajos, G.; Riedl, Z.; Viveiros, M.; Yalcin, I.; Aki-Sener, E.; Amaral, L. In vitro and ex vivo activity of thioridazine derivatives against Mycobacterium tuberculosis. Int. J. Antimicrob. Agents 2007, 29, 338–340. [CrossRef] [PubMed]

30. Amaral, L.; Kristiansen, J.; Lorian, V. Synergic effect of chlorpromazine on the activity of some antibiotics. J. Antimicrob. Chemother. 1992, 30, 556–558. [CrossRef] [PubMed]

31. Amaral, L.; Lorian, V. Effects of chlorpromazine on the cell envelope proteins of Escherichia coli. Antimicrob. Agents Chemother. 1991, 35, 1923–1924. [CrossRef] [PubMed]

32. Viveiros, M.; Amaral, L. Enhancement of antibiotic activity against poly-drug resistant Mycobacterium tuberculosis by phenothiazines. Int. J. Antimicrob. Agents 2001, 17, 225–228. [CrossRef] [PubMed]

33. Viveiros, M.; Jesus, A.; Brito, M.; Leandro, C.; Martins, M.; Ordway, D.; Molnar, A.M.; Molnar, J.; Amaral, L. Inducement and reversal of tetracycline resistance in Escherichia coli K-12 and expression of proton gradient-dependent multidrug efflux pump genes. Antimicrob. Agents Chemother. 2005, 49, 3578–3582. [CrossRef] [PubMed]
34. Viveiros, M.; Portugal, I.; Bettencourt, R.; Victor, T.C.; Jordaan, A.M.; Leandro, C.; Ordway, D.; Amaral, L. Isoniazid-induced transient high-level resistance in Mycobacterium tuberculosis. Antimicrob. Agents Chemother. 2002, 46, 2804–2810. [CrossRef] [PubMed]

35. Machado, D.; Couto, I.; Perdigão, J.; Rodrigues, L.; Portugal, I.; Baptista, P.; Veigas, B.; Amaral, L.; Viveiros, M. Contribution of efflux to the emergence of isoniazid and multidrug resistance in Mycobacterium tuberculosis. PLoS ONE 2012, 7, e34538. [CrossRef] [PubMed]

36. Rodrigues, L.; Machado, D.; Couto, I.; Amaral, L.; Viveiros, M. Contribution of efflux activity to isoniazid resistance in the Mycobacterium tuberculosis complex. Infect. Genet. Evol. 2012, 12, 695–700. [CrossRef] [PubMed]

37. Dutta, N.K.; Mazumdar, K.; Dastidar, S.G.; Karakousis, P.C.; Amaral, L. New patentable use of an old neuroleptic compound thioreidazine to combat tuberculosis: A gene regulation perspective. Recent Pat. Antinfec. Drug Discov. 2011, 6, 128–138. [CrossRef] [PubMed]

38. Dutta, N.K.; Mehra, S.; Kaushal, D. A Mycobacterium tuberculosis sigma factor network responds to cell-envelope damage by the promising anti-mycobacterial thioreidazine. PLoS ONE 2010, 5, e10069. [CrossRef] [PubMed]

39. De Keijzer, J.; Mulder, A.; de Haas, P.E.; de Ru, A.H.; Heerkens, E.M.; Amaral, L.; van Soolingen, D.; van Veelen, P.A. Thioreidazine alters the cell-envelope permeability of Mycobacterium tuberculosis. J. Proteom. Res. 2016, 15, 1776–1786. [CrossRef] [PubMed]

40. Yano, T.; Li, L.S.; Weinstein, E.; Teh, J.S.; Rubin, H. Steady-state kinetics and inhibitory action of antitubercular phenothiazines on Mycobacterium tuberculosis type-II NADH-menaquinone oxidoreductase (NDH-2). J. Biol. Chem. 2006, 281, 11456–11463. [CrossRef] [PubMed]

41. Teh, J.S.; Yano, T.; Rubin, H. Type II NADH: Menaquinone oxidoreductase of Mycobacterium tuberculosis. Infect. Disord. Drug Targets. 2007, 7, 169–181. [CrossRef] [PubMed]

42. Sharma, S.; Singh, A. Phenothiazines as anti-tubercular agents: Mechanistic insights and clinical implications. Expert Opin. Invest. Drugs 2011, 20, 1665–1676. [CrossRef] [PubMed]

43. Sohaskey, C.D. Nitrate enhances the survival of Mycobacterium tuberculosis during inhibition of respiration. J. Bacteriol. 2008, 190, 2981–2986. [CrossRef] [PubMed]

44. Sohaskey, C. Latent tuberculosis: Is there a role for thioreidazine? Recent Pat. Antiinfec. Drug Discov. 2011, 6, 139–146. [CrossRef] [PubMed]

45. Daniel, W.A.; Wojcikowski, J. Contribution of lysosomal trapping to the total tissue uptake of psychotropic drugs. Pharmacol. Toxicol. 1997, 80, 62–68. [CrossRef] [PubMed]

46. Daniel, W.A.; Wojcikowski, J. The role of lysosomes in the cellular distribution of thioreidazine and potential drug interactions. Toxicol. Appl. Pharmacol. 1999, 158, 115–124. [CrossRef] [PubMed]

47. Leonard, A.; Viveiros, M.; Kristiansen, J.E. “Non-Antibiotics”: Alternative therapy for the management of MDRTB and MRSA in economically disadvantaged Countries. Curr. Drug Targets 2006, 7, 887–891.

48. Reeves, E.P.; Lu, H.; Jacobs, H.L.; Messina, C.G.; Bolsover, S.; Gabella, G.; Potma, E.O.; Warley, A.; Roes, J.; Segal, A.W. Killing activity of neutrophils is mediated through activation of proteases by K+ flux. Nature 2002, 416, 291–297. [CrossRef] [PubMed]

49. Ahluwalia, J.; Tinker, A.; Clapp, L.H.; Duchen, M.R.; Abramov, A.Y.; Pope, S.; Nobles, M.; Segal, A.W. The large-conductance Ca2+-activated K+ channel is essential for innate immunity. Nature 2004, 427, 853–858. [CrossRef] [PubMed]

50. Viveiros, M.; Martins, M.; Rodrigues, L.; Machado, D.; Couto, I.; Ainsa, J.; Amaral, L. Inhibitors of mycobacterial efflux pumps as potential boosters for anti-tubercular drugs. Expert Rev. Anti. Infect. Ther. 2012, 10, 983–998. [CrossRef] [PubMed]

51. Torrelles, J.B.; Schlesinger, L.S. Diversity in Mycobacterium tuberculosis mannosylated cell wall determinants impacts adaptation to the host. Tuberculosis 2010, 90, 84–93. [CrossRef] [PubMed]

52. Józefowski, S.; Sobota, A.; Kwiatkowska, K. How Mycobacterium tuberculosis subverts host immune responses. Bioessays 2008, 30, 943–954. [CrossRef] [PubMed]

53. Yoshikai, Y. Immunological protection against mycobacterium tuberculosis infection. Crit. Rev. Immunol. 2006, 26, 515–526. [CrossRef] [PubMed]

54. Wall, A.A.; Condon, N.D.; Yeo, J.C.; Hamilton, N.A.; Stow, J.L. Dynamic imaging of the recycling endosomal network in macrophages. Methods Cell Biol. 2015, 130, 1–18. [PubMed]
55. Van der Wel, N.; Hava, D.; Houben, D.; Fluitsma, D.; van Zon, M.; Pierson, J.; Brenner, M.; Peters, P.J.M. Tuberculosis and M. leprae translocate from the phagolysosome to the cytosol in myeloid cells. *Cell* **2007**, *129*, 1287–1298. [CrossRef] [PubMed]

56. Ganley, I.G. Autophagosome maturation and lysosomal fusion. *Essays Biochem.* **2013**, *55*, 65–78. [CrossRef] [PubMed]

57. Harvey, W.R. Voltage coupling of primary H+ V-ATPases to secondary Na+- or K+-dependent transporters. *J. Exp. Biol.* **2009**, *212*, 1620–1629. [CrossRef] [PubMed]

58. Csonka, Á.; Spengler, G.; Martins, A.; Ocsovszki, I.; Christensen, J.B.; Hendricks, O.; Kristiansen, J.E.; Amaral, L.; Molnar, J. The mechanism by which the phenothiazine thioridazine stereoisomers on the drug accumulation of mouse lymphoma and human prostate cancer cell lines in vitro. *In Vivo* **2013**, *27*, 815–820. [PubMed]

59. Spengler, G.; Molnar, J.; Viveiros, M.; Amaral, L. Thioridazine induces apoptosis of multidrug-resistant mouse lymphoma cells transfected with the human ABCB1 and inhibits the expression of P-glycoprotein. *Anticancer Res.* **2011**, *31*, 4201–4205. [PubMed]

60. Spengler, G.; Takács, D.; Horváth, A.; Riedl, Z.; Hajós, G.; Amaral, L.; Molnár, J. Multidrug resistance reversing activity of newly developed phenothiazines on P-glycoprotein (ABCB1)-related resistance of mouse T-lymphoma cells. *Anticancer Res.* **2014**, *34*, 1737–1741. [PubMed]

61. Amaral, L.; Martins, M.; Viveiros, M. Enhanced killing of intracellular multidrug-resistant *Mycobacterium tuberculosis* by compounds that affect the activity of efflux pumps. *J. Antimicrob. Chemother.* **2007**, *59*, 1237–1246. [CrossRef] [PubMed]

62. Adams, K.N.; Takaki, K.; Connolly, L.E.; Wiedenhoft, H.; Winglee, K.; Humbert, O.; Edelstein, P.H.; Cosma, C.L.; Ramakrishnan, L. Drug tolerance in replicating mycobacteria mediated by a macrophage-induced efflux mechanism. *Cell* **2011**, *145*, 39–53. [CrossRef] [PubMed]

63. Gupta, S.; Tyagi, S.; Almeida, D.V.; Maiga, M.C.; Ammerman, N.C.; Bishai, W.R. Acceleration of tuberculosis treatment by adjunctive therapy with verapamil as an efflux inhibitor. *Am. J. Respir. Crit. Care Med.* **2013**, *188*, 600–607. [CrossRef] [PubMed]

64. Amaral, L.; Molnar, J. Mechanisms by which thioridazine in combination with antibiotics cures extensively drug-resistant infections of pulmonary tuberculosis. *In Vivo* **2014**, *28*, 267–271. [PubMed]

65. Amaral, L.; Martins, A.; Spengler, G.; Hunyadi, A.; Molnar, J. The mechanism by which the phenothiazine thioridazine contributes to cure problematic drug-resistant forms of pulmonary tuberculosis: Recent patents for “new use”. *Recent Pat. Antiinfect. Drug Discov.* **2013**, *8*, 206–212. [CrossRef] [PubMed]

66. Amaral, L.; Udwadia, Z.; Abbate, E.; van Soolingen, D. The added effect of thioridazine in the treatment of drug-resistant tuberculosis. *Int. J. Tuberc. Lung Dis.* **2012**, *16*, 1706–1708. [CrossRef] [PubMed]

67. Amaral, L.; Molnar, J. Why and how thioridazine in combination with antibiotics to which the infective strain is resistant will cure totally drug-resistant tuberculosis. *Expert Rev. Anti. Infect. Ther.* **2012**, *10*, 869–873. [CrossRef] [PubMed]

68. Amaral, L. Thioridazine: An old neuroleptic effective against totally drug resistant tuberculosis. *Acta Med. Port.* **2012**, *25*, 118–121. [PubMed]

69. Martins, M. Targeting the human macrophage with combinations of drugs and inhibitors of Ca²⁺ and K⁺ transport to enhance the killing of intracellular multi-drug resistant *Mycobacterium tuberculosis* (MDR-TB)—A novel, patentable approach to limit the emergence of XDR-TB. *Recent Pat. Antiinfect. Drug Discov.* **2011**, *6*, 110–117. [CrossRef] [PubMed]

70. Amaral, L.; Martins, A.; Molnar, J.; Kristiansen, J.E.; Martins, M.; Viveiros, M.; Rodrigues, L.; Spengler, G.; Couto, I.; Ramos, J.; et al. Phenothiazines, bacterial efflux pumps and targeting the macrophage for enhanced killing of intracellular XDR-TB. *In Vivo* **2010**, *24*, 409–424. [PubMed]

71. Gupta, S.; Salam, N.; Srivastava, V.; Singla, R.; Behera, D.; Khayyam, K.U.; Korde, R.; Malhotra, P.; Saxena, R.; Natarajan, K. Voltage gated calcium channels negatively regulate protective immunity to *Mycobacterium tuberculosis*. *PLoS ONE* **2009**, *4*, e5305. [CrossRef] [PubMed]

72. Martins, M.; Viveiros, M.; Kristiansen, J.E.; Molnar, J.; Amaral, L. The curative activity of thioridazine on mice infected with *Mycobacterium tuberculosis*. *In Vivo* **2007**, *21*, 771–775. [PubMed]

73. Dutta, N.K.; Pinn, M.L.; Karakousis, P.C. Sterilizing activity of thioridazine in combination with the first-line regimen against acute murine tuberculosis. *Antimicrob. Agents Chemother.* **2014**, *58*, 5567–5569. [CrossRef] [PubMed]
74. Dutta, N.K.; Pinn, M.L.; Karakousis, P.C. Reduced emergence of isoniazid resistance with concurrent use of thioridazine against acute murine tuberculosis. *Antimicrob. Agents Chemother.* 2014, 58, 4048–4053. [CrossRef] [PubMed]

75. Van Soolingen, D.; Hernandez-Pando, R.; Orozco, H.; Aguilar, D.; Magis-Escurra, C.; Amaral, L.; van Ingen, J.; Boeree, M.J. The antipsychotic thioridazine shows promising therapeutic activity in a mouse model of multidrug-resistant tuberculosis. *PLoS ONE* 2010, 5, e12640. [CrossRef] [PubMed]

76. Abbate, E.; Vescovo, M.; Natiello, M.; Cufré, M.; García, A.; Gonzalez Montaner, P.; Ambroggi, M.; Ritacco, V.; van Soolingen, D. Successful alternative treatment of extensively drug-resistant tuberculosis in Argentina with a combination of linezolid, moxifloxacin and thioridazine. *J. Antimicrob. Chemother.* 2012, 67, 473–477. [CrossRef] [PubMed]

77. Udwadia, Z.F.; Sen, T.; Pinto, L.M. Safety and efficacy of thioridazine as salvage therapy in Indian patients with XDR-TB. *Recent Pat. Infect. Drug Discov.* 2011, 6, 88–91. [CrossRef] [PubMed]

78. Amaral, L.; Martins, A.; Spengler, G.; Molnar, J. Efflux pumps of Gram-negative bacteria: What they do, how they do it, with what and how to deal with them. *Front. Pharmacol.* 2014, 4. [CrossRef] [PubMed]

79. Martins, A.; Hunyadi, A.; Amaral, L. Mechanisms of resistance in bacteria: An evolutionary approach. *Open Microbiol. J.* 2013, 7, 53–58. [CrossRef] [PubMed]

80. Spengler, G.; Rodrigues, L.; Martins, A.; Martins, M.; McCusker, M.; Cerca, P.; Machado, L.; Costa, S.S.; Ntoukou, E.; Couto, I.; et al. Genetic response of *Salmonella enterica* serotype Enteritidis to thioridazine rendering the organism resistant to the agent. *Int. J. Antimicrob. Agents.* 2012, 39, 16–21. [CrossRef] [PubMed]

81. Walter, N.D.; Dolganov, G.M.; Garcia, B.J.; Worodria, W.; Andama, A.; Musisi, E.; Ayakaka, I.; Van, T.T.; Voskuil, M.I.; de Jong, B.C.; et al. Transcriptional adaptation of drug-tolerant *Mycobacterium tuberculosis* during treatment of human tuberculosis. *J. Infect. Dis.* 2015, 212, 990–998. [CrossRef] [PubMed]

82. Amaral, L.; Cerca, P.; Spengler, G.; Machado, L.; Martins, A.; Couto, I.; Viveiros, M.; Fanning, S.; Pagès, J.M. Ethidium bromide efflux by *Salmonella*: Modulation by metabolic energy, pH, ions and phenothiazines. *Int. J. Antimicrob. Agents.* 2011, 38, 140–145. [CrossRef] [PubMed]

83. Martins, A.; Machado, L.; Costa, S.; Cerca, P.; Spengler, G.; Viveiros, M.; Amaral, L. Role of calcium in the efflux system of *Escherichia coli*. *Int. J. Antimicrob. Agents.* 2011, 37, 410–414. [CrossRef] [PubMed]

84. Black, P.A.; Warren, R.M.; Louw, G.E.; van Helden, P.D.; Victor, T.C.; Kana, B.D. Energy metabolism and drug efflux in *Mycobacterium tuberculosis*. *Antimicrob. Agents Chemother.* 2014, 58, 2491–2503. [CrossRef] [PubMed]

85. Schmaltzieg, A.M.; Srivastava, S.; Belkaya, S.; Deshpande, D.; Meek, C.; Leff, R.; van Oers, N.S.; Gummo, T. The antibiotic resistance arrow of time: Efflux pump induction is a general first step in the evolution of mycobacterial drug resistance. *Antimicrob. Agents Chemother.* 2012, 56, 4806–4815. [CrossRef] [PubMed]

86. Martins, A.; Spengler, G.; Martins, M.; Rodrigues, L.; Viveiros, M.; Dabiv-Regli, A.; Chevalier, J.; Couto, I.; Pagès, J.M.; Amaral, L. Physiological characterisation of the efflux pump system of antibiotic-susceptible and multidrug-resistant *Enterobacter aerogenes*. *Int. J. Antimicrob. Agents.* 2010, 36, 313–318. [CrossRef] [PubMed]

87. Martins, A.; Iversen, C.; Rodrigues, L.; Cerca, P.; Ramos, J.; Kern, W.V.; Couto, I.; Viveiros, M.; Fanning, S.; Pages, J.M.; et al. An AcrAB-mediated multidrug-resistant phenotype is maintained following restoration of wild-type activities by efflux pump genes and their regulators. *Int. J. Antimicrob. Agents.* 2009, 34, 602–604. [CrossRef] [PubMed]

88. Martins, A.; Spengler, G.; Rodrigues, L.; Viveiros, M.; Ramos, J.; Martins, M.; Couto, I.; Fanning, S.; Pagès, J.M.; Bolla, J.M.; et al. pH Modulation of efflux pump activity of multi-drug resistant *Escherichia coli*: Protection during its passage and eventual colonization of the colon. *PLoS ONE* 2009, 4, e6656. [CrossRef] [PubMed]

89. Martins, A.; Couto, I.; Aagaard, L.; Martins, M.; Viveiros, M.; Kristiansen, J.E.; Amaral, L. Prolonged exposure of methicillin-resistant *Staphylococcus aureus* (MRSA) COL strain to increasing concentrations of oxacillin results in a multidrug-resistant phenotype. *Int. J. Antimicrob. Agents.* 2007, 29, 302–305. [CrossRef] [PubMed]

90. Viveiros, M.; Dupont, M.; Rodrigues, L.; Couto, I.; Dabin-Regli, A.; Martins, M.; Pagès, J.M.; Amaral, L. Antibiotic stress, genetic response and altered permeability of *E. coli*. *PLoS ONE* 2007, 2, e126. [CrossRef] [PubMed]

91. Amaral, L.; van Soolingen, D. A novel advanced laboratory diagnosis to guide tuberculosis drug therapy. *Recent Pat. Antinfect. Drug Discov.* 2015, 10, 71–73. [CrossRef] [PubMed]

92. Beach, S.R.; Celano, C.M.; Noseworthy, P.A.; Januzzi, J.L.; Huffman, J.C. QTc prolongation, torsades de pointes, and psychotropic medications. *Psychosomatics* 2013, 54, 1–13. [CrossRef] [PubMed]
93. Thanacoody, R.H.; Daly, A.K.; Reilly, J.G.; Ferrier, I.N.; Thomas, S.H. Factors affecting drug concentrations and QT interval during thioridazine therapy. *Clin. Pharmacol. Ther.* 2007, 82, 555–565. [CrossRef] [PubMed]

94. Thanacoody, R.H. Thioridazine: The good and the bad. *Recent Pat. Antiinfect. Drug Discov.* 2011, 6, 92–98. [CrossRef] [PubMed]

95. Kwon, Y.S.; Koh, W.J. Synthetic investigational new drugs for the treatment of tuberculosis. *Expert Opin. Invest. Drugs* 2016, 25, 183–193. [CrossRef] [PubMed]

96. Marks, S.M.; Hirsch-Moverman, Y.; Salcedo, K.; Graviss, E.A.; Oh, P.; Seaworth, B.; Flood, J.; Armstrong, L.; Armitige, L.; Mase, S. TB epidemiologic studies consortium. Characteristics and costs of multidrug-resistant tuberculosis in-patient care in the United States, 2005–2007. *Int. J. Tuberc. Lung Dis.* 2016, 20, 435–441. [CrossRef] [PubMed]

97. Shah, N.S.; Pratt, R.; Armstrong, L.; Robison, V.; Castro, K.G.; Cegielski, J.P. Extensively drug-resistant tuberculosis in the United States, 1993–2007. *JAMA* 2008, 300, 2153–2160. [CrossRef] [PubMed]

98. Raviglione, M.; Sulis, G. Tuberculosis 2015: Burden, challenges and strategy for control and elimination. *Infect. Dis Rep.* 2016, 8. [CrossRef] [PubMed]

99. Sotgiu, G.; Centis, R.; D’ambrosio, L.; Migliori, G.B. Tuberculosis treatment and drug regimens. *Cold Spring Harb. Perspect. Med.* 2015, 5, a017822. [CrossRef] [PubMed]

100. Wasserman, S.; Meintjes, G.; Maartens, G. Linezolid in the treatment of drug-resistant tuberculosis: The challenge of its narrow therapeutic index. *Expert Rev. Anti. Infect. Ther.* 2016, 14, 901–915. [CrossRef] [PubMed]

101. Parumasivam, T.; Chan, J.G.; Pang, A.; Quan, D.H.; Triccas, J.A.; Britton, W.J.; Chan, H.K. In vitro evaluation of novel inhalable dry powders consisting of thioridazine and rifapentine for rapid tuberculosis treatment. *Eur. J. Pharm. Biopharm.* 2016, 107, 205–214. [CrossRef] [PubMed]

102. Vibe, C.B.; Fenaroli, F.; Pires, D.; Wilson, S.R.; Bogoeva, V.; Kalluru, R.; Speth, M.; Anes, E.; Griffiths, G.; Hildahl, J. Thioridazine in PLGA nanoparticles reduces toxicity and improves rifampicin therapy against mycobacterial infection in zebrafish. *Nanotoxicology* 2016, 10, 680–688. [CrossRef] [PubMed]

103. Winters, N.; Butler-Laporte, G.; Menzies, D. Efficacy and safety of World Health Organization group 5 drugs for multidrug-resistant tuberculosis treatment. *Eur. Respir. J.* 2015, 46, 1461–1470. [CrossRef] [PubMed]

© 2017 by the authors; licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC-BY) license (http://creativecommons.org/licenses/by/4.0/).