

Multi-bacillary leprosy under Chinese leprosy elimination program

Ge Li,^{1,3} Hong Zhang,^{2,3} Qingping Zhang,^{1*} Ping Chen,¹ Zhaoxing Lin,¹ Yaofei Wang,¹ Xiaodong Yang¹

¹Shaanxi Provincial Institute for Endemic Disease Control, Xi'an, 710003, China

²Shaanxi Provincial Institute for Skin Disease and STD Control, Xi'an, 710003, China

³Xi'an Jiaotong University, Xi'an, 710061, China

Ge Li and Hong Zhang contributed equally.

*Corresponding author: Qingping Zhang, E-mail: 1016009751@qq.com, Telephone: 13359182201, Fax numbers: 710003

Key words: Multibacillary; Factors; leprosy elimination

Author affiliations

¹Shaanxi Provincial Institute for Endemic Disease Control, Xi'an, 710003, China

²Shaanxi Provincial Institute for Skin Disease and STD Control, Xi'an, 710003, China

Word count : 1849

Abbreviations

MB: Multi-bacillary leprosy; LEPMIS: the leprosy management information system;

PB: pauci-bacillary leprosy; **OR:** odds ratio; **CI:** Concentration Index; **PCPs:** Primary Care Physicians

Abstract (words: 211)

Background: Objectives: To analyze the sociodemographic and clinical factors associated with multi-bacillary (MB) leprosy in elimination planning areas in Northwest China. **Design:** Retrospective observational study. **Setting:** Three specialized hospitals were included. **Participants:** The medical records of leprosy in Shaanxi Province from 2004 to 2020 were collected from the leprosy management information system (LEPMIS). **Primary and secondary outcome measures:** The basic situation of leprosy treatment and follow-up were investigated. **Results:** 305 new cases of leprosy were included in the study. 272 cases (89.18%) were MB leprosy,

29 and 33 (10.82%) were pauci-bacillary (PB) leprosy. Male patients were more likely to
30 have neurological injury ($P<0.001$; OR:0.35; 95%CI:0.194-0.630). Patients over 60
31 years old were more likely to have leprosy deformity ($P<0.001$; OR:0.113;
32 95%CI:0.027-0.478) and nerve injury ($P=0.035$; OR:0.333; 95%CI:0.115-0.965).
33 Patients with marriage histories were more likely to have leprosy deformities
34 ($P=0.018$; OR:0.842; 95%CI: 0.718-0.987). Patients with passive detection had a
35 lower probability of leprosy reaction ($P =0.011$; OR:4.268; 95%CI:1.276-14.272);
36 Patients with nerve damage, positive skin smear test or level I or level II disability
37 were more likely to be classified as MB leprosy. **Conclusions:** MB leprosy is related
38 to social and demographic factors (with or without marriage history, age at diagnosis,
39 discovery mode) and clinical factors (such as the number of skin lesions and nerve
40 lesions).

41 **Key words:** Multibacillary leprosy; Factors; Retrospective observational study

42 **Summary box:**

43 **What is already known about this subject:**

- 44 1. Leprosy is widespread throughout the world and seriously endangers human health
- 45 2. The condition of patients with MB leprosy is more complex, the treatment course
46 is longer and the prognosis is poorer, which seriously affect the life quality of the
47 patients
- 48 3. The incidence of polybacterial leprosy was different in different conditions.

49 **What are the new findings:**

- 50 1. Patients over 60 years old were more likely to have leprosy deformity and nerve

51 injury.

52 2. Patients with marriage histories were more likely to have leprosy deformities.

53 3. Patients with passive detection had a lower probability of leprosy reaction.

54 4. Patients with nerve damage, positive skin smear test or level I or level II disability
55 were more likely to be classified as MB leprosy.

56 **How might it impact on clinical practice in the foreseeable future?**

57 1. This study is the first time to analyze the differences in the classification of leprosy
58 patients in western China and the related influencing factors, aiming to provide
59 theoretical support for proposing better prevention and treatment measures.

60 2. 305 cases of leprosy patients in western China for 17 years were included, with
61 reliability and validity guaranteed. Medical records are clear and complete, and the
62 research value is high.

63 3. We can improve the ability of clinical diagnosis and treatment of MB leprosy by
64 intervening the related risk factors, and provide valuable demographic data for the
65 clinical study of leprosy.

66 **Introduction**

67 Leprosy, caused by mycobacterium tuberculosis, is a chronic infectious disease¹ and
68 mainly induces injuries of skin and peripheral nerves.² If not treated in time, leprosy
69 can cause permanent damage to peripheral nerves and may lead to amputation and
70 deformity.³ The diagnosis of leprosy is based on a series of clinical manifestations.

71 According to different therapeutic purposes, cases are divided into two types:

72 Paucibacillary (PB) leprosy and Multibacillary (MB) leprosy. MB leprosy is an

73 aggressive progression of the disease, which is mainly caused by unresponsiveness of
74 cellular immunity of leprosy bacilli,⁴ and is characterized by high infectivity and
75 functional disability rate.⁵ Leprosy disability severely affects the quality of life of
76 leprosy patients, and the social discrimination caused by it may induce psychological
77 problems. The World Health Organization (WHO) aimed to achieve the goal of global
78 eradication of leprosy.⁶ However, despite effective prevention and control measures
79 were extensively implemented, the number of new cases worldwide has remained
80 almost unchanged in the past ten years, with about 250,000 new diagnosed cases each
81 year.⁷ The way to the destination, is not easy and smooth, and the potential deformity⁸
82 caused by leprosy aggravates related social, health and economic loads.⁹ Although
83 leprosy is generally in a low epidemic state in northwest China,¹⁰ but the proportion
84 of MB types¹¹ and the rate of disability¹² are still at a high level.¹³

85 **Methods**

86 1. Criteria of diagnosis

87 The diagnoses of new and recurrent leprosy were based on Leprosy Prevention
88 Manual for Primary Care Physicians (PCPs) and Leprosy Diagnosis Standard
89 WS291-2008. Cases were classified according to the Leprosy Classification Standard,
90 WHO, 1987. Disability classification was bases on the Disability Classification
91 Standard for Leprosy, WHO, 1988.

92 2 Data sources

93 All new and recurrent leprosy cases were collected from the Leprosy Prevention and
94 Control Management Information System in China (LEPMIS). Population data came

95 from the Shaanxi Statistical Yearbook.

96 3 This study was an observational and retrospective study, involving 368 cases
97 collected in the LEMPIS from 2004 to 2015. The variables included gender,
98 nationality, occupation, education level, marital status, residence history, age, method
99 of detection, skin lesions, bacteria detection, nerve damage, and disability
100 classification. Cases with incomplete information (n=63) were excluded. This
101 retrospective cross-sectional study was reviewed and approved by the institutional
102 ethics committee.

103 4 Statistical analysis

104 EXCEL 2010 were used to establish a new and recurrent leprosy case report database.
105 Appropriate statistical methods including *U* test, χ^2 test, Fisher exact test, and
106 multivariate logistic regression were used to carry out statistical analysis in SPSS 19.0
107 software. Quantitative variables were expressed in descriptive statistics, and
108 qualitative variables were expressed in rates and 95% confidence intervals (CI). For
109 statistical purposes, we have divided the analysis variables into different categories.
110 We conducted Kolmogorov-Smirnov test and Levene test respectively to evaluate the
111 normality and variance homogeneity of quantitative variables. χ^2 test was used to
112 analyze the influence of various factors, and the results were expressed by dominance
113 ratio (socio-demographic factors and clinical factors). The significance level was
114 $P < 0.05$. Logistic regression analysis was used to adjust the confounding variables to
115 determine the independent risk factors of leprosy MB. In order to determine which
116 independent risks are related to the operational classification of leprosy, we used the

117 forward stepwise logistic regression method. This method used a model designed for
118 two-class qualitative variables to analyze the significance of each independent
119 variable, and a logistic regression model was obtained to classify the subjects into PB
120 type and MB type. The percentage of correct identification was 89.2%. The results
121 showed that the correct rate of this method is higher than the random correct rate
122 (50%). The global test of the model has statistical significance ($P < 0.001$). The
123 Hosmer-Lemeshow goodness-of-fit test showed that there is no statistical significance
124 ($P = 0.713$) between the expected frequency obtained from the prediction probability
125 and the observed frequency, which means the model fitted well.

126 **Results**

127 **1. Basic situation of leprosy cases**

128 It was shown in the system that there were 368 leprosy cases in Shaanxi province
129 between 2004 and 2020, of which 69.5% were male cases. 98.7% patients were Han
130 ethnics. Most of the new leprosy cases were farmers (91.1%). 6 cases (2.0%) received
131 education more than 9 years, 198 cases (64.9%) received education between 1 year
132 and 9 years, 101 cases (33.3%) received education less than 1 year. 216 cases (70.8%)
133 had marriage history, and 97.4% of the cases resided permanently in the locality. The
134 average age of diagnosed with leprosy was 45.79 ± 13.6 years. 16.1% cases were
135 diagnosed older than 60 years. 32.8% cases were infected from within the family,
136 12.5% cases were infected from outside of the family, and 54.8% cases had no
137 definite sources of infection. The detected cases were mostly MB (272 cases, 89.2%).
138 Compared with female patients, males were more prone to nerve injury ($P < 0.001$;

139 OR:0.35; 95%CI:0.194-0.630). Patients diagnosed older than 60 years were relatively
140 prone to disability (P<0.001; OR:0.113; 95%CI:0.027-0.478) and nerve injury
141 (P=0.035; OR:0.333; 95%CI:0.115-0.965). Patients with marriage histories were more
142 prone to leprosy disability (P=0.018; OR:0.842; 95%CI:0.718-0.987). The incidence
143 of leprosy reaction was lower in the patients detected passively than those detected
144 actively (P=0.011; OR:4.268; 95%CI:1.276-14.272).

145 **2. MB leprosy and socio-demographic factors**

146 The χ^2 test results showed that there were statistically significant differences in sex,
147 education level, marital status, residence history, age at diagnosis, mode of diagnosis,
148 and leprosy classification of patients (P<0.05), as shown in Table 1. Univariate
149 analysis showed that there were statistically significant relationships between sex
150 (p=0.043), education level (p=0.006), marital status (p<0.001), residence history
151 (p=0.014), age at diagnosis (p=0.019), mode of diagnosis (p=0.001) and MB leprosy.
152 The adjusted logistic regression analysis showed that marriage history (P=0.006;
153 OR=3.592; 95%CI:1.444-8.934), the age at diagnosis older than or equal to 60 years
154 (P<0.05; OR=1.034; 95%CI=1.001-1.068), and active detection (P=0.002; OR=3.640;
155 95%CI=1.590-8.333) were significantly correlated with multi-bacterial leprosy.

156

Table 1 Relationship between sociodemographic factors and leprosy types

| | MB leprosy N=272 | PB leprosy N=33 | P value | OR | 95%CI |
|---------------------------|-----------------------------|----------------------------|----------------|-----------|----------------|
| Sex | | | | | |
| Male | 184 (67.65) | 28 (84.85) | 0.043 | 0.373 | (0.139-1.000) |
| Female | 88 (32.35) | 5 (15.15) | | | |
| Ethnics | | | | | |
| Han | 268 (98.53) | 33 (100.00) | 0.483 | 0.89 | (0.856-0.926) |
| Others | 4 (1.47) | 0 (0.00) | | | |
| Profession | | | | | |
| Farmer | 24 (8.82) | 3 (9.09) | 0.959 | 0.968 | (0.275-3.407) |
| Others | 248 (91.18) | 30 (90.91) | | | |
| Education level, y | | | | | |
| More than 9 | 6 (2.21) | 0 (0.00) | | | |
| Between 1 and 9 | 184 (67.65) | 14 (42.42) | 0.002 | 3.145 | (1.504-6.573) |
| Less than 1 | 82 (30.15) | 19 (57.58) | | | |
| Marital status | | | | | |
| Unmarried | 69 (25.37) | 20 (60.61) | <0.001 | 0.221 | (0.104-0.468) |
| Married | 203 (74.63) | 13 (39.39) | | | |
| Living history | | | | | |
| Local | 267 (98.16) | 30 (90.91) | 0.014 | 5.34 | (1.215-23.465) |
| Nonlocal | 5 (1.84) | 3 (9.09) | | | |
| Age, y | | | | | |
| Older than 60 | 40 (14.71) | 9 (27.27) | | | |
| Between 20 and 60 | 224 (82.35) | 23 (69.70) | 0.063 | 0.460 | (0.199-1.061) |
| Younger than 20 | 8 (2.94) | 1 (3.03) | | | |
| Mode of diagnosis | | | | | |
| Active detection | 53 (19.49) | 15 (45.45) | 0.001 | 0.290 | (0.137-0.614) |
| Passive detection | 219 (80.51) | 18 (54.55) | | | |

157

158

159 **3. Analysis of MB leprosy and its clinical factors**

160 Of the cases of MB leprosy (89.2%), 21.32% cases had less than 5 skin lesions,
161 85.66% cases had no obvious leprosy reaction, 34.93% cases had no positive bacteria
162 detection, 19.49% had no nerve damage, and 22.43% cases had no deformity, as show
163 in Table 2. Analysis showed that patients with less than 1 skin lesion were less likely
164 to be classified as MB leprosy than those with more than 5 lesions. Patients with
165 nerve damage, positive skin smear results, or grade I or II disability were more likely
166 to be classified as MB leprosy. The adjusted logistic regression analysis showed that
167 clinical variables related to MB leprosy included >5 lesions (P<0.01; OR=7.880;
168 95%CI=2.506-24.775), and ≥ 2 nerve lesions (P<0.05; OR=3.516;

169 95%CI=1.076-11.491).

Table 2 Relationship between clinical factors and types of leprosy

| | MB leprosy (n=272) | PB leprosy (n=33) | P value | OR | 95% CI |
|--------------------------|-----------------------|----------------------|---------|--------|----------------|
| Skin lesion | | | <0.001 | 0.170 | (0.052-0.554) |
| None | 8 (2.94) | 5 (15.15) | | | |
| 1 lesion | 6 (2.21) | 7 (21.21) | | | |
| 2-5 lesions | 44 (16.18) | 15 (45.45) | | | |
| >5 lesions | 214 (78.68) | 6 (18.18) | | | |
| Leprosy reaction | | | 0.3 | 0.597 | (0.174-2.053) |
| No reaction | 233 (85.66) | 30 (90.91) | | | |
| I reaction | 15 (5.51) | 3 (9.09) | | | |
| II reaction | 20 (7.35) | 0 (0.00) | | | |
| Mixed reaction | 4 (1.47) | 0 (0.00) | | | |
| Skin smear result | | | <0.001 | 13.508 | (4.612-39.566) |
| Positive | 177 (65.07) | 4 (12.12) | | | |
| Negative | 95 (34.93) | 29 (87.88) | | | |
| Nerve damage | | | 0.003 | 1.355 | (0.500-3.678) |
| None | 53 (19.49) | 5 (15.15) | | | |
| 1 nerve | 31 (11.40) | 11 (33.33) | | | |
| ≥2 nerves | 188 (69.12) | 17 (51.52) | | | |
| Disability | | | 0.007 | 0.578 | (0.266-1.259) |
| None | 61 (22.43) | 11 (33.33) | | | |
| Grade I | 93 (34.19) | 3 (9.09) | | | |
| Grade II | 103 (37.87) | 19 (57.58) | | | |
| Not clear | 15 (5.51) | 0 (0.00) | | | |

170

171

172 Discussion

173 These results of our study showed high endemic characteristics of leprosy in this
174 investigated area, suggesting that the delay on leprosy diagnosis still existed in
175 northwest China,¹⁴ which led to more serious consequences and disabilities of
176 leprosy.¹⁵

177 Female leprosy patients were prone to have poor prognosis. MB leprosy had greater
178 impact on males than PB leprosy, even after adjusting the model. This result was
179 consistent with previous results conducted in other parts of our country.¹⁶ Leprosy
180 showed a low prevalence trend at this stage, and male patients still accounted for a

181 relatively high proportion of the new diagnosed cases in each year, which may be
182 related to different genetic susceptibility in different genders.¹⁷ Meanwhile, females
183 got more skin consultations than males, and males may be more easily exposed to
184 leprosy bacilli related to behavioral and cultural factors,¹⁸ which may partly explain
185 the dominant position of male cases. In addition, there may be some correlation
186 between androgen and MB leprosy, especially in adolescent males,¹⁹ and the immune
187 response stimulated by a large amount of androgen secreted had poor disease control
188 effect.²⁰ The susceptibility of the elderly to MB leprosy may be related to the
189 prolonged incubation period of leprosy bacilli, resulting to delayed response.²¹ In
190 addition, the aging of immune system in the elderly²² was an aggravating factor for
191 infection control.²³ A previous related study showed that the higher incidence of
192 leprosy changed from PB to MB, from the young to the elderly (especially people
193 over 60 years old), and from women to men.²⁴ Even after the adjustment of the model,
194 less years of education were related to MB leprosy. People with higher education are
195 more inclined to seek medical services to avoid delaying diagnosis and treatment.²⁵
196 Data showed that people with marriage history are the advantage group of MB leprosy,
197 which may indicate that close contact in the home was related to exposure to leprosy
198 bacilli, but we cannot rule out the importance of social contact in disease
199 transmission.²⁶ A related study abroad concluded that the spread of leprosy is not
200 limited to the indoor environment, and outdoor infection also affects the spread of the
201 disease.²⁷ Passive detection was a protective factor for MB leprosy, so it is necessary
202 to increase the publicity and education of leprosy prevention and control knowledge

203 in low-prevalence areas, and to improve the awareness rate of the masses and
204 self-care awareness.^{28, 29} More than 5 lesions were associated with MB leprosy, which
205 indicated that high concentration of leprosy bacilli infection can lead to more tissue
206 destruction, more skin damage and worse deformation. Related studies abroad have
207 analyzed the innate immune factors related to skin lesions of leprosy patients. For
208 example, in cases of MB infection, the activation of complement protein will
209 aggravate the inflammatory process and lead to peripheral nerve injury, resulting in
210 significantly higher incidence of skin lesions in MB patients than in PB patients.³⁰

211 Disability degree is an indicator of the ability to diagnose and monitor leprosy of
212 health service departments, and patients classified as Class I or Class II disability
213 were related to delayed diagnosis or monitoring failure.³¹

214 It was shown that MB cases have a higher probability of grade I or II disability after
215 model adjustment. By adding Class I disability (96 cases, 31.48%) to Class II
216 disability (122 cases, 40.0%), the number of patients with disabilities increased to 218
217 (71.48%), which represented a high percentage of the overall disability. In addition,
218 the incidence rate of Class II disability in our sample is far higher than the global
219 average of 6% reported by the World Health Organization in 2016,³² which indicated
220 that there is a delay in diagnosis or misdiagnosis in these patients.

221 This study was based on second-hand data obtained from the LEPMIS, so it has some
222 limitations, such as inconsistent information, prevalence bias and the defect of
223 cross-sectional design. Future studies need to consider longitudinal studies or
224 geographical distribution to clarify factors related to leprosy. MB patients were the

225 main infectious source of leprosy, so early detection and treatment can effectively
226 block the transmission of leprosy in the infectious source control link, which is of
227 great significance to reduce the probability of leprosy patients with disability. This
228 study elaborated the epidemic characteristics and regional characteristics of MB
229 leprosy in northwest China, which may be helpful to effectively prevent and control
230 leprosy.

231 **Statements**

232 **Acknowledgements**

233 We would like to thank all participants in the study on leprosy. In particular we would
234 like to express our thanks to National Health Commission of the People's Republic of
235 China and Health Commission of Shaanxi Province.

236 **Contributorship statement**

237 Ge Li and Hong Zhang co-wrote the article. Ping Chen has contributed to the
238 collection and acquisition of the article data. Yaofei Wang gave guidance on research
239 methods and case review. Xiaodong Yang was involved in data collection and
240 statistical analysis. Qingping Zhang, the corresponding author, reviewed the data and
241 results of this article.

242 Ge Li and Hong Zhang contributed equally.

243 *Corresponding author: Qingping Zhang, E-mail: 1016009751@qq.com

244 **Funding**

245 There is no funding to report for this submission. The authors have not declared a
246 specific grant for this research from any funding agency in the public, commercial or
247 not-for-profit sectors.

248 **Competing interests**

249 No competing interests exist in this study, all authors have approved the manuscript
250 for submission on your journal, the content of the manuscript has not been published
251 or submitted elsewhere, and the retrospective study was reviewed and approved by
252 the institutional ethics committee. All authors declare no known conflict of interest.

253 **Data sharing statement**

254 Data are available in a public, open access repository.

255 **Patient and public involvement statement**

256 Our research did not involve patients. LEPMIS is a legal and open information
257 platform for leprosy prevention and research personnel in China with a view to
258 strengthen Global Health and Global Health Research. The case information does not
259 involve the patient's personal information, but is collective and provides the basis for
260 the country's strategic prevention and control of leprosy No patient involved under the
261 sub-heading Patient and public involvement.

262

263 **Patient consent for publication** Not required.

264

265 **Ethics approval** Not required. Our research did not involve patients. LEPMIS is a
266 legal and open information platform for leprosy prevention and research personnel in
267 China with a view to strengthen Global Health and Global Health Research. The case
268 information does not involve the patient's personal information, but is collective and
269 provides the basis for the country's strategic prevention and control of leprosy No
270 patient involved under the sub-heading Patient and public involvement.

271

272 **Provenance and peer review** Not commissioned; externally peer reviewed.

273

274 **Data availability statement** The authors confirm that the data supporting the findings
275 of this article are available within the article.

276

277 **Open access** This is an open access article distributed in accordance with the Creative
278 Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits
279 others to distribute, remix, adapt, build upon this work non-commercially, and license
280 their derivative works on different terms, provided the original work is properly cited,
281 appropriate credit is given, any changes made indicated, and the use is
282 non-commercial.

283

284

285 **References**

286 1 Limeira OM, Gomes CM., Morais OO, *et al.* Active search for leprosy cases in Midwestern
287 Brazil: a serological evaluation of asymptomatic household contacts before and after prophylaxis
288 with bacillus Calmette-Guerin. *Rev Inst Med Trop Sao Paulo* 2013;55:173-177.

289 2 Queiros MI, Ramos AJ, Alencar CH, *et al.* Clinical and epidemiological profile of leprosy
290 patients attended at Ceara, 2007-2011. *An Bras Dermatol* 2016;91:311-317.

291 3 Camuset G, Lafarge S, Borgherini G, *et al.* Leprosy on Reunion Island, 2005-2013: Situation
292 and Perspectives. *PLoS Negl Trop Dis* 2016;10:e4612.

293 4 Moschioni C, Antunes C M, Grossi M A, *et al.* Risk factors for physical disability at diagnosis
294 of 19,283 new cases of leprosy. *Rev Soc Bras Med Trop* 2010;43:19-22.

295 5 Sales A M, Ponce D L A, Duppre N C, *et al.* Leprosy among patient contacts: a multilevel study
296 of risk factors. *PLoS Negl Trop Dis* 2011;5:e1013.

297 6 Sehgal V N, Sardana K, Dogra S. The imperatives of leprosy treatment in the pre- and
298 post-global leprosy elimination era: appraisal of changing the scenario to current status. J
299 Dermatolog Treat 2008;19:82-91.

300 7 Rodrigues L C, Lockwood D N. Leprosy now: epidemiology, progress, challenges, and research
301 gaps. The Lancet Infectious Diseases 2011;11:464-470.

302 8 Martins R J, Carloni M E, Moimaz S A, *et al.* Sociodemographic and epidemiological profile of
303 leprosy patients in an endemic region in Brazil. Rev Soc Bras Med Trop 2016;49:777-780.

304 9 Global leprosy update, 2016: accelerating reduction of disease burden. Wkly Epidemiol Rec
305 2017;92:501-519.

306 10 Siyu Long, Meiwen Yu, Liangbin Yan, *et al.* Epidemiological characteristics of leprosy in
307 China in 2011 and 2015. Chin J Derm 2017;50:400-403.

308 11 Global leprosy update, 2014: need for early case detection. Wkly Epidemiol Rec
309 2015;90:461-474.

310 12 Global leprosy update, 2013; reducing disease burden. Wkly Epidemiol Rec 2014;89:389-400.

311 13 Global leprosy update, 2016: accelerating reduction of disease burden. Wkly Epidemiol Rec
312 2017;92:501-519.

313 14 Yaofei Wang, Ningxia An, Huimin Wang, *et al.* Discussion on the causes of delayed diagnosis
314 of leprosy in Chenggu County, Shaanxi Province. Chin J Derm Venereol 2008;01:42-43.

315 15 Monteiro L D, Martins-Melo F R, Brito A L, *et al.* Physical disabilities at diagnosis of leprosy
316 in a hyperendemic area of Brazil: trends and associated factors. Lepr Rev 2015;86:240-250.

317 16 Yuanguai Li, Tiejue Wu. Analysis of 9 new cases of leprosy in Zunyi in 2013. Chin J Derm
318 Venereol 2016;32:496.

319 17 Shuxia Chang, Xiaohua Wang, Daocheng Zheng. Research Progress of leprosy susceptibility
320 Genes. J Diagn Ther Dermato-Venereol 2018;25:253-256.

321 18 Silva M E, de Souza C D, Costa E S S, *et al.* Epidemiological aspects of leprosy in
322 Juazeiro-BA, from 2002 to 2012. An Bras Dermatol 2015;90:799-805.

323 19 Guerra-Silveira F, Abad-Franch F. Sex bias in infectious disease epidemiology: patterns and
324 processes. PLoS One 2013;8:e62390.

325 20 Modlin R L. The innate immune response in leprosy. Curr Opin Immunol 2010;22:48-54.

326 21 Nobre M L, Illarramendi X, Dupnik K M, *et al.* Multibacillary leprosy by population groups in
327 Brazil: Lessons from an observational study. PLoS Negl Trop Dis 2017;11:e5364.

328 22 Chou J P, Effros R B. T cell replicative senescence in human aging. Curr Pharm Des
329 2013;19:1680-1698.

330 23 Hepper H J, Sieber C, Walger P, *et al.* Infections in the elderly. Crit Care Clin
331 2013;29:757-774.

332 24 Nobre M L, Illarramendi X, Dupnik K M, *et al.* Multibacillary leprosy by population groups in
333 Brazil: Lessons from an observational study. PLoS Negl Trop Dis 2017;11:e5364.

334 25 Teasdale K, De Wildt G, Das P K, *et al.* The patient perspective of the diagnostic process for
335 leprosy in Brazil. An exploratory study. Lepr Rev 2015;86:21-36.

336 26 Klusener S, Dribe M, Scalone F. Spatial and Social Distance at the Onset of the Fertility
337 Transition: Sweden, 1880-1900. Demography 2019;56:169-199.

338 27 Santos S D, Penna G O, Costa M C, *et al.* Leprosy in children and adolescents under 15 years
339 old in an urban centre in Brazil. Mem Inst Oswaldo Cruz 2016;111:359-364.

340 28 Fayang Kang, Lihua Ruan, Yongchao Jia, *et al.* Analysis of risk factors of abnormal deformity

341 in newly discovered leprosy patients in Guangyuan City from 1999 to 2008. *J Prev Med Inf*
342 2010;26:610-612.

343 29 Yongkui Hu, Zhiguang Zhou. Comparison of methods of leprosy detection. *Occup*
344 *Health*;2007;20:1837-1838.

345 30 Bahia E I N, Iyer A M, Ramaglia V, *et al.* In Situ complement activation and T-cell immunity in
346 leprosy spectrum: An immunohistological study on leprosy lesional skin. *PLoS One*
347 2017;12:e177815.

348 31 Global leprosy update, 2016: accelerating reduction of disease burden. *Wkly Epidemiol Rec*
349 2017;92:501-519.

350 32 Assis B P N, Lyon S, Grossi M A D F, *et al.* Risk factors for physical disability upon release
351 from multidrug therapy in new cases of leprosy at a referral center in Brazil. *Revista do Instituto*
352 *de Medicina Tropical de São Paulo* 2019,61:e13..