Multi-bacillary leprosy under the Chinese leprosy elimination program*

Dear Editor,

Leprosy, caused by Mycobacterium leprae, is a chronic infectious disease, which may cause permanent damage to peripheral nerves and deformity. According to the World Health Organization (WHO), leprosy cases are classified in PB (paucibacillary) or MB (multibacillary) leprosy based on the number of skin lesions: PB leprosy (2–5 skin lesions), and MB leprosy (more than 5 skin lesions). MB leprosy is mainly caused by the unresponsiveness of cellular immunity against leprosy bacilli, and is characterized by high infectivity and functional disability rate. Leprosy disability severely affects the life quality of leprosy patients and may induce psychological problems. However, despite effective measures were extensively implemented, the number of new cases worldwide has remained at almost 250,000 each year. Although leprosy is generally in a low endemic state in northwest China, the proportion of MB cases and the rate of disability are still at a high level.

To analyze the sociodemographic and clinical factors associated with MB leprosy in elimination planning areas in Northwest China, three specialized hospitals were included. The medical records of leprosy in province in northwest China from 2004 to 2020 were collected from the Leprosy Management Information System (LEPMIS).

This is an observational and retrospective study, involving 305 cases of MB and PB leprosy cases, collected in the LEMPS from 2004 to 2020. The variables included gender, nationality, occupation, and others. The statistical significance level was $p < 0.05$. Logistic regression analysis was used to adjust the confounding variables to determine the independent risk factors of MB leprosy cases.

Significant differences in sex and some other variables ($p < 0.05$) were shown comparing MB and PB cases, as shown in Table 1. Of the MB leprosy cases, 21.32% cases were documented as having less than 5 skin lesions, as shown in Table 2.

These results showed high endemic characteristics of leprosy, suggesting that the delay on leprosy diagnosis still exists in northwest China, which led to more serious consequences and disabilities. Leprosy showed a low prevalence trend at this stage, and male patients still accounted for a relatively high proportion of the newly diagnosed cases in each year, which may be related to different genetic susceptibility in different genders. Meanwhile, females got more skin consultations than males, and males may be more easily exposed to leprosy bacilli related to behavioral and cultural factors, which may partly explain the dominant position of male cases. The susceptibility of the elderly to MB leprosy may be related to the prolonged incubation period of leprosy bacilli, resulting to delayed response. In addition, the aging of the immune system in the elderly was an aggravating factor for infection control. People with higher education are more inclined to seek medical services to avoid delaying diagnosis and treatment. Data showed that people with a marriage history are the advantaged group of MB leprosy, which may indicate that close contact in the home was related to exposure to leprosy bacilli, but the we cannot rule out the importance of social contact in disease transmission.

Passive detection was a protective factor for MB leprosy, so it is necessary to increase the publicity and education of leprosy prevention and control knowledge in low-prevalence areas, and to improve the awareness rate of the masses and self-care awareness. More than 5 lesions were associated with MB leprosy, which indicated that a high concentration of leprosy bacilli infection can lead to more tissue destruction, more skin damage, and worse deformation. MB cases have a higher probability of grade I or II disability after model adjustment. The incidence rate of Class II disability in the present sample is far higher than the global average of 6% reported by the WHO in 2016, which indicated that there is a delay in diagnosis or misdiagnosis in these patients.

This study was based on second-hand data obtained from the LEPMIS, so it has some limitations, such as inconsistent information, prevalence bias and the defect of cross-sectional design. Future longitudinal studies or geographical distribution are desirable to clarify factors related to leprosy.

In conclusion, MB patients were the main infectious source of leprosy, so early detection and treatment can effectively block the transmission of leprosy in the infectious source control link, which is of great significance to reduce the probability of leprosy in patients with disability. This study has shown the epidemic characteristics and regional characteristics of MB leprosy in northwest China, which may be helpful in effectively preventing and controlling leprosy.

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Authors’ contributions

Ge Li: Co-wrote the article; the corresponding author, reviewed the data and results of this article; contributed to interpretation of the data, commented on the manuscript, revised the manuscript, and approved the final version for publication.

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### Table 1  Relationship between sociodemographic and behavioral factors and leprosy types.

| Factor                  | MB leprosy (n = 272) | PB leprosy (n = 33) | p-value | OR     | 95% CI      |
|-------------------------|-----------------------|---------------------|---------|--------|-------------|
| **Sociodemographic factors** |                       |                     |         |        |             |
| **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)            | 0.002   | 3.145  | (0.137–6.573) |
| Between 1 and 9         | 184 (67.65)           | 14 (42.42)          | 0.959   | 0.968  | (0.275–3.407) |
| 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)           | 0.063   | 0.460  | (0.199–1.061) |
| Between 20 and 60       | 224 (82.35)           | 23 (69.70)          |         |        |             |
| Younger than 20         | 8 (2.94)              | 1 (3.03)            |         |        |             |
| **Behavioral factors**  |                       |                     |         |        |             |
| **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)          |         |        |             |

### Table 2  Relationship between clinical factors and leprosy types.

| Factor                  | MB leprosy (n = 272) | PB leprosy (n = 33) | p-value | OR     | 95% CI      |
|-------------------------|-----------------------|---------------------|---------|--------|-------------|
| **Skin lesion**         |                       |                     |         |        |             |
| None                    | 8 (2.94)              | 5 (15.15)           | < 0.001 | 0.170  | (0.052–0.554) |
| 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**    |                       |                     |         |        |             |
| No reaction             | 233 (85.66)           | 30 (90.91)          | 0.3     | 0.597  | (0.174–2.053) |
| 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**   |                       |                     |         |        |             |
| Positive                | 177 (65.07)           | 4 (12.12)           | < 0.001 | 13.508 | (4.612–39.566) |
| Negative                | 95 (34.93)            | 29 (87.88)          |         |        |             |
| **Nerve damage**        |                       |                     |         |        |             |
| None                    | 53 (19.49)            | 5 (15.15)           | 0.003   | 1.355  | (0.500–3.678) |
| 1 nerve                 | 31 (11.40)            | 11 (33.33)          |         |        |             |
| ≥ 2 nerves              | 188 (69.12)           | 17 (51.52)          |         |        |             |
| **Disability**          |                       |                     |         |        |             |
| None                    | 61 (22.43)            | 11 (33.33)          | 0.007   | 0.578  | (0.266–1.259) |
| Grade I                 | 93 (34.19)            | 3 (9.09)            |         |        |             |
| Grade II                | 103 (37.87)           | 19 (57.58)          |         |        |             |
| Not clear               | 15 (5.51)             | 0 (0.00)            |         |        |             |
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Conflicts of interest

None declared.

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Negative patch tests: what should we think about these results? a

Dear Editor,

Patch tests are the best tools to identify the etiological agents of allergic contact dermatitis (ACD). They are indicated in suspected ACD, in chronic eczemas with no defined etiology, in occupational contact dermatitis, and for the investigation of drug reactions with a delayed hypersensitivity mechanism. Responses to patch tests are evaluated using morphological criteria already described by the International Contact Dermatitis Research Group (ICDRG). A negative result in a patient with some type of eczema can be frustrating; thus, it is important to know the differential clinical diagnoses and the reasons why a test can be negative.

The present study aims to determine the frequency of negative patch tests in patients with clinical suspicion of ACD, their epidemiological profile and final diagnoses. This study was approved by the Human Research Ethics Committee under number 20285919.1.0000.5479.

This is a descriptive retrospective study, carried out with the analysis of medical records of patients with ACD from 2013 to 2018 who had negative results or irritative reactions to the tested substances. The patch test series used were selected according to the suspected diagnosis: Brazilian Standard, Cosmetics and Corticosteroids (FDA Allergenic/Rio de Janeiro, Brazil); Latin American, Expanded Series Patch Testing, Phototest, Footwear, Metals, Ulcers (Chemotechnique Diagnostics/Vellinge, Sweden); Hair, Nails, and Anti-Inflammatoryes (IPI-ASAC/São Paulo, Brazil). The tests were applied to the patients upper back region using AlergoChamber™ (Neoflex/Sertãozinho, São Paulo, Brazil). The obtained data were tabulated and analysed.

Of the 694 patients submitted to patch tests with a diagnostic hypothesis of ACD, 116 (16.7%) were negative for all tested substances, 72 of which (62.1%) were female. Age

a Study conducted at the Dermatology Clinic, Santa Casa de São Paulo, São Paulo, SP, Brazil.