First interchromosomal insertion in a patient with cerebral and spinal cavernous malformations

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Supplementary Table S1 Clinical characteristics of the index patient from family 1 and seven additional CCM patients analyzed in this study.

|                                | N = 8 |
|--------------------------------|-------|
| Mean age at genetic analysis ± standard deviation in years (range) | 40.0 ± 19.4 (7 - 63) |
| Sex (male : female)            | 2 : 6 |
| Family historya (familial : sporadic) | 1 : 7 |

a One CCM index patient was classified as familial case based on the presence of family members with clinical symptoms that were suggestive of CCM disease (seizures, hemorrhagic stroke).