



Supplementary Figure 3. Genotype–phenotype correlation of *RNF213* R4810K variant in moyamoya disease. (A) All patients with moyamoya disease. (B) Patients with adult moyamoya disease. (C) Patients with pediatric moyamoya disease. *RNF213*, ring finger protein 213; GG, wild type; GA, heterozygote; AA, homozygote; FHx, family history; TIA, transient ischemic attack; ICH/IVH, intracranial cerebral hemorrhage/intraventricular hemorrhage; PCA, posterior cerebral artery.