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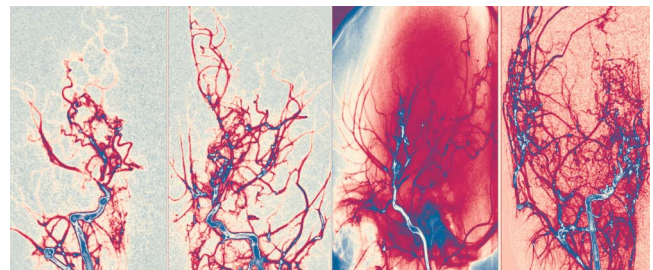
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Cerebral arteriopathy associated with a novel heterozygous mutation in MYH11 in a child with Moyamoya-like cerebrovascular disease. Colorized by Cindy Abair, *Neurology* Senior Graphics Editor.

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