

McLeod Syndrome: Report of an Indian family with phenotypic heterogeneity

Sir,

We thank the authors^[1] of the letter for their keen interest in our published report on McLeod Syndrome (MLS)^[2]. It appears that the report mentioned by the authors of the letter had not been adequately cited in PUBMED. Moreover, a lack of molecular genetic data on the case would fall short of making a definitive diagnosis of MLS in the case mentioned. The family reported by us, therefore, appears to be the first genetically proved case of MLS reported from India.

A. Chakravarty, P. Bhattacharya¹, D. Banerjee, S. Mukherjee¹

Departments of Neurology and Hematology,
Vivekananda Institute of Medical Sciences,
¹Departments of Transfusion Medicine and
Hematology, AMRI Hospitals
Kolkata, India

For correspondence:

Dr. Ambar Chakravarty, 1E 1202 AVISHIKTALI,
Kolkata - 700 078, India. E-mail: saschakra@yahoo.com

1. Mehndiratta M. McLeod syndrome and Acanthocytosis. Ann Indian Acad Neurol 2011. Acad Neurol 2011;14:143
2. Chakravarty A, Bhattacharya P, Banerjee D, Mukherjee S. McLeod Syndrome: Report of an Indian Family with Phenotypic Heterogeneity. Ann Indian Acad Neurol 2011;14:53-5.

Access this article online

Quick Response Code:



Website:

www.annalsofian.org

DOI:

10.4103/0972-2327.82827