

CORRESPONDENCE ▼

Unanswered questions on the safety of MDT-U - Reply*

Gerson Oliveira Penna^{1,2}, Heitor de Sá Gonçalves³,
Maria Araci de Andrade Pontes³

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Dear Dr. Barve,

Thank you very much for your comments regarding our paper "Clinical trial for uniform multidrug therapy for leprosy patients in Brazil (U-MDT/CT-BR): adverse effects approach".¹

Let us clarify some points:

1. Indeed, both pigmentation and xerosis are caused by clofazimine. We definitely did not imply that these were due to rifampicin and/or dapsone.
2. It is clear that paucibacillary (PB) patients treated with R-MDT do not use clofazimine. However, as mentioned in reference 2, the inclusion of clofazimine in the treatment of PB patients did not lead to an increase in non-compliance when we used U-MDT.²
3. Definitely, we cannot compare data from leprosy control programs with a randomized and controlled clinical trial.³ It would be a fundamental and serious mistake. However, it is very important to stress that only 24 patients had to interrupt treatment due to adverse effects (AE).¹
4. Despite your question about the moment of AE onset, for us it is clear and elementary that the shorter the treatment is, the less AE we are likely to find.

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¹ Center for Tropical Medicine, Universidade de Brasília, Brasília, DF, Brazil.

² Escola Fiocruz de Governo, Fundação Oswaldo Cruz, Brasília, DF, Brazil.

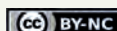
³ Dermatology Outpatient Clinic, Centro de Dermatologia Dona Libânia, Fortaleza, CE, Brazil.

MAILING ADDRESS:

Gerson Oliveira Penna

E-mail: gerson.penna@fiocruz.br


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AUTHORS' CONTRIBUTIONS

Gerson Oliveira Penna  ORCID 0000-0001-8967-536X
Elaboration and writing of the manuscript; approval of the final version of the manuscript.

Heitor de Sá Gonçalves  ORCID 0000-0001-9062-2180
Elaboration and writing of the manuscript; approval of the final version of the manuscript.

Maria Araci de Andrade Pontes  ORCID 0000-0002-7292-8452
Elaboration and writing of the manuscript; approval of the final version of the manuscript.

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