

Determinants of Multidrug Resistance Tuberculosis: Systemic Review and Meta-analysis

Kaleab Tesfaye Tegegne¹⁺, Andualem Zenebe¹, Abiyu Ayalew Assefa¹, Mekibib Kassa Tessema², Eleni Tesfaye Tegegne³, Alelign Tadele Abebe⁴, Berhanu Bifato¹, Muse Rike⁵

ABSTRACT

Objective: The aim of this systemic review and meta-analysis was to determine demographic, behavioral and clinical risk factors for multiple drug-resistant 19 studies were included in the systemic review and meta-analysis.

Materials and Methods: Search engines; PubMed/Medline, Web of Science, Scopus and Google scholar were used to search related literatures on MDR TB and selected determinates of MDRTB. All data were independently extracted. We used a random-effects model according to the DerSimonian and Laird method.

Results: The risk of having MDR-TB in patients who have informal education was 0.67 times lower than that of patients who have formal education (RR=0.67; 95% CI, 0.46 to 0.96). The risk of having MDR-TB in patients who have Diabetes Mellitus was 1.33 times higher than that of patients who have no Diabetes Mellitus (RR of 1.33 and 95% CI, 1.01 to 1.77)

Conclusion: This systemic review and meta-analysis indicated that the risk of having MDR-TB in patients who have informal education was 0.67 times lower than that of patients who have formal education and the risk of having MDR-TB in patients who have Diabetes Mellitus was 1.33 times higher than that of patients who have no Diabetes Mellitus. Our review did not show the presence of an association between MDR TB and other selected variables gender, smoking and alcohol. We recommend interested researchers on MDR TB to do further research on patient's educational level as a risk factor for MDR-TB in order to explain the variation in MDR TB among patients with formal education and patients with informal education.

Keywords: Multidrug resistance tuberculosis; Alcohol; Smoking; Diabetes mellitus; Gender; Educational status

Introduction

Multidrug resistance tuberculosis is a tuberculosis caused by a tubercle bacillus that is resistant to isoniazid and rifampicin [1]. Worldwide, an estimated 10.0 million people diseased with TB in 2018 [2]. In 2018, WHO reported that there were 484,000 incident cases of MDRTB cases and 44.21 (214,000 deaths) [3]. In the world, MDR-TB among TB patients who are new cases and previously treated cases was estimated to be 3.5% and 20.5% respectively [4]. MDR-TB has received increased attention in Africa and it is under-reported [5-6]. In Sub-Saharan Africa the magnitude reaches 14% of the world new MDR-TB cases [5].

In six countries of Africa including (new/ retreatment % respectively) Angola (2.6/18%), DR Congo (2.2/17%), Kenya (1.3/9.4%), Nigeria (4.3/25%), Somalia (8.7/47%) and Zimbabwe (4.6/14%) [4].

In the African Region, the frequency of resistance per country ranged Burundi in the third position with 66% after Rwanda (87%), Democratic Republic of the Congo (68%) [7].

In Ethiopia prevalence of MDR TB is 2.7% and 14.0% in newly and treated previously TB patients respectively [4].

In Ethiopia, many of the MDR-TB patients are undiagnosed due to the poor socioeconomic

¹Department of Public Health, Hawassa College of Health Science, Hawassa, Ethiopia

²Leishmania Research and Treatment Center, University of Gondar, Gondar, Ethiopia

³College of Medicine and Health Science, School of Nursing, University of Gondar, Gondar, Ethiopia

⁴Department of Medical Laboratory Technology, Hawassa College of Health Science, Hawassa, Ethiopia

⁵Hawassa College of Health Science, Research and publication Directorate

[†]Author for correspondence: Kaleab Tesfaye Tegegne, Department of Public Health, Hawassa College of Health Science, Hawassa, Ethiopia; Email: kaleabtesfaye35@gmail.com

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status of people, low level of knowledge and decreased access to health service.

In Ethiopia the number of patients tested for MDR-TB was <1% of new cases and <4% of retreatment cases [8].

Previous studies identified some risk factors associated with MDRTB like gender [9,10], smoking or other substance misuse and diabetes [11-14].

Previous studies in Ethiopia and China also reported that HIV/AIDS smoking cigarette, and drinking alcohol, overcrowding, and poor adherence to DOTS (Directly Observed Treatment Short-course) program are factors associated with MDR-TB [15-20].

Few authors worldwide had the concern of investigating the patient's educational level as a risk factor for MDR-TB [21,22].

The aim of this systemic review and meta-analysis was to determine demographic, behavioral and clinical risk factors for multiple drug-resistant TB and for this we have selected two demographic variables gender and educational status, two behavioral factors alcohol and smoking and one clinical factor that is Diabetes Mellitus.

Methodology

Literature search strategy

These systematic reviews and meta-analyses were done in line with PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) Guideline [23] and we have done the search for different literatures that include data on magnitude of MDR-TB in relation to two socio demographic variables gender and educational status, two behavioral variables alcohol and smoking and one clinical variable, Diabetes Mellitus.

Search engines; PubMed/Medline, Web of Science, Scopus and Google scholar were used to search related literatures on MDR TB and selected determinates of MDRTB.

Combination of words used to search were; "MDRTB", "Gender and MDRTB", "Diabetes Mellitus and MDRTB", "alcohol and MDRTB", "educational status and MDRTB" and "smoking and MDRTB."

In addition, the reference of selected articles was used to search additional studies not found in our initial search.

Selection/eligibility criteria

Studies that reported five selected determinants of MDR-TB and this are gender, educational status, alcohol, smoking and Diabetes Mellitus and published in English language regardless of the design and setting were included. Studies were included in the review irrespective of their study year.

Outcome of interest

MDR-TB is TB that caused by strains of M. *tuberculosis* that are resistant to isoniazid and RMP [4,24].

Data extraction and quality assessment

The titles and abstracts were assessed for reporting the outcome and for fulfilling the eligible criteria. From studies that fulfill the eligible criteria the following data was extracted using prepared form: Name of author, prevalence of MDR-TB in the selected variables gender, smoking, alcohol, educational status and Diabetes Mellitus.

All data were independently extracted. Returned abstracts were reviewed to check whether they contained relevant information.

Each selected research was assessed using the Newcastle-Ottawa Scale (NOS) for assessing the quality of non randomized studies in meta-analyses [25].

Data analysis

An outcome of interest was MDR-TB. We used a random-effects model according to the Der Simonian and Laird method [26].

Heterogeneity was assessed by the I² and values greater than 50% considered representing significant heterogeneity.

When there is heterogeneity between studies we used random-effect models [27]. Results were presented in forest plots. Publication bias was assessed using funnel plots and Beggs and eggers regression test. Analysis was performed using Rev Man software version 5.3 [28].

Results

Studies included

Our initial search using the search terms gave 1170 articles. Out of these, 559 researches which do not match with the aim of the study were excluded after reviewing their titles and abstracts. Then the remaining articles were checked for duplications and 77 articles were selected for full text review. Of the 77 articles reviewed in full text, 19 articles [29-47] were included in the final analysis based on the inclusion-exclusion criteria and quality assessment and 58 studies were removed prior to analysis for reasons of not reporting at least one variable from the five selected variables to be studied as determinants of MDRTB. The selection process of study is presented in (Figure 1).

Description of findings

Nineteen studies were included in the systemic review and meta-analysis [29-47]. Ten studies measured magnitude of MDR-TB among Informal and/or formal educated patients [29,30,32,33,35,37,38,44,46,47].

Sixteen studies measured magnitude of MDR-TB among male and/or female patients [29-40, 43, 45-47].

Seven studies measured magnitude of MDR-TB among Diabetes mellitus and/or no diabetes mellitus patients [29, 32, 36, 38, 40, 42, 45].

Eight studies measured magnitude of MDR-TB among alcohol drinking patients and/or no alcohol drinking patients [29-33,35,45,47].

Thirteen studies measured magnitude of MDR-TB among Smoking patients and/or no Smoking patients [29-33,35,37,38,41,42,45-47].

Gender and MDR TB

The meta-analysis of gender and MDR-TB did not demonstrate a statistically significant association of MDR-TB with Gender (An overall RR of 0.96 and 95% CI, 0.86 to 1.07)

As presented in Figure 2, except for three studies [30,38,39], all other studies showed

no association of MDR-TB to male or female patients.

Heterogeneity testing showed significant variation among the studies $(I^2=53\%)$.

Educational status and MDR TB

The risk of having MDR-TB in patients who have informal education was 0.67 times lower than that of patients who have formal education (overall RR=0.67; 95% CI, 0.46 to 0.96).

The combined effect size indicated that there is a statistically significant association of MDR-TB with educational status of patients, five studies [30,32,37,46,47] were exceptions.

There is heterogeneity between the studies $(I^2=84\%)$ (Figure 3).

Source of heterogeneity for educational status and MDRTB

There is a statistically significant association between effect size and magnitude of informal education as shown in the moderator analysis as prevalence of informal education increase, the effect size for MDRTB decrease (B= -3.67, p value <0.0001)

Assessment of publication bias for educational status and MDRTB

We used Beggs test for asymmetry of funnel plot (p value 0.531) and Eggers regression test (p value 0.903) to test publication bias and both indicated that there is no publication bias for the meta-analysis of educational status and MDRTB.

Alcohol and MDR TB

The meta-analysis of Alcohol and MDR-TB did not also demonstrate a statistically significant

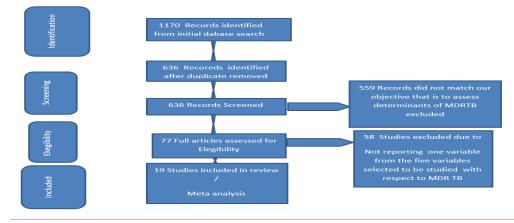


Figure 1: Flow chart for the process of searching, selection and screening of the systemic review and meta - analysis

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	Male	el	Fema	ile		Risk Ratio	Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% Cl
Arnaud IRADUKUNDA et al	41	123	19	57	4.1%	1.00 [0.64, 1.56]	- - -
Assefa et al	109	387	120	323	8.6%	0.76 [0.61, 0.94]	-
B. Baya et al	102	163	32	51	7.9%	1.00 [0.78, 1.27]	+
Elmi et al.	70	217	35	97	5.9%	0.89 [0.64, 1.24]	
Fregona G, et al	132	1123	41	453	5.8%	1.30 [0.93, 1.81]	+
G. Mulisa et al	50	154	38	111	5.6%	0.95 [0.67, 1.34]	-
Glasauer S, et al	354	16556	202	9627	9.8%	1.02 [0.86, 1.21]	+
Herman Owuor Weyenga	50	146	31	97	5.2%	1.07 [0.74, 1.55]	+
LEMA.N et al	71	143	31	61	6.6%	0.98 [0.73, 1.31]	-+
Liang.L et al	171	1385	70	610	7.4%	1.08 [0.83, 1.40]	+
Mekonnen et al	6	85	1	44	0.3%	3.11 [0.39, 25.00]	
Nigus DM, et al	57	363	53	243	5.7%	0.72 [0.51, 1.01]	
Rifat et al	167	609	83	391	8.2%	1.29 [1.03, 1.63]	-
Tenzin C, et al	33	85	46	112	5.5%	0.95 [0.67, 1.34]	-
Vashakidze et al.	189	478	72	127	9.3%	0.70 [0.58, 0.84]	+
Wijayanto et al.	30	60	15	30	4.1%	1.00 [0.65, 1.55]	-
Total (95% CI)		22077		12434	100.0%	0.96 [0.86, 1.07]	•
Total events	1632		889				
Heterogeneity: Tau ² = 0.02; C	hi ² = 31.7	7. df = 1	5 (P = 0.0	07); l ² =	53%		0.01 0.1 1 10 100
Test for overall effect: Z = 0.75		0.01 0.1 1 10 100 Male Female					

Figure 2: Risk ratio of multidrug resistant TB (MDR-TB) in relation to Gender status (Male vs Female), a meta-analysis on Determinates of MDR TB.

	Informal Edu	cation	Formal Edu	cation		Risk Ratio		Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% Cl		M-H, Random, 95% Cl
Arnaud IRADUKUNDA et al	5	34	146	237	8.0%	0.24 [0.11, 0.54]		
Assefa et al	31	105	198	605	12.0%	0.90 [0.66, 1.24]		
Baghaei P, et al.	13	76	34	51	10.3%	0.26 [0.15, 0.44]		
Fregona G, ET AL.	9	88	119	1057	9.4%	0.91 [0.48, 1.73]		_ _
G. Mulisa et al	13	61	75	204	10.5%	0.58 [0.35, 0.97]		
Herman Owuor Weyenga	11	19	70	224	11.1%	1.85 [1.21, 2.85]		
LEMA.N et al	16	35	86	169	11.5%	0.90 [0.61, 1.33]		
Mekonnen et al	2	64	5	60	3.7%	0.38 (0.08, 1.86)		
Rifat M, et al	55	353	195	647	12.3%	0.52 [0.39, 0.68]		-
Tenzin C et al	18	49	61	148	11.3%	0.89 [0.59, 1.35]		
Total (95% CI)		884		3402	100.0%	0.67 [0.46, 0.96]		•
Total events	173		989					-
Heterogeneity: Tau ² = 0.26; C	Chi² = 54.79, df:	= 9 (P < 0	.00001); I ² =	84%				
Test for overall effect: Z = 2.1							0.01	0.1 1 10 100 Informal Education Formal Education

Figure 3: Risk ratio of multidrug resistant TB (MDR-TB) in relation to Gender status (Male vs Female), a meta-analysis on Determinates of MDR TB.

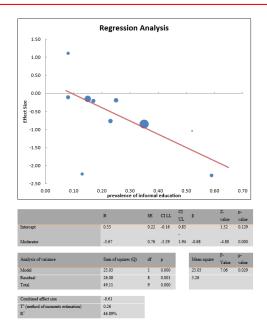


Figure 4: Moderator analysis for educational status and MDRTB.

association of MDR-TB with Drinking Alcohol (overall RR of 0.97 and 95% CI, 0.75 to 1.25).

As presented in Figure 6, except for two studies [32,33], all other studies showed no association of MDR-TB to Alcohol drinking patients or No alcohol drinking patients.

There is Heterogeneity between the studies $(I^2=74\%)$.

Smoking and MDR TB

with smoking status (an overall RR of 1.18 and 95% CI, 0.87 to 1.60)

As presented in Figure 7, except for five studies [29, 32, 38, 42, 47], all other studies showed no association of MDR-TB to smoking patients or no smoking patients.

There is Heterogeneity between the studies $(I^2=87\%)$.

Diabetes Mellitus and MDR TB

The meta-analysis of smoking and MDR-TB did not also demonstrate association of MDR-TB

The meta-analysis of Diabetes Mellitus and MDR-TB showed that there is a statistically

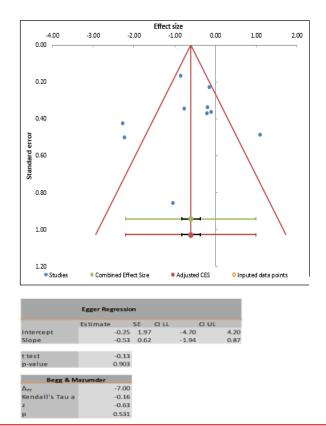


Figure 5: Funnel plot for educational status and MD RTB.

	Alcoh	lol	No Alco	ohol		Risk Ratio		Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% Cl		M-H, Random, 95% Cl
Assefa et al	53	191	176	519	15.1%	0.82 [0.63, 1.06]		
B. Baya et al.	11	18	123	196	12.7%	0.97 [0.66, 1.43]		
Elmi et al.	13	30	92	283	11.7%	1.33 [0.86, 2.07]		
Fregona G,et al	54	321	67	588	13.7%	1.48 [1.06, 2.06]		
G. Mulisa et 58 al	10	72	76	190	9.0%	0.35 [0.19, 0.63]		_ -
Herman Owuor Weyenga	33	109	48	134	13.1%	0.85 [0.59, 1.22]		
RADUKUNDA,A et al	48	126	12	54	9.9%	1.71 [0.99, 2.96]		
LEMA.N et al	46	100	56	104	14.8%	0.85 [0.65, 1.13]		-
Total (95% CI)		967		2068	100.0%	0.97 [0.75, 1.25]		•
Total events	268		650					
Heterogeneity: Tau ² = 0.10;	Chi ² = 26	.61. df:	= 7 (P = 0	.0004);	I² = 74%		<u> </u>	
Test for overall effect: Z = 0.		, -			0.01	0.1 1 10 1 Alcohol No Alcohol		

Figure 6: Risk ratio of multidrug resistant TB (MDR-TB) in relation to drinking alcohol (drinking alcohol vs no drinking alcohol), a meta-analysis on Determinates of MDR-TB.

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	Smok	ing	No Smo	king		Risk Ratio	Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% Cl
Assefa et al et al	42	154	187	556	9.4%	0.81 [0.61, 1.08]	
B. Baya et a	41	68	93	146	9.7%	0.95 [0.75, 1.19]	+
Baghaei.P et al	37	215	10	66	7.0%	1.14 [0.60, 2.16]	_ +
BARROSO.E et al	78	156	56	163	9.5%	1.46 [1.12, 1.89]	-
Elmi. Et al	41	143	64	171	9.2%	0.77 [0.55, 1.06]	
Fregona G, et al	32	80	62	529	9.0%	3.41 [2.39, 4.87]	
G. Mulisa et 58 al.	10	34	77	229	7.6%	0.87 [0.50, 1.52]	
Herman Owuor Weyenga et al	15	52	66	191	8.2%	0.83 [0.52, 1.33]	
Hirama T et al	16	145	30	340	7.5%	1.25 [0.70, 2.22]	- +
IRADUKUNDA.A et al	22	37	38	143	8.8%	2.24 [1.53, 3.27]	
LEMA.N et al	53	78	49	126	9.5%	1.75 [1.34, 2.28]	-
Mekonnen et al.	2	35	5	89	2.7%	1.02 [0.21, 5.00]	
Rifat M, et al	1	83	125	534	2.0%	0.05 [0.01, 0.36]	·
Total (95% CI)		1280		3283	100.0%	1.18 [0.87, 1.60]	+
Total events	390		862				ſ
Heterogeneity: Tau ² = 0.24; Chi ²	= 90.13, c	df = 12	(P < 0.000	001); I ^z =	= 87%		
Test for overall effect: Z = 1.05 (F							0.01 0.1 1 10 100 No Smoking Smoking

Figure 7: Risk ratio of Multidrug Resistant TB (MDR-TB) in relation to smoking status (smoking vs no smoking), a metaanalysis on Determinates of MDR TB.

	Diabetes M	ellites	no Diabetes M	lellites		Risk Ratio	Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% Cl	M-H, Random, 95% Cl
BARROSO.E et al	11	29	123	290	13.6%	0.89 [0.55, 1.45]	
Elmi et al.	28	88	77	226	16.8%	0.93 [0.65, 1.33]	-+-
Fregona G, et al	11	65	99	704	11.8%	1.20 [0.68, 2.13]	
IRADUKUNDA.A	4	6	56	174	11.1%	2.07 [1.13, 3.80]	
Liang.L et al	10	79	231	1916	11.3%	1.05 [0.58, 1.90]	
Rifat M, et al	34	83	216	917	18.6%	1.74 [1.31, 2.31]	+
Wijayanto et al.	10	12	35	78	16.8%	1.86 [1.30, 2.64]	
Total (95% CI)		362		4305	100.0%	1.33 [1.01, 1.77]	◆
Total events	108		837				
Heterogeneity: Tau ² =	= 0.09; Chi ² = 1	17.45, df	= 6 (P = 0.008)	; l² = 66%			
Test for overall effect	Z = 2.02 (P =	0.04)					0.01 0.1 1 10 100 no Diabetes Mellites Diabetes Mellites

Figure 8: Risk ratio of Multidrug-Resistant TB (MDR-TB) in relation to Diabetes Mellitus (Diabetes Mellitus *vs* No Diabetes Mellitus), a meta-analysis on Determinates of MDR TB.

significant association of MDR-TB with Diabetes Mellitus.

The risk of having MDR-TB in patients who have Diabetes Mellitus was 1.33 times higher than that of patients who have no Diabetes Mellitus (an overall RR of 1.33 and 95% CI, 1.01 to 1.77).

As presented in Figure 8, except for three studies [29, 38, 40], all other studies showed no association of MDR-TB to Diabetes Mellitus patients or no Diabetes Mellitus patients.

Heterogeneity testing showed significant variation among the studies ($I^2=66\%$).

Discussion

MDRTB and GENDER

The finding of this systemic review and metaanalysis showed that there is no statistically significant difference between male and female with respect to Multi Drug Resistance Tuberculosis and our study is similar with previous studies [29,30,32,33,37,38,48-50]. Our finding is not similar with previous studies [39,43,51-53].

The explanations for the association between gender and MDRTB in previous studies are socio economic factors, health seeking behavior, prolonged delays in female patients (due to lack of control at household levels on financial resources), increased immune suppression with pregnancy, lactation and increased magnitude of HIV.

MDRTB and SMOKING

This systemic review and meta-analysis did not indicate the presence of association between smoking and Multi Drug Resistance Tuberculosis and this is consistent with previous studies [54].

Further studies need to be done on smoking and resistance to anti-tuberculosis drugs [32]. This systemic review and meta-analysis are not similar with previous studies [13,38]

The reason for the difference in findings might be variation in sample size, study design and study participants characteristics

ALCOHOL and MDRTB

Our finding showed that there is no statistical significant association between MDRTB and drinking alcohol and we could not found similar findings in our search for literature but our findings is not similar with previous studies [49,55-57].

The explanation for the observed association between alcohol and MDRTB in previous studies is that alcohol drinking is associated with elevated probability of MDR-TB due to decreased adherence immunosuppression and adverse drug effects.

Drinking Alcohol was associated with increased risk of MDR-TB [37,58-59].

Diabetes Mellitus and MDRTB

In our systemic review and meta-analysis there is an association between Diabetes Mellitus and MDR TB and our finding is similar with previous studies [29,60].

The findings of this systemic review and metaanalysis are not similar with previous study [61].

The possible explanation for the variation in results might be difference in sample size and study participant's characteristics.

Educational Status and MDRTB

Fewer researchers had investigating the educational level of patients and MDR-TB [61-62].

Our systemic review and meta-analysis indicated that the risk of having MDR-TB in patients who have informal education was 0.67 times lower than that of patients who have formal education (overall RR=0.67; 95% CI, 0.46 to 0.96) and our finding is consistent with previous studies [38,39,42]. This study is not similar with previous studies [49,58,59,63-66].

The possible explanation for difference in findings between our review and other studies might be difference in study participants characteristics, sample size and study areas.

Conclusion

This systemic review and meta-analysis indicated that the risk of having MDR-TB in patients who

have informal education was 0.67 times lower than that of patients who have formal education and the risk of having MDR-TB in patients who have Diabetes Mellitus was 1.33 times higher than that of patients who have no Diabetes Mellitus.

Our review did not show the presence of association between MDR-TB and other selected variables gender, smoking and alcohol.

We recommend interested researchers on MDRTB to do further research on patient's educational level as a risk factor for MDR-TB in order to explain the variation in MDR-TB among patients with formal education and patients with informal education.

Data Availability

All data are included in the paper.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Authors' Contributions

Kaleab Tesfaye Tegegne was responsible for conceptualization, project administration, software, supervision, and development of the original drafting of the manuscript. Kaleab Tesfaye Tegegne, Eleni Tesfaye Tegegne, Abiyu Ayalew Assefa, Mekibib Kassa Tessema, Berhanu Bifato, Alelign Tadele Abebe, Muse Rike and Andualem Zenebe were participated in quality assessment of articles, methodology, validation, and screening of research papers. All authors contributed with data analysis, critically revised the paper, and agreed to be accountable for their contribution.

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Competing of interest

The authors have declared that there is no competing interest.

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