Macroscopical and microscopical findings of polycystic kidney disease in a cat

Shokrpoor S.^{1*}; Ebrahimi Toori N.²; Torjani N.²; Jarideh M.²

Received: March 2022

Accepted: May 2022

Abstract

Autosomal dominant polycystic kidney disease (AD-PKD) is a genetic feline disease characterized by fluid-filled cysts formation in one or both kidneys. Persians and Persian-related cats can be affected. A five-year-old male Persian cat weighing 6.5 kg with severe bilateral abdominal swelling was referred to the Clinic. Radiography and ultrasonography were performed. Finally, the cat was euthanized due to the severity of clinical signs, and necropsy was performed. Histopathologically, cystic structures filled with eosinophilic material lined by cuboidal or squamous epithelium were seen. Based on ultrasonographic images, clinical, radiological and histopathological findings, this disease was diagnosed polycystic kidney disease.

Keywords: Cat, Necropsy, Pathology, PKD

¹⁻Department of Pathology, Faculty of Veterinary Medicine, University of Tehran, Tehran, Iran

²⁻Faculty of Veterinary Medicine, University of Tehran, Tehran, Iran

^{*}Corresponding author's Email: shokrpoor@ut.ac.ir

Introduction

Polycystic kidney disease is a common hereditary disease in human being and is one of the leading causes of end-stage renal disease that is characterized by progressive formation of multiple liquid-filled cysts in kidney (Rahmati-Holasoo et al., 2015). Pathogenesis of cystic lesions of animals is usually congenital or parasitic (Rahmati-Holasoo et al., 2020). Autosomal dominant polycystic kidney disease (AD-PKD) is one of the most common life-threatening inherited disorders in cats, characterized by abnormal fluidfilled cysts formation in one or both kidneys, sometimes, in other organs such as liver and pancreas.

The affected cats exhibit clinical signs, including vomiting, anorexia, fatigue, polvuria and polydipsia (Michel-Regalado et al., 2022). The renal cysts are congenital, can arise from any part of the nephron, are highly variable in number and are initially very small, but tend to increase in size with age (Bosje et al., 1998; Norman, 2011; Bilgen et al., 2020; Guerra et al., 2020). The Persian cat is affected by AD-PKD. Other breeds such as the Chartreux, Maine Coon and Neva Masquerade breeds can be affected (Jasik and Kulesza, 2014). Recently, in America and Germany were found that 95-100% respectively of Persian cats diagnosed with polycystic kidney disease (PKD) by ultrasound scanning had the mutation identified by polymerase chain reaction restriction fragment polymorphism (PCR-RFLP) length (Lyons et al., 2004; Kappe et al., 2005).

Clinical findings, radiography, ultrasonography and histopathological examination can guide the diagnosis of disease.

Materials and methods

In August 2021, a five-year-old male Persian cat weighing 6.5 kg with severe bilateral abdominal swelling and a clinical history of lethargy, anorexia, polyuria, polydipsia and vomiting was referred to the Clinic. Physical examination, blood cell counts, serum biochemistry tests and urinalysis were performed. Cat was restrained by his owner in the awake state and sedated with ketamine (10 mg/kg), administered intramuscularly, and ultrasound examination was performed on cat. Dorsoventral digital radiograph was taken. Finally, the cat was euthanized due to the severity of clinical signs, and necropsy was performed. For histopathological examinations. samples of kidneys and internal organs were dissected and preserved in 10% buffered formalin, dehydrated and embedded in paraffin (with paraffin tissue processor and paraffin dispenser), sectioned at 5 μ m, stained with hematoxylin and eosin, and observed with light microscopy.

Result

Hematology and biochemistry results showed mild leukocytosis, mild anemia and severe azotemia. Dorsoventral radiograph showed bilateral abdominal distention (Fig. 1a). Ultrasonography revealed significant enlargement of both kidneys with multiple anechoic or

cysts hypoechoic (Fig. 1b). At surface, the necropsy, on cut sonographically diagnosed polycystic structure was apparent, and the cystic cavities were contained varying amounts of water-like fluid (Fig. 1c-d). Histopathological examination of the kidneys revealed cystic structures of tubules. The cystic tubules were lined by cuboidal or squamous epithelium and separated by a band of fibrous connective tissue (Fig. 2a-b). Some

cysts were filled with eosinophilic and amorphic material (Fig. 2c). Birefringent crystals were observed with polarized and non-polarized light (Fig. 2b, d).



Figure 1: (a-d) Radiography, ultrasonography and Necropsy findings of polycystic kidney disease in a Persian cat. (a) Dorsoventral digital radiograph showing abdominal distention, left kidney (L), (b) Ultrasonographic image showing multiple fluid-filled cysts of varying size, (c) Gross Appearance of affected kidneys, (d) Numerous variably sized cysts in the cortex and medulla.

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Figure 2: (a-d) Histopathological findings of polycystic kidney disease in a Persian cat. (a) Multiple cysts (C) of varying size, (b) Some cysts (C) contain crystals (arrowheads), (c) cysts contain proteinaceous material (arrowheads), (d) Note the birefringent radiating crystal (arrowhead) in renal tubule. Polarized light. H&E stain.

Discussion

The important clinical most manifestation of PKD in cats is chronic renal failure. As cysts enlarge and increase in size, they compress surrounding normal renal parenchyma and lesions of chronic interstitial nephritis occur. For these reasons, PKD is considered a progressive disease, and the prognosis is guarded (Biller et al., 1996; Barrs et al., 2001). Ultrasonography is one of the main methods for detection of PKD in human and cat (Chapman, 2007; Domanjko-Petrič et al., 2008). Ultrasound as a noninvasive technique is the most commonly used imaging modality for diagnosis of PKD in cat (Guerra et al., 2019; Noori et al., 2019). Similar to our case, ultrasound was of value in the diagnosis of PKD in a Chartreux cat (Volta et al., 2010). Ultrasonographic radiological images. clinical. and histopathological findings in this case are similar to the other studies (Volta et al., 2010; Gendron et al., 2013; Jasik and Kulesza, 2014; Schirrer et al., 2021). In conclusion, the progressive nature of PKD that leads to irreversible renal failure, should raise the interest of veterinarians and breeders of cats that have current or past links with Persians. Although renal cysts may be identified in kittens, their absence does not preclude detection at an older age. In fact, the sensitivity of renal cyst detection by ultrasonography increases with the age, because in affected cats, cysts enlarge with age, and this is an imperative factor in screening and breeding programs. This report points out that PKD should be considered as a possible cause of renal failure and Persian cats should be included in the screening program of PKD.

Acknowledgment

Mr. Samani is thanked for assisting with the histopathological sections.

References

- Barrs, V.R., Gunew, M., Foster, S.F., Beatty, J.A. and Malik, R., 2001. Prevalence of autosomal dominant polycystic kidney disease in Persian cats and related-breeds in Sydney and Brisbane. *Australian Veterinary Journal*, 79(4), pp.257-259. https://doi.org/10.1111/j.1751-0813.2001.tb11977.x
- Bilgen, N., Bişkin Türkmen, M., Çınar Kul, B., Isparta, S., Şen, Y., Akkurt, M.Y., Çıldır, Ö.Ş. and Bars, Z., 2020. Prevalence of PKD1 gene mutation in cats in Turkey and pathogenesis of feline polycystic kidney disease. Journal of veterinary diagnostic investigation, 32(4), pp.549-555.
- https://doi.org/10.1177/1040638720935 433
- Biller, D.S., DiBartola, S.P., Eaton, K.A., Pflueger, S., Wellman, M.L. and Radin, M.J., 1996. Inheritance of polycystic kidney disease in Persian cats. *Journal of Heredity*, 87(1), pp.1-5. https://doi.org/10.1093/oxfordjournal s.jhered.a022945
- Bosje, J.T., Van den Ingh, T.S.G.A.M. and Van der Linde-Sipman, J.S., 1998. Polycystic kidney and liver disease in cats. *Veterinary Quarterly*, 20(4), pp.136-139.

https://doi.org/10.1080/01652176.19 98.9694858

Chapman, A.B., 2007. Autosomal dominant polycystic kidney disease:

time for a change? *Journal of the American Society of Nephrology*, 18(5), pp.1399-1407. https://doi.org/10.1681/ASN.200702 0155

- Domanjko-Petrič, A., Černec, D. and Cotman, M., 2008. Polycystic kidney disease: а review and occurrence in Slovenia with comparison between ultrasound and genetic testing. Journal of Feline Medicine & Surgery, 10(2), pp.115-119.https://doi.org/10.1016/j.jfms.20 07.07.004
- Gendron, K., Owczarek-Lipska, M., Lang, J. and Leeb, T., 2013. Maine Coon renal screening: ultrasonographical characterisation and preliminary genetic analysis for common genes in cats with renal cysts. *Journal of Feline Medicine & Surgery*,15(12), pp.1079-1085. https://doi.org/10.1177/1098612X13 492164
- Guerra, J.M., Cardoso, N.C., Daniel, Onuchic, A.G.T., L.F. and Cogliati, B., 2020. Prevalence of polycystic autosomal dominant kidney disease in Persian and Persian-related cats in Brazil. Brazilian journal of biology, 81, pp.392-397.

https://doi.org/10.1590/15196984.22 7131

- Guerra, J.M., Freitas, M.F., Daniel, A.G., Pellegrino, A., Cardoso, N.C., de Castro, I., Onuchic, L.F. and Cogliati, B., 2019. Age-based ultrasonographic criteria for diagnosis of autosomal dominant polycystic kidney disease in Persian cats. Journal of Feline Medicine and Surgery, 21(2), pp.156-164. https://doi.org/10.1177/1098612X18 764591
- Jasik, A. and Kulesza, M., 2014. Polycystic kidney disease in a Neva Masquerade cat. *Journal of Small*

Animal Practice, 55(**7**), pp.387-387. https://doi.org/10.1111/jsap.12240

- Kappe, E.C., Hecht, W., Gerwing, M., Michele, U. and Reinacher, M., 2005. Polycystic kidney disease in the German population of Persian cats. A comparative study of ultrasonographical examination and genetic testing. *Tieraerztliche Praxis Ausgabe Kleintiere Heimtiere*, 33(6), pp.413-418.
- Lyons, L.A., Biller, D.S., Erdman, C.A., Lipinski, M.J., Young, A.E., Roe, B.A., Qin, B. and Grahn, R.A., 2004. Feline polycystic kidney disease mutation identified in PKD1. *Journal of the American Society of Nephrology*, 15(10), pp.2548-2555. https://doi.org/10.1097/01.ASN.0000 141776.38527.BB
- Noori, Z., Moosavian, H.R., Esmaeilzadeh, H., Vali, Y. and Fazli, M., 2019. Prevalence of polycystic kidney disease in Persian and Persian related-cats referred to Small Animal Hospital, University of Tehran, Iran. *Iranian Journal of Veterinary Research*, 20(2), pp. 151-154.
- Michel-Regalado, N.G., Ayala-Valdovinos, M.A., Galindo-García, J., Duifhuis-Rivera, T. and Virgen-Méndez, A., 2022. Prevalence of polycystic kidney disease in Persian and Persian-related cats in western Mexico. Journal of Feline Medicine and Surgery, p.1098612X221114043.

https://doi.org/10.1177/1098612X22 1114043

Norman, J., 2011. Fibrosis and progression of autosomal dominant polycystic kidney disease (ADPKD). *Biochimica et Biophysica Acta* (*BBA*)-*Molecular Basis of Disease*, 1812(10), pp.1327-1336. https://doi.org/10.1016/j.bbadis.2011 .06.012

- Rahmati- Holasoo, H., Masoudifard, M., Ebrahimzadeh Mousavi, H., Shokrpoor, S., Tavakkoli, A. and Farazandemehr, M. S. (2015). Cystic lesions in the kidney of flower horn fish. hvbrid cichlid. Journal of Fish Diseases, 38(9), pp. 833-838. https://doi.org/10.1111/jfd.12292
- Rahmati-Holasoo, H., Shahbazi, M., Ebrahimzadeh Mousavi, H.A. Shokrpoor, S., **Pourmortazavi** Bahambari, M. and Mokhatari, A. 2020. Polycystic kidney disease in discus (Symphysodon aequifasciatus) and Siamese fighting fish (Betta splendens): А histopathological study. Bulletin of the European Association Fish ofPathologists, 40(1), pp. 39-44.
- Schirrer, L., Marín-García, P. J., and Llobat, L. 2021. Feline polycystic kidney disease: an update. *Veterinary Sciences*, 8(11), pp. 269. https://doi.org/10.3390/vetsci811026 9
- Volta, A., Manfredi, S., Gnudi, G., Gelati, A. and Bertoni, G., 2010. Polycystic kidney disease in a Chartreux cat. *Journal of Feline Medicine and Surgery*, 12(2), pp.138-140. https://doi.org/10.1016/j.jfms.2009.0 6.001