Reviewer’s report

Title: SCL20A2 mutation presenting with acute ischemic stroke: a case report

Version: 0 Date: 19 Nov 2017

Reviewer: Fatma Silan

Reviewer’s report:

This manuscript presents acute ischemic stroke in a Primary Familial Brain Calcification patient with SLC20A2 gene missense mutation. This is a very rare condition and genotype phenotype correlations are very helpful for medical professionals.

Authors didn't give any information about patient's parents: Are they living? with or without any neurological problem? If they are dead, what is the cause of death?

Is there any other family member (uncle? aunt? grandmother/father et cet) with neurological problems, and/or stroke?

One of his daughters has brain calcification without any clinical symptom. How old is she?

Are the methods appropriate and well described?
If not, please specify what is required in your comments to the authors.

Yes

Does the work include the necessary controls?
If not, please specify which controls are required in your comments to the authors.

Yes

Are the conclusions drawn adequately supported by the data shown?
If not, please explain in your comments to the authors.

Yes

Are you able to assess any statistics in the manuscript or would you recommend an additional statistical review?
If an additional statistical review is recommended, please specify what aspects require further assessment in your comments to the editors.

Not relevant to this manuscript

Quality of written English
Please indicate the quality of language in the manuscript:
Acceptable

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