

# Outcomes of Bedaquiline Treatment in Patients with Multidrug-Resistant Tuberculosis

Lawrence Mbuagbaw, Lorenzo Guglielmetti, Catherine Hewison, Nyasha Bakare, Mathieu Bastard, Eric Caumes, Mathilde Fréchet-Jachym, Jérôme Robert, Nicolas Veziris, Naira Khachatryan, Tinatin Kotrikadze, Armen Hayrapetyan, Zaza Avaliani, Holger J. Schünemann, Christian Lienhardt

Bedaquiline is recommended by the World Health Organization for the treatment of multidrug-resistant (MDR) and extensively drug-resistant (XDR) tuberculosis (TB). We pooled data from 5 cohorts of patients treated with bedaquiline in France, Georgia, Armenia, and South Africa and in a multicountry study. The rate of culture conversion to negative at 6 months (by the end of 6 months of treatment) was 78% (95% CI 73.5%–81.9%), and the treatment success rate was 65.8% (95% CI 59.9%–71.3%). Death rate was 11.7% (95% CI 7.0%–19.1%). Up to 91.1% (95% CI 82.2%–95.8%) of the patients experienced  $\geq 1$  adverse event, and 11.2% (95% CI 5.0%–23.2%) experienced a serious adverse event. Lung cavitations were consistently associated with unfavorable outcomes. The use of bedaquiline in MDR and XDR TB treatment regimens appears to be effective and safe across different settings, although the certainty of evidence was assessed as very low.

In 2017, there were  $\approx 10$  million (range, 9.0–11.1 million) new cases of tuberculosis (TB) worldwide, of which  $\approx 558,000$  were rifampin-resistant TB (RR TB) or multidrug-resistant TB (MDR TB) (1). MDR TB refers to resistance to isoniazid and rifampin, 2 of the most powerful TB drugs, with or without resistance to other first-line drugs. Extensively drug-resistant tuberculosis (XDR TB), a more severe form of drug-resistant TB, is defined as MDR TB with additional resistance to any fluoroquinolone and to any of the 3 second-line injectables (amikacin, capreomycin, or kanamycin) (2). Treatment outcomes in patients with MDR TB are generally poor, with treatment success in about half of those who receive treatment (56.4%), and much worse in patients with XDR TB (3). In 2017, MDR TB and RR TB caused  $\approx 230,000$  deaths (1). The treatment of MDR and XDR TB is complex and expensive, requiring the use of  $\geq 4$  medications considered to be active in longer regimens (18–20 months) (4–6), and is fraught with many adverse events that can be debilitating or life threatening (7,8).

Bedaquiline is a new compound belonging to the diarylquinoline class used to treat MDR TB; cure and culture conversion rates using bedaquiline are promising (9,10). A recent cost-effectiveness analysis showed that bedaquiline added to a background MDR TB regimen would improve health outcomes and reduce costs in high TB burden countries (11). Bedaquiline received accelerated approval in the United States in 2012 for the treatment of pulmonary MDR TB as part of an appropriate combination therapy in adult patients with resistance or intolerance to other treatment regimens. However, in 2013, limited data and concerns about higher death rates among patients who received bedaquiline in the phase II randomized controlled trial (10) led the World Health Organization (WHO) to issue an interim conditional recommendation on its use under specific conditions: proper patient inclusion, signed informed consent, adherence to the WHO-recommended principles of designing an MDR TB regimen, close monitoring, and active pharmacovigilance (12). Since 2013, many countries

Author affiliations: St. Joseph's Healthcare Hamilton, Hamilton, Ontario, Canada (L. Mbuagbaw); Centre for the Development of Best Practices in Health, Yaoundé, Cameroon (L. Mbuagbaw); McMaster University, Hamilton (L. Mbuagbaw, H.J. Schünemann); Centre d'Immunologie et des Maladies Infectieuses, INSERM, Paris (L. Guglielmetti, E. Caumes, J. Robert, N. Veziris); Centre Hospitalier de Bligny, Bris-sous-Forges, France (L. Guglielmetti, M. Fréchet-Jachym); Sorbonne Université, Paris, France (L. Guglielmetti, J. Robert, N. Veziris); Médecins Sans Frontières, Paris (C. Hewison); Janssen Research & Development, LLC, Titusville, New Jersey, USA (N. Bakare); Epicentre, Paris (M. Bastard); Hôpitaux Universitaires de l'Est Parisien, Paris (N. Veziris); Médecins Sans Frontières, Yerevan, Armenia (N. Khachatryan); Médecins Sans Frontières, Tbilisi, Georgia (T. Kotrikadze); National Tuberculosis Control Centre, Yerevan (A. Hayrapetyan); National Centre for Tuberculosis and Lung Disease, Tbilisi (Z. Avaliani); World Health Organization, Geneva, Switzerland (C. Lienhardt); Université de Montpellier, Montpellier, France (C. Lienhardt)

DOI: <https://doi.org/10.3201/eid2505.181823>

have introduced bedaquiline as part of their management strategy for MDR TB, and in 2018, WHO updated its guidance for the use of bedaquiline in MDR TB, including children  $\geq 6$  years of age (6).

In 2016, to update the interim guidance, the WHO Guideline Development Group (GDG) conducted a review of newly available data on the use of bedaquiline in the treatment of MDR TB (13). After the publication of the GDG report, further outcome data were retrieved from cohorts in Armenia and Georgia. We report the results of the updated analysis from 5 cohorts of patients with MDR TB treated with bedaquiline, taking into account both study-level and patient-level characteristics on outcomes, including deaths. We also report on the use of bedaquiline in research and nonresearch settings and on adverse events.

## Materials and Methods

### Data Sources

We pooled data from 5 cohorts of patients with MDR or XDR TB treated with bedaquiline as part of compassionate use, programmatic use, expanded access, or research programs. The processes and methods used to search, screen, and select studies have been reported previously (13). In brief, studies and datasets were considered only if they met the following inclusion criteria: participants received a diagnosis of MDR TB and were treated with bedaquiline for  $\geq 6$  months as part of an anti-TB regimen. We excluded studies of bedaquiline-only therapy, studies not providing details of the background regimens, studies not providing outcome information, and studies with  $< 10$  participants. We contacted national TB programs, nongovernmental organizations, and the drug manufacturer, Janssen Therapeutics (<https://www.janssen.com>), for unpublished data that fit the criteria.

Our search retrieved 674 studies, of which only 5 were eligible. The 5 datasets that were finally included for the individual patient data (IPD) meta-analyses originated from Médecins Sans Frontières, which contributed 2 cohorts on behalf of the national TB programs of Armenia and Georgia; the national TB program of South Africa; the Hospital of Bligny, France; and Janssen Therapeutics. All the cohort studies have been published in complete form (14–17) (Table 1).

### Data Management

We invited the investigators of each study to provide data on the basis of a formal data sharing agreement. We used only anonymized data in this study. We cleaned, recoded, and merged data from the 5 cohorts and saved the data on a secure server at the biostatistics unit of St Joseph's Healthcare Hamilton/McMaster University (Hamilton, Ontario, Canada). We contacted investigators of each study to ensure accuracy after recoding. We modified categorical variables to match the most commonly used format to ensure consistency across studies. For example, for data from chest radiography, the presence or absence of lung cavitation was the most commonly reported format for findings, so we collapsed data pertaining to the site of the cavitation (left or right lung) or the number of cavitations. We corrected QT intervals for heart rate using the Fridericia formula (QTcF) (18). We categorized drug resistance in order of increasing severity as MDR TB (resistance to isoniazid and rifampin, with or without resistance to other first-line drugs), MDR TB + FLQ (additional resistance to fluoroquinolones), MDR TB + INJ (additional resistance to second-line injectables), and XDR (resistance to at least isoniazid and rifampin, and to any fluoroquinolone, and to any second-line injectables).

### Outcomes

Using standard WHO definitions, we measured the following treatment outcomes: cure, treatment completion, treatment success (the sum of cure and treatment completion), loss to follow-up, and death (19). We computed culture conversion at 6 months as 2 consecutive cultures, taken  $\geq 30$  days apart, found to be negative before or at the end of the sixth month of treatment. Adverse event severity and seriousness were defined as by the US Food and Drug Administration (20), or as reported by investigators. We measured the following adverse event outcomes: any adverse event, any serious adverse event, number of adverse events by body system, and QT interval prolongation (highest recorded QTcF value and increase from baseline).

### Statistical Methods

We summarized baseline data as mean ( $\pm$  SD) for continuous variables and frequency (%) for categorical variables.

**Table 1.** Characteristics of cohorts in study of bedaquiline treatment for multidrug-resistant tuberculosis

Cohort	Design	Location	Sample size	Inception date	Type of care
Pym 2013 (14)	Phase II, single-arm open-label trial	31 sites, 11 countries*	233	2009 Aug	Research
Guglielmetti 2017 (15)	Retrospective cohort	France	45	2010 Jan	Expanded access
Ndjeka 2018 (16)	Prospective cohort	South Africa	195	2013 Mar	Compassionate use
Hewison 2018 (17)	Prospective cohort	Armenia	62	2013 Apr	Compassionate use
	Prospective cohort	Georgia	30		Compassionate use (20), programmatic use (10)

\*China, Estonia, Republic of Korea, Latvia, Peru, Philippines, Russian Federation, South Africa, Thailand, Turkey, Ukraine.

We conducted a random effects meta-analysis of proportions in the first instance to pool effect sizes for effectiveness and safety. The random effects model incorporates the heterogeneity between studies and redistributes the weights of the studies based on this heterogeneity. We assessed statistical heterogeneity using the  $I^2$  statistic, a measure of heterogeneity between studies. We reported variables and outcomes with various levels of completeness and highlighted them where appropriate. For example, we computed culture conversion only if cultures were examined at the sixth month and not later. Similarly, we used baseline QTcF data only if data were collected within 1 month of starting bedaquiline. We used generalized estimation equations to model the effect of individual- and study-level characteristics on outcomes. We built separate models for the dependent binary (yes/no) variables, culture conversion at 6 months, treatment success, and death, using an unstructured correlation matrix and the logit link. The independent variables (age, sex, HIV status, presence of lung cavitations, severity of drug resistance, and previous use of second-line drugs) are all known to affect outcomes in TB (21,22). We assessed model fit using the quasi-likelihood under independence model criterion and set the level of statistical significance at  $\alpha = 0.05$ . We used different numbers of patients for each analysis because of variations in completeness and availability of data. Samples used for each analysis are shown in Appendix Table 1 (<https://wwwnc.cdc.gov/EID/article/25/5/18-1823-App1.pdf>).

### Certainty Evaluation

We assessed the certainty of the evidence using the Grading of Recommendations, Assessment, Development, and Evaluation (GRADE) approach, which categorizes each outcome by how confident we are that the effect estimate is close to the quantity of interest (23). Using this approach, the certainty rating across studies can be high, moderate, low, or very low. We summarized the results and certainty as evidence profiles.

### Results

We included a total of 537 participants in the data analysis. Baseline characteristics are shown in detail by cohort and overall in Table 2. The mean age was 36.4 years (SD 11.8). Two thirds of the participants were men (342; 63.7%); 138 (25.7%) were HIV positive; 341 (99.7%) had pulmonary TB, 253 (73.9%) with lung cavities; and 188 (35.0%) had XDR TB. The key differences between the datasets were a higher proportion of male participants in the cohorts from France and Armenia; 63.1% participants having concurrent HIV in the South Africa cohort; and complete outcome data being available from only 51% of all participants because others were still receiving treatment at the time we collected data. Of note, 36 (6.7%) patients received bedaquiline for  $\geq 6$  months.

The baseline regimens we used in the cohorts varied according to local treatment guidelines, drug susceptibility results, or both. Lamivudine, nevirapine, efavirenz, and tenofovir were the most frequently used drugs

**Table 2.** Baseline characteristics of participants in study of bedaquiline treatment for multidrug-resistant tuberculosis\*

Variable	Cohort					Total, n = 537
	South Africa, n = 195	France, n = 45	Janssen, n = 205	Armenia, n = 62	Georgia, n = 30	
Mean age, y (SD)	35.8 (11.2)	37.4 (12.1)	34.9 (12.2)	41.6 (12.6)	38.7 (11.9)	36.4 (11.8)
Sex, no. (%)						
M	98 (50.3)	36 (80.0)	132 (64.4)	55 (88.7)	21 (70.0)	342 (63.7)
F	97 (49.7)	9 (20.0)	73 (35.6)	7 (11.3)	9 (30.0)	195 (36.3)
Mean no. months on BDQ (SD)	5.8 (1.2)	12.3 (7.0)	5.9 (1.1)	5.6 (1.6)	6.0 (1.3)	6.37 (2.3)
No. on BDQ >6 mo (%)	4 (2.1)	32 (71.1)	0.0	0.0	0.0	36 (6.7)†
Mean total treatment duration, mo (SD)	14.9 (6.7)	19.4 (4.7)	21.8 (7.6)	20.2(7.4)	14.0 (6.1)	18.47 (6.9)†
No. (%) with treatment outcome available	101 (51.8)‡	45 (100.0)	205 (100.0)	62 (100.0)	30 (100.0)	443 (82.5)
No. (%) HIV positive‡	120 (63.1)	2 (4.4)	8 (4.0)	4 (6.5)	1 (3.3)	135 (25.1)
No. (%) on antiretroviral therapy	110 (56.4)	2 (4.4)	0.0	0	0	112 (20.9)
Type of TB, no. (%)						
Pulmonary	NR	44 (97.8)	205 (100.0)	62 (100.0)	30 (100.0)	341 (99.7)
Extrapulmonary	NR	8 (17.8)	0	0	0	8 (2.3)
No. (%) with previous TB treatment	NR	34 (75.6)	193 (94.1)	62 (100.0)	29 (96.7)	271 (79.2)
No. (%) with previous second-line TB treatment	NR	27 (60.0)	177 (86.3)	62 (100.0)	29 (96.7)	295 (86.3)
No. (%) with lung cavities on chest radiograph	NR	39 (86.7)	135 (65.8)	55 (88.7)	24 (80.0)	253 (73.9)
Resistance profile, no. (%)§						
MDR TB	0	7 (15.6)	93 (45.4)	6 (9.7)	0	100 (18.6)
MDR TB + FQ	73 (37.4)	8 (17.8)	31 (15.1)	26 (41.9)	5 (16.7)	147(27.3)
MDR TB + INJ	29 (14.9)	6 (13.3)	13 (6.3)	7 (11.3)	0	55 (10.2)
XDR TB	77 (39.5)	24 (53.3)	37 (18.0)	23 (37.1)	25 (83.3)	188 (35.0)

\*BDQ, bedaquiline; FQ, fluoroquinolone; INJ, injectable; MDR, multidrug resistant; NR, not reported; TB, tuberculosis; XDR, extensively drug resistant.

†Missing data: South Africa = 15.

‡Missing data: South Africa = 17; Janssen (drug manufacturer) = 7.

§Missing data: South Africa = 16; Janssen (drug manufacturer) = 31.

**Table 3.** Multivariable analyses for key outcomes in study of bedaquiline treatment for MDR TB\*

Covariate	Culture conversion at 6 mo, n = 318		Success, n = 325		Death, n = 325	
	Adjusted OR (95% CI)	p value	Adjusted OR (95% CI)	p value	Adjusted OR (95% CI)	p value
Male sex	1.25 (0.65–2.41)	0.499	1.27 (0.74–2.15)	0.382	0.60 (0.24–1.47)	0.264
Age, y	1.01 (0.99–1.04)	0.342	0.99 (0.98–1.01)	0.550	1.05 (1.01–1.09)	0.010
HIV positive	0.42 (0.13–1.39)	0.155	0.35 (0.12–0.99)	0.050	0.97 (0.09–10.05)	0.982
Resistance profile†	0.57 (0.43–0.76)	<0.001	0.84 (0.68–1.04)	0.110	1.14 (0.73–1.79)	0.562
Presence of lung cavitation	0.30 (0.13–0.70)	0.004	0.38 (0.21–0.68)	0.001	5.31 (1.25–22.52)	0.023
Previous use of second-line drugs	0.67 (0.22–2.01)	0.437	0.73 (0.33–1.59)	0.423	1.22 (0.29–5.15)	0.783

\*Absence of data on cavitation precluded use of data from South Africa. MDR, multidrug resistant; TB, tuberculosis; XDR, extensively drug resistant.

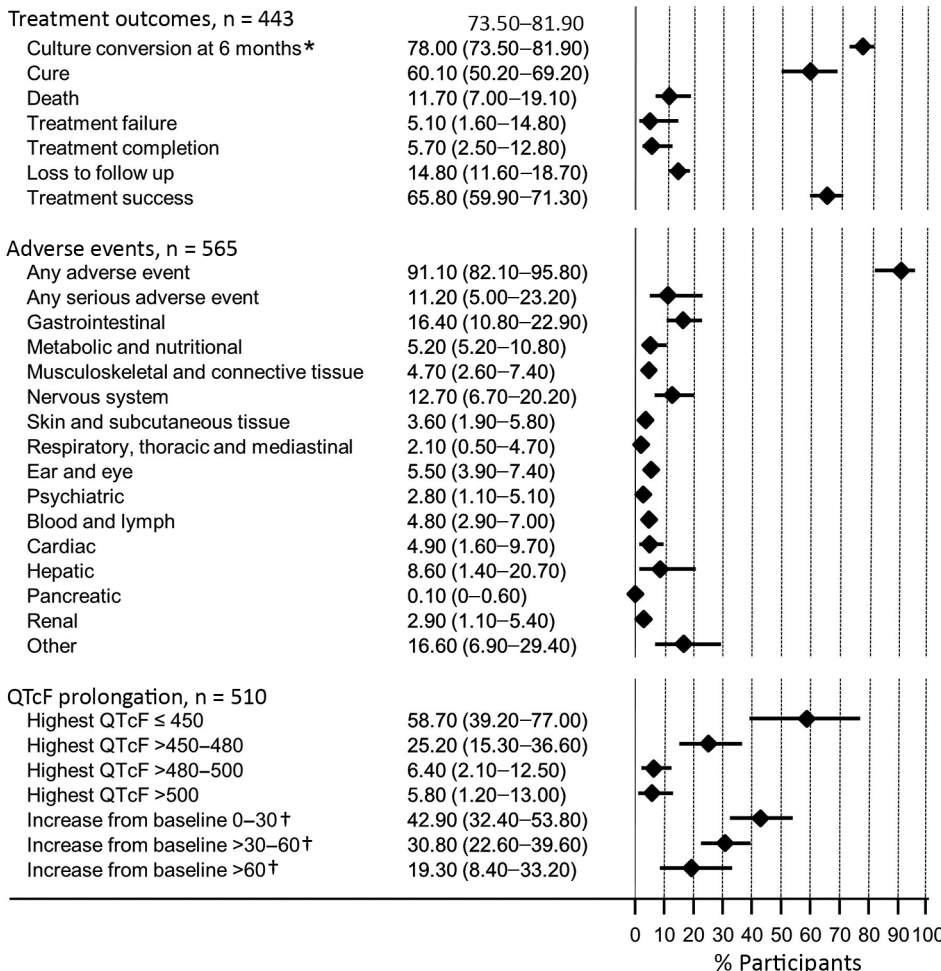
†Resistance profiles used in this study were MDR TB (reference), MDR TB plus fluoroquinolone, MDR TB plus injectable drugs, and XDR TB.

among the patients on antiretroviral therapy (Appendix Tables 2, 3).

We computed culture conversion at 6 months only for patients who had a positive sputum culture at baseline and 2 consecutive culture readings  $\geq 30$  days apart, the last taken at the end of the sixth month. Thirty-seven patients did not have sufficient culture data. Of 406 patients with sufficient culture data, the overall culture conversion rate at 6 months was 78.0% (95% CI 73.5%–81.9%;  $I^2 = 46\%$ ). A total of 443/537 (82.5%) participants had end-of-treat-

ment outcome data: cure, 60.1% (95% CI 50.2%–69.2%;  $I^2 = 66\%$ ); treatment success, 65.8% (95% CI 59.9%–71.3%;  $I^2 = 38\%$ ); death, 11.7% (95% CI 7.0%–19.1%;  $I^2 = 71\%$ ); treatment failure, 5.1% (95% CI 1.6%–14.8%;  $I^2 = 73\%$ ); and loss to follow-up, 14.8% (95% CI 11.6%–18.7%;  $I^2 = 7\%$ ).

Safety data were available from a total of 565 participants, including an additional 28 participants from the Janssen cohort who were not eligible for efficacy analyses because they were missing a confirmation of



**Figure.** Summary of treatment outcomes and adverse events in study of bedaquiline treatment for multidrug-resistant tuberculosis. Values are shown as percent with 95% CI, shown in the graph as horizontal bars. QTcF indicates QT intervals corrected for heart rate using the Fridericia formula. \*A total of 406 study participants with culture data at the 6-month point; † a total of 509 participants with baseline QTcF data.

MDR ( $n = 3$ ) or a positive culture at baseline ( $n = 25$ ). Of these participants, 91.1% (95% CI 82.2%–95.8%) experienced occurrence of any adverse event and 11.2% (5.0%–23.2%) occurrence of any serious adverse event. The most frequent adverse events were gastrointestinal (16.4%; 95% CI 10.8%–22.9%), nervous system (12.7; 95% CI 6.7%–20.2%), and hepatic (8.6%; 95% CI 1.4%–20.7%) (Figure).

We found that 5.8% of 510 participants (95% CI 1.2%–13.0%;  $I^2 = 84\%$ ) had a highest QTcF  $>500$  ms and 19.3% of 509 (95% CI 8.4%–33.2%;  $I^2 = 93\%$ ) had an increase in QTcF from baseline of more  $>60$  ms. Despite the small sample of patients receiving bedaquiline for a prolonged period (i.e.,  $>6$  months), data seem to indicate an absence of effect of exposure to bedaquiline for  $>6$  months on QTc prolongation  $>500$  ms.

### Adjusted Analyses

Culture conversion at 6 months was less likely in patients with a more severe resistance profile (aOR 0.57, 95% CI 0.43–0.76;  $p < 0.001$ ) and with lung cavitations (aOR 0.30, 95% CI 0.13–0.70;  $p = 0.004$ ). Treatment success was less likely in patients with lung cavitations (aOR 0.38, 95% CI 0.21–0.68;  $p = 0.001$ ) and in those with HIV infection (aOR 0.35, 95% CI 0.12–0.99;  $p = 0.05$ ). The presence of lung cavitations was associated with death (aOR 5.31, 95% CI 1.25–22.52;  $p = 0.023$ ) (Table 3).

The GRADE evidence profile is reported in Appendix Table 4. The GDG judged evidence for all outcomes to be of very low certainty; reasons were the risk for bias (lack of control data), risk for inconsistency (considerable statistical heterogeneity), imprecision (wide confidence intervals), and indirectness (variations in adverse event definition) (13).

### Discussion

Inclusion of bedaquiline for  $\geq 6$  months in the treatment regimen was associated with good outcomes in these cohorts, with 78% culture conversion at 6 months and a 65.8% treatment success rate, indicating a favorable efficacy of this medicine. Observed death rate was 11.7%. Although almost all patients experienced at least an adverse event (91.1%), only 11.2% experienced a serious one. Only 5.8% of patients had a highest recorded value of QTcF  $>500$  ms and 19.3% had an increase QT from baseline of more than 60 ms. Being older, having a more severe resistance profile, concurrent HIV, and lung cavitation were associated with unfavorable outcomes.

Such results compare favorably with those observed in large cohorts of patients with MDR TB in the prebedaquiline era (with success rates of 54%–58% and death rates of 13.8%–15%) (7,24), thus indicating a beneficial effect of the addition of bedaquiline to background MDR/XDR TB

regimens. Our findings are in line with those from a large South Africa cohort in which the additional use of bedaquiline reduced the risk for death in patients with MDR TB, compared with standard regimens (25). In a matched subset of patients from the South Africa cohort, switching from second-line injectables (because of intolerance) to bedaquiline led to fewer unfavorable outcomes (death, loss to follow-up, or treatment failure) (26). Likewise, an individual patient data meta-analysis of 50 studies (including this study's cohort data from France, South Africa, and Janssen Therapeutics) reported lower odds of death with the use of bedaquiline (27). The lower mortality rate we observed strengthens the case for the use of bedaquiline in patients with MDR TB. On the basis of data cumulated since 2012 from various observational and programmatic studies, including the South Africa cohort (25), WHO has now consolidated its recommendation for the use of bedaquiline, now proposed as a Group A drug (medicines to be prioritized) in longer MDR TB regimens (6).

Very few patients in our study had a highest recorded value of QTcF  $>500$  ms. Other studies have found similarly low rates of cardiotoxicity (28). This finding could therefore alleviate some of the concerns around the risk for cardiotoxicity related with the use of bedaquiline (28).

Our study adds some information about the use of bedaquiline in persons living with HIV and indicates that favorable outcomes may be more challenging to achieve in these patients, bearing in mind that not all of them were on antiretroviral therapy as recommended by WHO. This information is important for countries with a high prevalence of HIV/TB co-infection. However, other reports indicate that the reductions in number of deaths are similar or better in persons living with HIV (13,25).

Our findings are similar to those observed in other large multicenter studies in terms of success, death, and adverse events (4), although higher culture conversion rates at 6 months were reported from studies in Belarus and Germany (29,30). Our findings should be interpreted with due consideration of the severity of illness in this cohort, in which up to 35% of patients had XDR TB, 86.3% previous second-line treatment, and 73.9% lung cavitations, suggesting that a success rate of 65.8% is probably higher than would otherwise be expected.

Currently, WHO does not recommend the routine use of bedaquiline for longer than 6 months (6). However, WHO acknowledges that “clinicians and national TB programs may be compelled to use [bedaquiline] beyond 24 weeks in selected MDR TB patients (including those with additional drug resistance), if the regimen is unlikely to achieve cure or poses a risk [of] creating additional drug resistance” (31). We were unable to properly investigate the effect of prolonged bedaquiline use in this study, given the small number of patients (6.7%) who received the

medicine for >6 months and the risk for bias by indication (bedaquiline offered to patients with more severe disease). However, we found no correlation between QTcF prolongation and extended bedaquiline use.

Overall, our data confirm that bedaquiline should be a Group A drug in the treatment of MDR TB, as currently recommended by WHO (6), for not only its additive value in culture conversion and treatment success, but also its safety in varied settings.

This study has some limitations. First, cohorts differed greatly in completeness and quality of data, because they were not all initially designed for research purposes. Some variables were defined differently and the timing and number of follow-up visits varied. To maximize the use of data, we tried to work with variables that were reported across all datasets. These methodological differences, in addition to baseline differences by settings (prevalence of HIV infection, provision of baseline regimens and antiretroviral therapy), probably caused high levels of heterogeneity reflected in the meta-analyses. Second, the absence of certain variables from given datasets precluded their use in the adjusted analyses. For example, at the time of these analyses, data on pulmonary cavitation or prior use of second-line drugs were not available from the South Africa cohort, and we had end-of-treatment data for only 51.8% of those patients. Third, the absence of comparative data from patients who did not receive bedaquiline limits the inferences that can be drawn from these data. Finally, we were unable to report a causality assessment of the adverse events. Because of these limitations, the certainty of evidence was rated by the independent GDG as very low for all outcomes for the purpose of GRADE evaluation in the June 2016 meeting (13). Although these concerns appear to limit the credibility of these findings, they represent a real-world picture of the use of bedaquiline under programmatic conditions, outside of research settings.

The strength of this work lies in the use of patient-level data and in the random effects approach used for analyses that embraces the heterogeneity across cohorts. Our results represent data from many countries with different income levels, suggesting that the findings are generalizable. However, whereas heterogeneity is duly accounted for, it is not fully explained, and  $I^2$  values >50% warrant further investigation (32). Some study-level characteristics may contribute to the high levels of heterogeneity, such as the prevalence of HIV, extent of drug resistance, delivery of care, and data quality. Another strength of this work is the detailed information on adverse events by system, particularly the QTcF measurements, which are not usually captured in large databases and registries.

Despite the study strengths, some questions still remain unanswered, such as the role of prolonged use of

bedaquiline and how best to report safety data on ECG measurements, given the heterogeneity in timing of QTcF measurements. Further studies including data on patients using bedaquiline for >6 months are warranted, as well as studies on the use of bedaquiline in shorter treatment regimens.

In conclusion, these pooled data from 5 cohorts of patients treated with bedaquiline suggest that this drug is effective and safe across different modalities of delivery and in different settings, when added to standard background regimens. Outcomes are less favorable, however, in patients with lung cavitations and more severe drug resistance. The overall certainty of the evidence is very low.

### Acknowledgments

We thank L. Thabane, M. Loeb, J. Beyene, and D. Mertz for their guidance on statistics and reporting. We thank Nobert Ndjeka, Francesca Conradie, and Kate Schnippel for supporting the acquisition and transfer of data from the South Africa cohort.

For their contributions to data collection and patient care, we thank L. Yegiazaryan, T. Aydinyan, N. Saribekyan, H. Atchemyan, O. Kirakosyan, N. Danielyan, A. Melikyan, O. Petrosyan, A. Serobyanyan, N. Kiria, K. Barbakadze, L. Mikiashvili, M. Oniani, N. Chumburidze, M. Kikvidze, M. Gergedava, G. Salia, G. Zaridze, and A. Gegeshidze.

Part of this study was carried out under YODA project no. 2016-0734, using data obtained from the Yale University Open Data Access Project, which has an agreement with Janssen Research & Development, LLC. The interpretation and reporting of research using these data are solely the responsibility of the authors and does not necessarily represent the official views of the Yale University Open Data Access Project or Janssen Research & Development, LLC. The Bill and Melinda Gates Foundation supported this study as part of the WHO guidelines development process through grant project no. OPP1126615.

Conflict of interest: L.G., J. R., and N.V. work in a laboratory that has received grants from Johnson & Johnson to perform preclinical studies on bedaquiline. N.B. is an employee of Johnson & Johnson, the manufacturer of bedaquiline.

Authors' contributions: L. M. contributed to data acquisition, data analyses, interpretation; wrote and revised the manuscript; and approved the final version. L.G., C.H., N.B., M.B., M.F.J., J.R., N.V., N.K., T.K., A.H., and Z.A. contributed to data acquisition from the primary studies, interpretation of data, manuscript writing, and approval of the final version. H.J.S. and C.L. contributed to the conception and design of this study, interpretation of the data, and review and approval of the final version. All authors agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

## About the Author

Dr. Mbuagbaw is an assistant professor in the department of health research methods, evidence, and impact at McMaster University and a research methods scientist at St Joseph's Healthcare Hamilton. His primary research interests are infectious diseases and health research methods.

## References

- World Health Organization. Global tuberculosis report 2018. 2018 [cited 2018 Dec 11]. [https://www.who.int/tb/publications/global\\_report](https://www.who.int/tb/publications/global_report)
- World Health Organization. Drug-resistant TB: totally drug-resistant TB FAQ. 2018 [cited 2018 12 Feb]. <http://www.who.int/tb/areas-of-work/drug-resistant-tb/totally-drug-resistant-tb-faq>
- Bonnet M, Bastard M, du Cros P, Khamraev A, Kimenye K, Khurkhumal S, et al. Identification of patients who could benefit from bedaquiline or delamanid: a multisite MDR-TB cohort study. *Int J Tuberc Lung Dis*. 2016;20:177–86. <http://dx.doi.org/10.5588/ijtld.15.0962>
- Borisov SE, Dheda K, Enwerem M, Romero Leyet R, D'Ambrosio L, Centis R, et al. Effectiveness and safety of bedaquiline-containing regimens in the treatment of MDR- and XDR TB: a multicentre study. *Eur Respir J*. 2017;49:1700387. <http://dx.doi.org/10.1183/13993003.00387-2017>
- Falzon D, Schünemann HJ, Harausz E, González-Angulo L, Lienhardt C, Jaramillo E, et al. World Health Organization treatment guidelines for drug-resistant tuberculosis, 2016 update. *Eur Respir J*. 2017;49:1602308. <http://dx.doi.org/10.1183/13993003.02308-2016>
- World Health Organization. WHO treatment guidelines for multidrug- and rifampicin-resistant tuberculosis 2018 update. 2018 [cited 2019 Jan 15]. <https://www.who.int/tb/publications/2018/WHO.2018.MDR-TB.Rx.Guidelines.prefinal.text.pdf>
- Ahuja SD, Ashkin D, Avendano M, Banerjee R, Bauer M, Bayona JN, et al.; Collaborative Group for Meta-Analysis of Individual Patient Data in MDR-TB. Multidrug resistant pulmonary tuberculosis treatment regimens and patient outcomes: an individual patient data meta-analysis of 9,153 patients. *PLoS Med*. 2012;9:e1001300. <http://dx.doi.org/10.1371/journal.pmed.1001300>
- Nathanson E, Gupta R, Huamani P, Leimane V, Pasechnikov AD, Tupasi TE, et al. Adverse events in the treatment of multidrug-resistant tuberculosis: results from the DOTS-Plus initiative. *Int J Tuberc Lung Dis*. 2004;8:1382–4.
- Diacon AH, Dawson R, Von Groote-Bidlingmaier F, Symons G, Venter A, Donald PR, et al. Randomized dose-ranging study of the 14-day early bactericidal activity of bedaquiline (TMC207) in patients with sputum microscopy smear-positive pulmonary tuberculosis. *Antimicrob Agents Chemother*. 2013;57:2199–203. <http://dx.doi.org/10.1128/AAC.02243-12>
- Diacon AH, Pym A, Grobusch MP, de los Rios JM, Gotuzzo E, Vasilyeva I, et al.; TMC207-C208 Study Group. Multidrug-resistant tuberculosis and culture conversion with bedaquiline. *N Engl J Med*. 2014;371:723–32. <http://dx.doi.org/10.1056/NEJMoa1313865>
- Lu X, Smare C, Kambili C, El Khoury AC, Wolfson LJ. Health outcomes of bedaquiline in the treatment of multidrug-resistant tuberculosis in selected high burden countries. *BMC Health Serv Res*. 2017;17:87. <http://dx.doi.org/10.1186/s12913-016-1931-3>
- World Health Organization. The use of bedaquiline in the treatment of multidrug-resistant tuberculosis: interim policy guidance. 2013 [cited 2018 Mar 19]. [http://apps.who.int/iris/bitstream/10665/84879/1/9789241505482\\_eng.pdf](http://apps.who.int/iris/bitstream/10665/84879/1/9789241505482_eng.pdf)
- World Health Organization. Report of the Guideline Development Group Meeting on the use of bedaquiline in the treatment of multidrug-resistant tuberculosis 2016 [cited 2018 Mar 19]. <http://apps.who.int/iris/bitstream/10665/254712/1/WHO-HTM-TB-2017.01-eng.pdf>
- Pym AS, Diacon AH, Tang SJ, Conradie F, Danilovits M, Chuchottaworn C, et al.; TMC207-C209 Study Group. Bedaquiline in the treatment of multidrug- and extensively drug-resistant tuberculosis. *Eur Respir J*. 2016;47:564–74. <http://dx.doi.org/10.1183/13993003.00724-2015>
- Guglielmetti L, Jaspard M, Le Dù D, Lachâtre M, Marigot-Outtandy D, Bernard C, et al.; French MDR-TB Management Group. Long-term outcome and safety of prolonged bedaquiline treatment for multidrug-resistant tuberculosis. *Eur Respir J*. 2017;49:1601799. <http://dx.doi.org/10.1183/13993003.01799-2016>
- Ndjeka N, Schnippel K, Master I, Meintjes G, Maartens G, Romero R, et al. High treatment success rate for multidrug-resistant and extensively drug-resistant tuberculosis using a bedaquiline-containing treatment regimen. *Eur Respir J*. 2018;52:1801528. <http://dx.doi.org/10.1183/13993003.01528-2018>
- Hewison C, Bastard M, Khachatryan N, Kotrikadze T, Hayrapetyan A, Avaliani Z, et al. Is 6 months of bedaquiline enough? Results from the compassionate use of bedaquiline in Armenia and Georgia. *Int J Tuberc Lung Dis*. 2018;22:766–72. <http://dx.doi.org/10.5588/ijtld.17.0840>
- Vandenberk B, Vandael E, Robyns T, Vandenberghe J, Garweg C, Foulon V, et al. Which QT correction formulae to use for QT monitoring? *J Am Heart Assoc*. 2016;5:e003264. <http://dx.doi.org/10.1161/JAHA.116.003264>
- World Health Organization. Companion handbook to the 2011 WHO guidelines for the programmatic management of drug-resistant tuberculosis. 2014 [cited 2018 Mar 16]. [http://www.who.int/tb/publications/pmdt\\_companionhandbook/](http://www.who.int/tb/publications/pmdt_companionhandbook/)
- US Food and Drug Administration. Code of Federal Regulations Title 21. 2017 [cited 2018 Mar 16]. <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfCFR/CFRSearch.cfm?fr=312.32>
- Ali MK, Karanja S, Karama M. Factors associated with tuberculosis treatment outcomes among tuberculosis patients attending tuberculosis treatment centres in 2016–2017 in Mogadishu, Somalia. *Pan Afr Med J*. 2017;28:197.
- Guglielmetti L, Le Dù D, Jachym M, Henry B, Martin D, Caumes E, et al.; MDR-TB Management Group of the French National Reference Center for Mycobacteria and the Physicians of the French MDR-TB Cohort. Compassionate use of bedaquiline for the treatment of multidrug-resistant and extensively drug-resistant tuberculosis: interim analysis of a French cohort. *Clin Infect Dis*. 2015;60:188–94. <http://dx.doi.org/10.1093/cid/ciu786>
- Guyatt G, Oxman AD, Akl EA, Kunz R, Vist G, Brozek J, et al. GRADE guidelines: 1. Introduction—GRADE evidence profiles and summary of findings tables. *J Clin Epidemiol*. 2011;64:383–94. <http://dx.doi.org/10.1016/j.jclinepi.2010.04.026>
- Cegielski JP, Kurbatova E, van der Walt M, Brand J, Ershova J, Tupasi T, et al.; Global PETTS Investigators. Multidrug-resistant tuberculosis treatment outcomes in relation to treatment and initial versus acquired second-line drug resistance. *Clin Infect Dis*. 2016;62:418–30.
- Schnippel K, Ndjeka N, Maartens G, Meintjes G, Master I, Ismail N, et al. Effect of bedaquiline on mortality in South African patients with drug-resistant tuberculosis: a retrospective cohort study. *Lancet Respir Med*. 2018;6:699–706. [http://dx.doi.org/10.1016/S2213-2600\(18\)30235-2](http://dx.doi.org/10.1016/S2213-2600(18)30235-2)
- Zhao Y, Fox T, Manning K, Stewart A, Tiffin N, Khomo N, et al. Improved treatment outcomes with bedaquiline when substituted for second-line injectable agents in multidrug-resistant tuberculosis: a retrospective cohort study. *Clin Infect Dis*. 2018

- Aug 24 [Epub ahead of print]. <https://doi.org/10.1093/cid/ciy727>
27. Ahmad N, Ahuja SD, Akkerman OW, Alffenaar JC, Anderson LF, Baghaei P, et al.; Collaborative Group for the Meta-Analysis of Individual Patient Data in MDR-TB treatment–2017. Treatment correlates of successful outcomes in pulmonary multidrug-resistant tuberculosis: an individual patient data meta-analysis. *Lancet*. 2018;392:821–34. [http://dx.doi.org/10.1016/S0140-6736\(18\)31644-1](http://dx.doi.org/10.1016/S0140-6736(18)31644-1)
  28. Guglielmetti L, Tiberi S, Burman M, Kunst H, Wejse C, Togonidze T, et al. QT prolongation and cardiac toxicity of new tuberculosis drugs in Europe: a Tuberculosis Network European Trialsgroup (TBnet) study. *Eur Respir J*. 2018;52:1800537. <http://dx.doi.org/10.1183/13993003.00537-2018>
  29. Skrahina A, Hurevich H, Falzon D, Zhilevich L, Rusovich V, Dara M, et al. Bedaquiline in the multidrug-resistant tuberculosis treatment: Belarus experience. *Int J Mycobacteriol*. 2016;5 (Suppl 1):S62–3. <http://dx.doi.org/10.1016/j.ijmyco.2016.11.014>
  30. Olaru ID, Heyckendorf J, Andres S, Kalsdorf B, Lange C. Bedaquiline-based treatment regimen for multidrug-resistant tuberculosis. *Eur Respir J*. 2017;49:1700742. <http://dx.doi.org/10.1183/13993003.00742-2017>
  31. World Health Organization. WHO best-practice statement on the off-label use of bedaquiline and delamanid for the treatment of multidrug-resistant tuberculosis. 2017 [cited 2018 May 28]. <http://apps.who.int/iris/bitstream/handle/10665/258941/WHO-HTM-TB-2017.20-eng.pdf>
  32. Higgins JPT, Thompson SG, Deeks JJ, Altman DG. Measuring inconsistency in meta-analyses. *BMJ*. 2003;327:557–60. <http://dx.doi.org/10.1136/bmj.327.7414.557>

Address for correspondence: Lawrence Mbuagbaw, St. Joseph's Healthcare, The Research Institute, Biostatistics Unit, 50 Charlton Ave E, 3rd Fl, Martha Wing, Room H321, Hamilton, Ontario L8N 4A6, Canada; email: mbuagblc@mcmaster.ca



## EMERGING INFECTIOUS DISEASES®

April 2018

# Antimicrobial Resistance

- Seroprevalence of Chikungunya Virus in 2 Urban Areas of Brazil 1 Year after Emergence
- Two Infants with Presumed Congenital Zika Syndrome, Brownsville, Texas, USA, 2016–2017
- Reemergence of Intravenous Drug Use as Risk Factor for Candidemia, Massachusetts, USA
- Rickettsial Illnesses as Important Causes of Febrile Illness in Chittagong, Bangladesh
- Influence of Population Immunosuppression and Past Vaccination on Smallpox Reemergence
- Emerging Coxsackievirus A6 Causing Hand, Foot and Mouth Disease, Vietnam
- Influenza A(H7N9) Virus Antibody Responses in Survivors 1 Year after Infection, China, 2017
- Genomic Surveillance of 4CMenB Vaccine Antigenic Variants among Disease-Causing *Neisseria meningitidis* Isolates, United Kingdom, 2010–2016
- Evolution of Sequence Type 4821 Clonal Complex Meningococcal Strains in China from Prequinolone to Quinolone Era, 1972–2013
- Avirulent *Bacillus anthracis* Strain with Molecular Assay Targets as Surrogate for Irradiation-Inactivated Virulent Spores
- Phenotypic and Genotypic Characterization of *Enterobacteriaceae* Producing Oxacillinase-48–Like Carbapenemases, United States
- Artemisinin-Resistant *Plasmodium falciparum* with High Survival Rates, Uganda, 2014–2016
- Carbapenem-Nonsusceptible *Acinetobacter baumannii*, 8 US Metropolitan Areas, 2012–2015
- Cooperative Recognition of Internationally Disseminated Ceftriaxone-Resistant *Neisseria gonorrhoeae* Strain
- Imipenem Resistance in *Clostridium difficile* Ribotype 017, Portugal
- Enhanced Replication of Highly Pathogenic Influenza A(H7N9) Virus in Humans
- Multidrug-Resistant *Salmonella enterica* 4,[5],12:i:-Sequence Type 34, New South Wales, Australia, 2016–2017
- Genetic Characterization of Enterovirus A71 Circulating in Africa

To revisit the April 2018 issue, go to:

<https://wwwnc.cdc.gov/eid/articles/issue/24/4/table-of-contents>



# Outcomes of Bedaquiline Treatment in Patients with Multidrug-Resistant Tuberculosis

## Appendix

**Appendix Table 1.** Description of samples used for the various analyses of the use of bedaquiline in treating multidrug-resistant tuberculosis

Analysis	Study cohort	Total	Description
Description of baseline characteristics	South Africa = 195 France = 45 Janssen = 205 Armenia = 62 Georgia = 30	537	Total sample with baseline characteristics available
Composition of antiretroviral therapy	South Africa = 110  France = 2 Janssen = 8	120	Data available only for HIV-infected patients on antiretroviral therapy
Effectiveness (sputum culture conversion at 6 mo)	South Africa = 72  France = 41 Janssen = 205 Armenia = 60 Georgia = 28	406	Data on patients who had a culture done at 6 mo
Treatment outcomes (cure, death, lost to follow-up, treatment complete, treatment failure)	South Africa = 101  France = 45 Janssen = 205 Armenia = 62 Georgia = 30	443	Data on cohorts of patients with complete follow-up ( $\geq 18$ mo) and available outcome data
Safety (adverse events)	South Africa = 195 France = 45 Janssen = 233 Armenia = 62 Georgia = 30	565*	Total no. patients who received bedaquiline
Safety (QT prolongation)	South Africa = 141†  France = 45 Janssen = 232 Armenia = 62 Georgia = 30	510	Total number of patients with baseline and follow-up QT data

\* Includes additional data from 28 patients from the Janssen cohort who received bedaquiline, but were later found to be ineligible

† Only 141 from South Africa, 1 of whom did not have baseline data

**Appendix Table 2.** Composition of optimized baseline regimen in studies of the use of bedaquiline in treating multidrug-resistant tuberculosis

Drug	No. in cohort on drug (%)				
	South Africa n = 195	France n = 45	Janssen n = 205	Armenia n = 62	Georgia n = 30
Aminoglycosides	56 (28.7)	45 (100.0)	152 (74.1)	17 (27.4)	1 (3.3)
Amikacin sulfate	1 (0.5)	32 (71.1)	47 (22.9)	1 (1.6)	0 (0.0)
Kanamycin	55 (28.2)	0 (0.0)	103 (50.2)	16 (25.8)	1 (3.3)
Streptomycin	0 (0.0)	45 (100.0)	3 (1.5)	0 (0.0)	0 (0.0)
Fluoroquinolones	158 (81.0)	26 (57.7)	180 (87.8)	28 (45.2)	7 (23.3)
Ciprofloxacin	0 (0.0)	0 (0.0)	7 (3.4)	0 (0.0)	0 (0.0)
Gatifloxacin	0 (0.0)	0 (0.0)	1 (0.5)	0 (0.0)	0 (0.0)
Levofloxacin	158 (81.0)	8 (17.8)	66 (32.2)	28 (45.2)	7 (23.3)
Moxifloxacin	0 (0.0)	24 (53.3)	1 (0.5)	0 (0.0)	0 (0.0)
Ofloxacin	0 (0.0)	0 (0.0)	101 (49.3)	0 (0.0)	0 (0.0)
Sparfloxacin	0 (0.0)	0 (0.0)	5 (2.4)	0 (0.0)	0 (0.0)
Macrolide	19 (9.7)	0 (0.0)	22 (10.7)	0 (0.0)	0 (0.0)
Azithromycin	9 (4.6)	0 (0.0)	16 (7.8)	0 (0.0)	0 (0.0)
Clarithromycin	10 (5.1)	0 (0.0)	6 (2.9)	0 (0.0)	0 (0.0)
Miscellaneous anti-TB drugs	170 (87.2)	45 (100.0)	205 (100.0)	62 (100.0)	30 (100.0)
Ethambutol	103 (52.8)	20 (44.4)	109 (53.2)	3 (4.8)	0 (0.0)
Isoniazid	34 (17.4)	0 (0.0)	30 (14.6)	0 (0.0)	0 (0.0)
Pyrazinamide	159 (81.5)	0 (0.0)	152 (74.1)	7 (11.3)	5 (16.7)
Rifampin	1 (0.5)	19 (42.2)	1 (0.5)	0 (0.0)	0 (0.0)
Amoxicilin clavunate	5 (2.6)	0 (0.0)	20 (9.8)	50 (80.6)	29 (96.7)
Capreomycin	38 (19.5)	3 (6.7)	45 (22.0)	22 (35.5)	11 (36.7)
Clofazimine	151 (77.4)	20 (44.4)	13 (6.3)	51 (82.3)	24 (80.0)
Cycloserine	0 (0.0)	32 (71.1)	53 (25.9)	45 (72.6)	13 (43.3)
Dapsone	0 (0.0)	0 (0.0)	1 (0.5)	0 (0.0)	0 (0.0)
Ethionamide	99 (50.8)	11 (24.4)	84 (41.0)	0 (0.0)	0 (0.0)
Imipenem	3 (1.5)	28 (62.2)	1 (0.5)	0 (0.0)	0 (0.0)
Imipenem-cilastatin	0 (0.0)	0 (0.0)	0 (0.0)	44 (71.0)	27 (90.0)
Linezolid	121 (62.1)	43 (95.6)	12 (5.9)	62 (100.0)	30 (100.0)
Para-aminosalicylic acid	157 (80.5)	40 (88.9)	97 (47.3)	31 (50.0)	12 (40.0)
Meropenem	1 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)
Meropenem/amoxicilin clavunate	0 (0.0)	2 (4.4)	0 (0.0)	0 (0.0)	0 (0.0)
Para-aminosalicylic acid salt	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	1 (3.3)
Prothionamide	0 (0.0)	0 (0.0)	76 (37.1)	10 (16.1)	1 (3.3)
Terizidone	161 (82.6)	0 (0.0)	60 (29.3)	0 (0.0)	0 (0.0)
Thioacetazone	0 (0.0)	0 (0.0)	3 (1.5)	0 (0.0)	0 (0.0)

**Appendix Table 3.** Composition of regimen in patients who were on antiretroviral therapy in study of the use of bedaquiline in treating multidrug-resistant tuberculosis\*

Antiretroviral drug	No. in cohort on drug (%)		
	South Africa, n = 110	France, n = 2	Janssen (n = 8)
Lamivudine	93 (47.7)	2 (4.4)	8 (100.0)
Abacavir	7 (3.6)	1 (2.2)	0 (0.0)
Zidovudine	21 (10.8)	0 (0.0)	2 (25.0)
Stavudine	27 (13.8)	0 (0.0)	0 (0.0)
Efavirenz	7 (3.6)	2 (4.4)	8 (100.0) <sup>†</sup>
Emtricitabine	15 (7.7)	1 (2.2)	0 (0.0)
Lopinavir/ritonavir	27 (14.3)	0 (0.0)	0 (0.0)
Nevirapine	74 (37.9)	0 (0.0)	1 (12.5)
Tenofovir	56 (28.7)	1 (2.2)	6 (75%)

\*No patients in 2 of the 5 study cohorts were on antiretroviral therapy.

<sup>†</sup>In the Drug manufacturer cohort, EFV was used only after BDQ was stopped

**Appendix Table 4.** Grading of Recommendations, Assessment, Development, and Evaluation evidence profile for study of whether addition of bedaquiline to WHO-recommended second-line drug therapy safely improves outcomes for patients with multidrug-resistant tuberculosis

No. and design of studies	Certainty assessment					No. patients (weighted proportions)*	Relative effect, % at 95% CI†	Certainty	Importance
	Risk for bias	Inconsistency	Indirectness	Imprecision	Other considerations				
Culture conversion (Proportion with sputum conversion at 6 mo of bedaquiline (follow up: mean 6 mo))									
5 observational studies	Serious‡	Not serious	Not serious	Not serious	None	322/405 (78.0%)	73.5–81.9	⊕○○○ Very low	Critical
Mortality (follow-up: mean 18.5 mo)									
5 observational studies	Serious‡	Serious§	Not serious	Serious¶	None	49/443 (11.7%)	7.0–19.1	⊕○○○ Very low	Critical
Treatment success (Proportion with treatment complete + cure) (follow-up: mean 18.5 mo)									
5 observational studies	Serious‡	Not serious	Not serious	Not serious	None	290/443 (65.8%)	59.9–71.3	⊕○○○ Very low	Critical
Serious adverse events (no. patients experiencing at least one SAE over total no. patients) (follow-up: mean 18.5 mo)									
5 observational studies	Serious‡	Serious#	Serious**	Serious¶	None	47/565 (11.2%)	5.0–23.2	⊕○○○ Very low	Critical
QTcF prolongation >60 ms from baseline (follow-up: mean 18.5 mo)									
5 observational studies	Serious‡	Not serious††	Not serious	Serious¶	None	75/509 (19.3%)	8.4–33.2	⊕○○○ Very low	Critical
Highest recorded QTcF prolongation >500 (follow-up: mean 18.5 mo)									
5 observational studies	Serious‡	Serious‡‡	Not serious	Not serious	None	24/510 (5.8%)	1.2–13.0	⊕○○○ Very low	Critical

\*Patients were given bedaquiline with background MDR-TB treatment. No patients in the study were on background MDR-TB treatment (regimen drugs recommended by World Health Organization) alone.

†No comparator data were available to determine absolute effect.

‡Downgrading for lack of control data

§Downgrading for considerable statistical heterogeneity:  $I^2 = 71\%$

¶Downgrading for wide confidence intervals

#Downgrading for considerable statistical heterogeneity:  $I^2 = 88\%$

\*\*Downgrading for indirectness because the definition and (inconsistency in) reporting of all adverse events.

††Downgrading for statistical heterogeneity:  $I^2 = 93\%$

‡‡Downgrading for statistical heterogeneity:  $I^2 = 84\%$