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Moyamoya syndrome associated with polycystic kidney disease – a rare case report and literature review

Syndrom moyamoya doprovázený polycystickou chorobou ledvin – kazuistika vzácného onemocnění a přehled literatury

Dear editorial office,
Moyamoya disease is a chronic cerebrovascular disorder which is characterized by progressive stenoses and/or occlusions of the intracranial internal carotid artery (ICA) and its proximal branches, involving the development of a basal collateral network [1]. Conditions such as atherosclerosis, polycystic kidney disease (PKD), neurofibromatosis, or meningitis have been well recognized to be associated with moyamoya syndrome [2]. PKD is characterized by the presence of bilateral renal cysts and is always associated with vascular abnormalities including intracranial aneurysms, dilatation of the aortic root, and dissection of the thoracic aorta [3]. We herein report a rare case of moyamoya syndrome associated with PKD.

A 34-year-old male patient suddenly suffered from an intensive headache 2 months ago and lost consciousness 2 days later. CT revealed intraventricular hemorrhage. The headache reoccurred 1 month after the first episode and CT revealed intraventricular hemorrhage again. Digital subtraction angiography revealed typical bilateral findings of moyamoya disease with extensive cortical brain collateralization (Fig 1). The abdominal CT scan showed the presence of bilateral multiple renal cysts (Fig 2). The patient was treated with combined revascularization surgery (superficial temporal artery-middle cerebral artery anastomosis and encephalo-duro-arterio-synangiosis) without neurologic deterioration. His father, older sister, and older brother were also diagnosed with PKD.

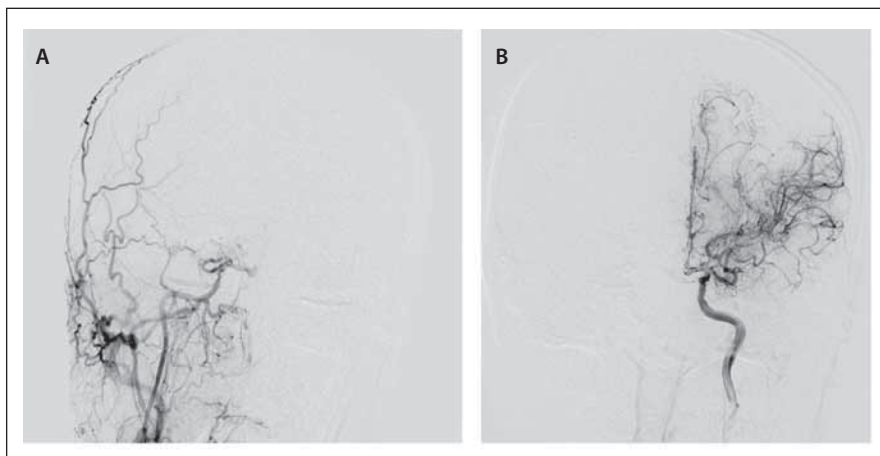


Fig. 1. DSA images of bilateral carotid arteries in anteroposterior view. Occlusion of the distal segment of the internal carotid arteries with moyamoya vessels at the skull base (A, B).

Obr. 1. DSA karotid v předozadním zobrazení. Oboustranná okluze distálních segmentů vnitřních karotid s cévněním typu moyamoya u baze lebni (A, B).

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In moyamoya syndrome, vessel stenosis or occlusion is accompanied by an abnormal vascular network associated with an underlying disease [4]. The annual incidence of moyamoya syndrome is 0.11/100,000, and its prevalence is 4/100,000 [5]. Underlying diseases include atherosclerosis (29%), Down syndrome (15.1%), von Recklinghausen disease

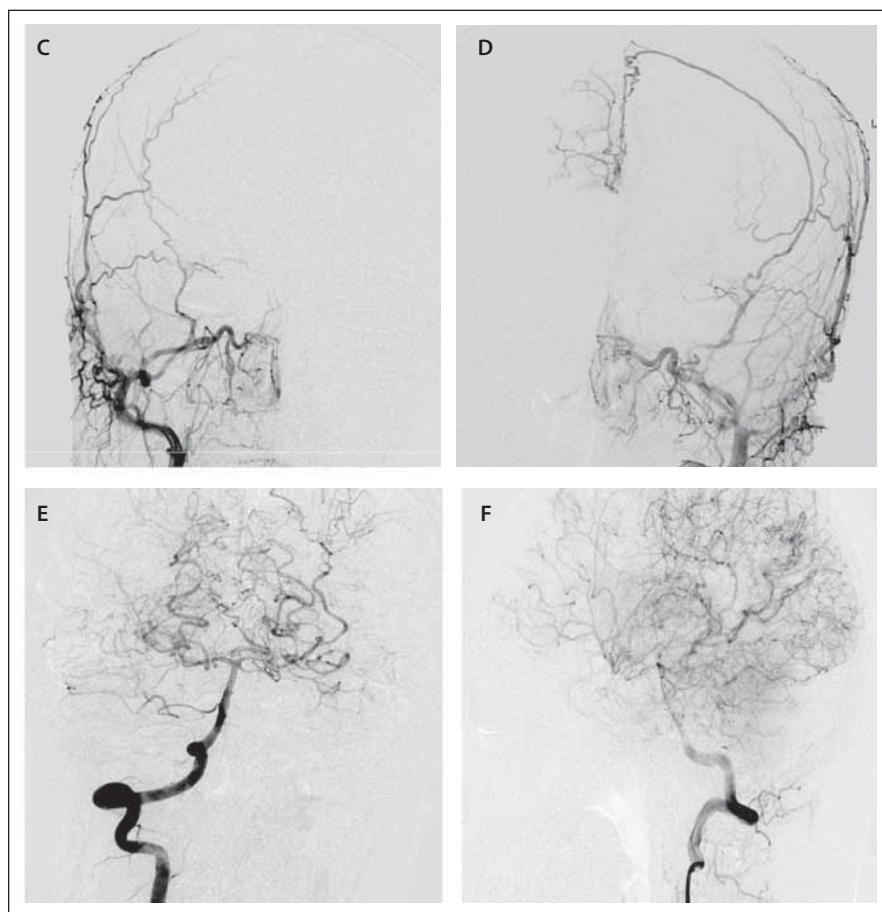


Fig. 1 – continuing. Anteroposterior view of bilateral external carotid arteries suggesting partial compensation of cerebral blood supply by the left external carotid artery (C, D). Angiograms of vertebral arteries demonstrated bilateral vertebral arteries partially compensate cerebral blood supply (E, F).

Obr. 1 – pokračování. Předozadní zobrazení vnějších karotid na obou stranách svědčí pro částečnou kompenzaci krevního zásobení mozku levou vnější karotidou (C, D). Angiogramy vertebrálních tepen prokázaly, že vertebrální tepny na obou stranách částečně kompenzovaly krevní zásobení mozku.

(14%), brain tumor or brain irradiation (7.5%), autoimmune disease (7.5%), and hyperthyroidism (7.5%) [6]. PKD is characterized by bilateral renal cysts and other abnormalities, including vascular disease. Vascular manifestations include intracranial and other arterial aneurysms and, more rarely, dolichoectasia, dilation of the aortic root, dissection of the thoracic aorta, and, possibly, coronary artery aneurysms [7]. PKD is also known as one of the underlying diseases of moyamoya syndrome which was rarely reported. Only 2 cases of moyamoya syndrome associated with PKD have been reported in the literature. In 1988, Pracyk et al reported the first case of a female patient whose arteriogram showed bilateral occlusion of the ICAs with “puff of smoke” collateralization arising from the circle of Willis. She was also diagnosed with arterial hypertension and eosinophilic granuloma. The patient finally died of the right putaminal hemorrhage with extension into the ventricle [8]. The second case was reported in 2012. The patient was diagnosed with moyamoya syndrome with PKD. Neurologic examination showed choreoathetosis in the right side of the body. Her twin sister was diagnosed with moyamoya syndrome with an intestinal duplication cyst [9]. Generally, moyamoya disease presents with 2 main types of symptoms: ischemia and hemorrhage [2]. For hemorrhagic moyamoya disease, a cause of hemorrhage may be cerebral aneurysms [10]. PKD confers the highest risk of intracranial berry aneurysm. Pracyk et al believed that rupture of an aneurysm could account for the cerebral hemorrhage found in their patient [8]. Our case included intraventricular hemorrhage which may also be due to a microaneurysm rupture. Only 1 case included choreoathetosis

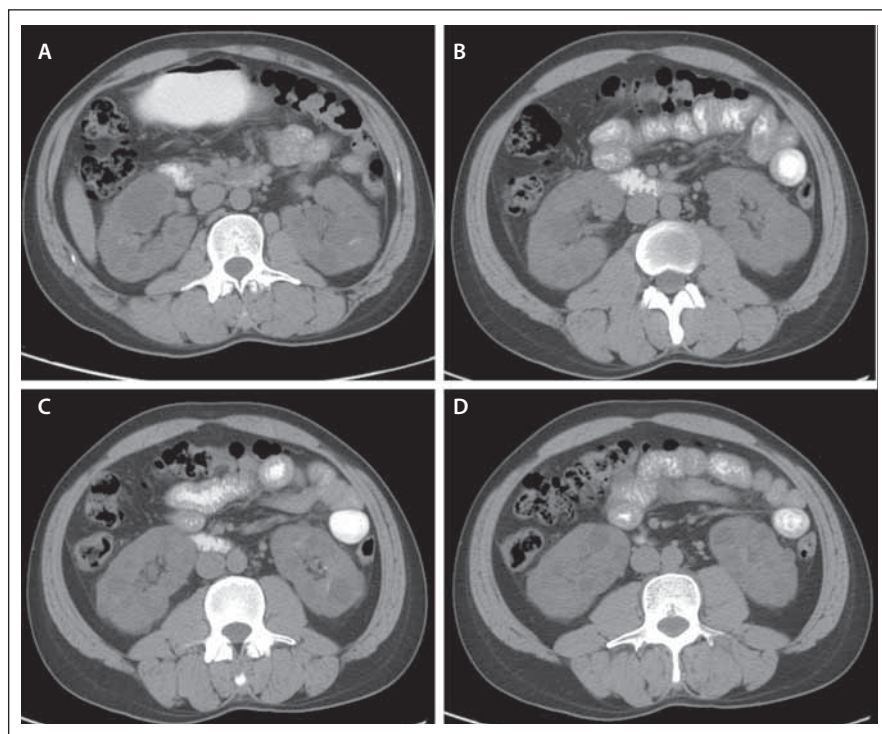


Fig. 2. Abdominal CT images. Bilateral renal polycystosis.

Obr. 2. CT snímky břišní dutiny. Polycystóza ledvin oboustranně.

which may be due to ischemia. This may indicate that patients with moyamoya syndrome associated with PKD were prone to cerebral hemorrhage.

The prevalence of moyamoya disease has an ethnic bias, with a high incidence of the disease in Eastern Asian countries such as Japan, Korea and China. And this is the first report of an Asian person with PKD associated with moyamoya syndrome.

Disclosures

The authors declare they have no potential conflicts of interest concerning drugs, products, or services used in the study.

References

1. Suzuki J, Takaku A. Cerebrovascular "moyamoya" caucasian disease: disease showing abnormal net-like vessels in base of brain. *Arch Neurol* 1969; 20(3): 288–299. doi: 10.1001/archneur.1969.00480090076012.
2. Scott RM, Smith ER. Moyamoya disease and moyamoya syndrome. *N Engl J Med* 2009; 360(12): 1226–1237. doi: 10.1056/NEJMra0804622.
3. Harris PC, Torres VE. Polycystic kidney disease, autosomal dominant synonym: ADPKD. Seattle, WA, USA: GeneReviews Initial 2002.
4. Hashimoto N, Tominaga T, Miyamoto S et al. Guidelines for diagnosis and treatment of moyamoya disease (spontaneous occlusion of the circle of Willis). *Neurol Med Chir (Tokyo)* 2012; 52(5): 245–266. doi: 10.2176/nmc.52.245.
5. Hayashi K, Horie N, Suyama K et al. An epidemiological survey of moyamoya disease, unilateral moyamoya disease and quasi-moyamoya disease in Japan. *Clin Neurol Neurosurg* 2013; 115(7): 930–933. doi: 10.1016/j.clineuro.2012.09.020.
6. Hayashi K, Horie N, Izumo T et al. Nationwide survey on quasi-moyamoya disease in Japan. *Acta Neurochir (Wien)* 2014; 156(5): 935–940. doi: 10.1007/s00701-014-2013-0.
7. Pirson Y, Chauveau D, Torres V. Management of cerebral aneurysms in autosomal dominant polycystic kidney disease. *J Am Soc Nephrol* 2002; 13(1): 269–276.
8. Pracyk JB, Massey JM. Moyamoya disease associated with polycystic kidney disease and eosinophilic granuloma. *Stroke* 1989; 20(8): 1092–1094. doi: 10.1161/01.str.20.8.1092.
9. Nzwalo H, Santos V, Gradil C et al. Caucasian familial moyamoya syndrome with rare multisystemic malformations. *Pediatr Neurol* 2013; 48(3): 240–243. doi: 10.1016/j.pediatrneuro.2012.11.009.
10. Kawaguchi S, Sakaki T, Morimoto T et al. Characteristics of intracranial aneurysms associated with moyamoya disease: a review of 111 cases. *Acta Neurochir (Wien)* 1996; 138(11): 1287–1294. doi: 10.1007/BF01411057.

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