Aortic Dissection in Familial Patients with Autosomal Dominant Polycystic Kidney Disease

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Autosomal dominant polycystic kidney disease (ADPKD) is the most common congenital kidney disease. However, reports on occasional cases of aortic dissection in PKD familial patients remain scarce. Herein, we describe rare aortic dissection cases in PKD familial patients (i.e., mother and daughter) and our successful treatment experience. The mother (84 years old) and daughter (53 years old) had a referral to us to treat type A acute aortic dissection. We performed emergency surgery and successfully treated the patients with an artificial graft. For comprehensive evaluation and treatment, ADPKD patients and their families should be screened for aortic diseases.

Keywords: aortic dissection, ADPKD

Case Reports

A 53-year-old woman came to the emergency department for a sudden onset of back pain. The patient had a medically managed hypertension from five years ago. Computed tomography (CT) showed acute type B aortic dissection, polycystic kidney, and multiple hepatic cysts. Blood tests on admission revealed a creatinine level of 0.95 mg/dl and an estimated glomerular filtration rate (eGFR) of 48.5 ml/min/1.73 m². The patient received a referral to our center for medical treatment. Sixteen days later, the patient suffered from severe chest pain. A CT showed acute type A aortic dissection (Figs. 1A and 1B), and the patient underwent emergency surgery. A dissection entry was detected from the ascending aorta to the distal aorta. The patient underwent a total aortic arch replacement with a 3-branched artificial graft (J-Graft, Japan Lifeline, Tokyo, Japan) and a Frozen elephant trunk (FROZENIX, Japan Lifeline). She did not require postoperative hemodialysis. The patient was discharged one month postoperatively. Histological analysis showed rupture of the internal elastic lamina and tunica media and inflow of blood. There was a high degree of atherosclerosis in the tunica intima and media. Elastica van Gieson staining showed no medial cystic necrosis (Figs. 2A and 2B).

One week from discharge, the 84-year-old patient’s mother was referred by another hospital to our center for type A aortic dissection. CT showed type A aortic dissection, polycystic kidney, and multiple hepatic cysts (Figs. 1C and 1D). A dissection entry was found in the ascending aorta. Blood tests on admission revealed a creatinine level of 3.82 mg/dl and an eGFR of 9.3 ml/min/1.73 m². The patient had medically managed hypertension from 50 years of age with a history of PKD and renal failure. An urgent ascending aortic replacement was performed using a straight artificial graft (J-Graft, Japan Lifeline). Although the patient required transient hemodialysis in the intensive care unit, it was unnecessary at discharge. The patient was moved to another hospital for rehabilitation one month postoperatively. Histological analysis showed aortic dissection, rupture of elastic fibers, and medial cystic necrosis (Figs. 2C and 2D). The postoperative magnetic resonance
imaging showed no cerebral aneurysms in both patients. Her brother had died of aortic dissection several years earlier, but his detailed medical history was unknown.

Discussion

ADPKD is the most common congenital renal cystic disease with an estimated prevalence of between one in 1000 and one in 2500 individuals. Its course is characterized by the development and inexorable expansion of multiple cysts scattered throughout the kidney parenchyma. ADPKD patients have a 10% incidence of intracranial aneurysms. Mitral valve prolapse occurs in up to 26% of PKD-1 patients. Aortic dissection is a rare complication of ADPKD, and there are few reports on the frequencies of aortic aneurysm and aortic dissection. ADPKD is genetically heterogeneous, and the PKD 1 and PKD 2 genes’ mutation contributes to its development. Mutations in PKD genes that encode polycystin, which is often expressed in vascular smooth muscle, including the kidney, are suggested to cause comorbid cysts and cardiovascular abnormalities. In a mouse model research, PKD 1 has been implicated in maintaining vessel wall structural integrity.

This is a rare report of aortic dissection in ADPKD familial patients successfully repaired with an artificial graft to the best of our knowledge. The congenital polycystic kidney and multiple renal cysts detected by CT fulfill the ADPKD criteria of Ravine et al. An interesting finding is that the three family members similarly suffered from an aortic dissection. The reason for the histological difference between the mother and the daughter is unclear. Although medial cystic necrosis appeared in the mother’s aortic wall, it was undetected in the daughter. Cystic medial necrosis may not necessarily cause aortic dissection. Other factors, including changes in the extracellular matrix or cellular interaction from PKD gene mutation, may cause aortic dissection.

The mother and daughter were diagnosed with ADPKD before the onset of acute aortic dissection (AAD), implying an important finding that the cause of their AAD was ADPKD. Hypertension and renal failure associated with ADPKD were considered significant risk factors for aortic dissection following arteriosclerosis. A systematic review described a markedly higher frequency of hypertension and younger age in aortic dissection patients with ADPKD than the overall aortic dissection population. The earlier aortic dissection manifestation and the possible lack of symptoms suggestive of aortic dissection in ADPKD patients underline the importance of performing close clinical screening from a young age. Antihypertensive therapy may prevent renal failure in ADPKD patients.

Regarding the emergency treatment of acute aortic dissection, antihypertensive therapy is crucial for preventing the progress of the dissection and organ malperfusion. The renin angiotensin system’s activity is reportedly increased in ADPKD. Antihypertensive therapy using an angiotensin-converting-enzyme inhibitor may be effective in acute aortic dissection patients with ADPKD.

In conclusion, we successfully treated acute type A aortic dissection in ADPKD familial patients. For comprehen-
sive evaluation and treatment, ADPKD patients and their families should be screened regularly for aortic diseases.

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References
1) Cornec-Le Gall E, Alam A, Perrone RD. Autosomal dominant polycystic kidney disease. Lancet 2019; 393: 919-35.
2) Zhou Z, Xu Y, Delcourt C, et al. Is regular screening for intracranial aneurysm necessary in patients with autosomal dominant polycystic kidney disease? A systematic review and meta-analysis. Cerebrovasc Dis 2017; 44: 75-82.
3) Lumiaho A, Ikaheimo R, Miettinen R, et al. Mitral valve prolapse and mitral regurgitation are common in patients with polycystic kidney disease type 1. Am J Kidney Dis 2001; 38: 1208-16.
4) Hassane S, Claij N, Lantinga-van Leeuwen IS, et al. Pathogenic sequence for dissecting aneurysm formation in a hypomorphic polycystic kidney disease 1 mouse model. Arterioscler Thromb Vasc Biol 2007; 27: 2177-83.
5) Ravine D, Sheffield L, Danks DM, et al. Evaluation of ultrasonographic diagnostic criteria for autosomal dominant polycystic kidney disease 1. Lancet 1994; 343: 824-7.
6) Silverio A, Prota C, Di Maio M, et al. Aortic dissection in patients with autosomal dominant polycystic kidney disease: a series of two cases and a review of the Literature. Nephrology (Carlton) 2015; 20: 229-35.
7) Schrier RW, McFann KK, Johnson AM. Epidemiological study of kidney survival in autosomal dominant polycystic kidney disease. Kidney Int 2003; 63: 678-85.
8) Loghman-Adham M, Soto CE, Inagami T, et al. Expression of components of the renin-angiotensin system in autosomal recessive polycystic kidney disease. J Histochem Cytochem 2005; 53: 979-88.