Reviewer's report

Title: Two novel connexin32 mutations cause early onset X-linked Charcot-Marie-Tooth

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Reviewer: Moritz Meins

Reviewer’s report:

I find the paper by Braathen et al. is a well-written report and expanding the knowledge of genotype-phenotype correlations for CX32 mutations. It should be considered for publication by BMC neurology.

There are a few points that should be addressed by the authors:

1. The numbering of nucleotides should refer to the open reading frame, following the current recommendations for mutation nomenclature.
2. I can not recognize from the manuscript which family members were available for molecular genetic analysis, who carried the mutation and who did not.
3. Furthermore, did the authors check for the absence of the mutation in a control cohort?
4. The figure legends are very short.