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### Prevalence of Spinal Meningeal Diverticula in Autosomal Dominant Polycystic Kidney Disease

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#### ABSTRACT

**BACKGROUND AND PURPOSE:** Patients with autosomal dominant polycystic kidney disease (ADPKD) develop cysts in the kidneys, liver, spleen, pancreas, prostate, and arachnoid spaces. In addition, spinal meningeal diverticula have been reported. To determine whether spinal meningeal diverticula are associated with ADPKD, we compared their prevalence in subjects with ADPKD with a control cohort without ADPKD.

**MATERIALS AND METHODS:** Subjects with ADPKD and age- and sex-matched controls without ADPKD undergoing abdominal MRI from the midthorax to the pelvis from 2003 to 2023 were retrospectively evaluated for spinal meningeal diverticula by 4 blinded observers. The prevalence of spinal meningeal diverticula in ADPKD was compared with that in control subjects, using *t* tests and correlated with clinical and laboratory data and MR imaging features, including cyst volumes and cyst counts.

**RESULTS:** Identification of spinal meningeal diverticula in ADPKD (n = 285, median age, 47; interquartile range [IQR], 37–56 years; 54% female) and control (n = 285, median age, 47; IQR, 37–57 years; 54% female) subjects had high interobserver agreement (pairwise Cohen  $\kappa = 0.74$ ). Spinal meningeal diverticula were observed in 145 of 285 (51%) subjects with ADPKD compared with 66 of 285 (23%) control subjects without ADPKD (P < .001). Spinal meningeal diverticula in ADPKD were more prevalent in women (98 of 153 [64%]) than men (47 of 132 [36%], P < .001). The mean number of spinal meningeal diverticula per affected subject with ADPKD was 3.6  $\pm$  2.9 compared with 2.4  $\pm$  1.9 in controls with cysts (P < .001). The median volume (IQR, 25%–75%) of spinal meningeal diverticula was 400 (IQR, 210–740) mm<sup>3</sup> in those with ADPKD compared with 250 (IQR, 180–440) mm<sup>3</sup> in controls (P < .001). The mean spinal meningeal diverticulum diameter was greater in the sacrum (7.3 [SD, 4.1] mm) compared with thoracic (5.4 [SD, 1.8] mm) and lumbar spine (5.8 [SD, 2.0] mm), (P < .001), suggesting that hydrostatic pressure contributed to enlargement.

CONCLUSIONS: ADPKD has a high prevalence of spinal meningeal diverticula, particularly in women.

 $\label{eq:abstraction} \textbf{ABBREVIATIONS:} \ \textbf{ADPKD} = \textbf{autosomal dominant polycystic kidney disease; } \textbf{IQR} = \textbf{interquartile range}$ 

A utosomal dominant polycystic kidney disease (ADPKD) is characterized by multiorgan cystic (eg, kidney, liver) and noncystic (eg, pericardial, pleural) fluid accumulations.<sup>1–7</sup> Although most ADPKD imaging focuses on the kidneys, the spine is within the abdominal field of view, enabling the evaluation of nerve roots.

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Indicates article with online supplemental data. http://dx.doi.org/10.3174/ajnr.A8407 Patients with ADPKD are known to develop intracranial arachnoid cysts.<sup>8–11</sup> We have also noticed that spinal meningeal diverticula appear to be more prevalent in patients with ADPKD, and there have been multiple case reports.<sup>12–18</sup>

Spinal meningeal diverticula, also known as nerve root cyst or perineural cyst, are focal fluid-filled outpouchings along the nerve root that directly communicate with the subarachnoid space and are covered with arachnoid and dura mater. The etiology is unknown, but increased prevalence in connective tissue disorders and spinal injuries has been reported.<sup>19–21</sup> One hypothesis is that inflammation and a ball valve mechanism between the nerve root sheath and subarachnoid space result from pulsatile hydrodynamic forces on the CSF.<sup>21</sup> Most spinal meningeal diverticula are asymptomatic, though the clinical presentation can include headache exacerbated by upright posture,<sup>12,14,16,17</sup> low back pain,<sup>22–25</sup> sciatic pain,<sup>23–25</sup> leg weakness,<sup>23–25</sup> bowel and bladder dysfunction,<sup>23–25</sup> claudication,<sup>23,24</sup> and sacral insufficiency fractures.<sup>26</sup> Spinal

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#### **SUMMARY**

**PREVIOUS LITERATURE:** Schievink and Torres first reported spinal meningeal diverticula in 3 women with ADPKD being evaluated for headache, visual blurring aggravated by upright positioning, and low CSF pressure. Since then, there have been additional reports of spinal CSF leaks related to spinal meningeal diverticula in subjects with ADPKD. Asymptomatic spinal meningeal diverticula have also been reported in patients with ADPKD, all of whom were women. Aşik et al reported more and larger cysts in 50 subjects with ADPKD compared with 37 controls without ADPKD and correlated the number and size of spinal meningeal diverticula with a headache severity score.

**KEY FINDINGS:** Spinal meningeal diverticula were observed in 51% of patients with ADPKD on abdominal MRI, which is 2.2 times greater than that in a control population without ADPKD, (P < .001) and 1.8 times more common in women (98 of 153 [64%]) than men (47 of 132 [36%], P < .001).

**KNOWLEDGE ADVANCEMENT:** Spinal meningeal diverticula are prevalent in subjects with ADPKD and should be included in the differential diagnosis of headache in ADPKD, especially when the headache is exacerbated or provoked by upright posture.

meningeal diverticulum formation may also be associated with idiopathic intracranial hypertension,<sup>27</sup> in which there is elevated intracranial pressure in the absence of hydrocephalus or mass lesions. Additionally, spinal meningeal diverticula can also be associated with spontaneous spinal CSF leaks, with CSF flowing directly into the surrounding epidural space (type 2 CSF leak) or flowing into an aberrant connection between the subarachnoid space and an adjacent paraspinal vein (type 3 CSF leak), resulting in spontaneous intracranial hypotension.<sup>28–30</sup>

In this STrengthening the Reporting of OBservational studies in Epidemiology checklist–guided study, we identified the prevalence of spinal meningeal diverticula in subjects with ADPKD compared with age- and sex-matched control subjects without ADPKD to determine whether these spinal meningeal diverticula are also associated with ADPKD.

#### MATERIALS AND METHODS

#### **Study Design and Populations**

This retrospective, cross-sectional study of existing data and images acquired from 2003 to 2023 is compliant with the Health Insurance Portability and Accountability Act (HIPAA) and was approved by the Weill Cornell Medicine Institutional Review Board. We retrospectively analyzed 285 patients with ADPKD and 285 age- and sex-matched control subjects without ADPKD. All subjects with ADPKD were enrolled in the Rogosin PKD Repository (https://clinicaltrials.gov/study/NCT01988038), signed informed consent, and underwent MR imaging as outpatients. The requirement for informed consent was waived for analysis of existing data from control patients.

Inclusion criteria for subjects with ADPKD were the following: 1) a diagnosis of ADPKD based on the Pei-Ravine criteria,<sup>31</sup> and 2) MRI with T2-weighted images covering from midthorax extending below the kidneys. Exclusion criteria were the following: 1) medical conditions associated with spinal meningeal diverticula including connective tissue disorder or spinal deformity, 2) an incomplete T2 MRI, and 3) laboratory data unavailable within 12 months of the MRI (Fig 1). For each patient with ADPKD, we identified age- (within 2.5 years) and sex-matched control patients without ADPKD, connective tissue disorder, or spinal deformity from the PACS with contemporaneous examination dates.

#### **Data Extraction**

ADPKD and control subjects' demographic information, clinical data, laboratory data, and genetic data were extracted from



FIG 1. Patient flow chart.

the PKD Repository and the electronic medical records for the date closest to the MRI date. We analyzed MRI reports prepared prospectively at the time of imaging to obtain data on kidney, liver, and spleen volumes, cyst counts/fractions, pancreatic cysts, prostate cysts, seminal megavesicles, and

# pleural effusions.

MRI examinations without contrast enhancement were obtained at 1.5T or 3T using a body array coil with the parameters shown in the Online Supplemental Data. Pulse sequences used for this analysis were coronal and axial T2-weighted single-shot FSE (Signa family; GE Healthcare) or HASTE (Magnetom Skyra; Siemens).



**FIG 2.** Coronal T2-weighted single-shot FSE images in 3 typical patients with ADPKD including a 60-year-old woman with T10-12 spinal meningeal diverticula bilaterally (*red arrows, A*) and an L1 spinal meningeal diverticulum on the right (*B*) in red (thoracic) and green (lumbar); *C*, A 46-year-old woman with lumbar spinal meningeal diverticula (*red arrows, C*) with a corresponding annotation (*D*) in green for L2 (right), L3 (*bilateral*), L4 (*left*), and L5 (*bilateral*); *E*, a 28-year-old female with sacral spinal meningeal diverticula (*red arrows, F*) in blue. Also, note renal cysts (*yellow arrows*) and liver with multiple cysts (*green arrow*).

#### **Image Analysis**

Axial and coronal T2 MR images were analyzed by 4 independent, experienced observers (U.S., X.Y., X.L., M.R.P.) blinded to the patient information. Observers independently counted the number of spinal meningeal diverticula present in each subject. Disagreements were resolved by consensus. Spinal meningeal diverticula were defined as present when there was a well-defined T2 bright signal corresponding to fluid intensity along the course of the normal nerve root on T2-weighted images at least doubling the normal nerve root diameter. For size measurement and region distribution, one of the observers (U.S.) annotated spinal meningeal diverticula on coronal images using ITK-SNAP software, Version 3.8.0 (www.itksnap. org) (Fig 2).

#### **Descriptive Statistics**

For normally distributed continuous variables (assessed by the Shapiro-Wilk test), means (SDs) were reported. For non-normal distributions, median and interquartile range (IQR) were reported. Frequency and percentage were calculated for categoric variables.

For 2-group matched continuous variables, a paired *t* test was used to assess the statistical significance. For multigroup continuous variables, the ANOVA was used to assess statistical significance. For categoric variables, a  $\chi^2$  or Fisher exact test was used to assess statistical significance, depending on sample size. Interobserver agreement for identifying spinal meningeal diverticula were assessed using the pair-wise Cohen  $\kappa$  and intraclass correlation coefficients, respectively.

#### **Regression Models**

Bivariate analysis was used to estimate the correlation between the presence of spinal meningeal diverticula and age, sex, height, weight, body mass index, body surface area, blood pressure, blood urea nitrogen level, creatinine level, estimated glomerular filtration rate, albumin level, aspartate transaminase and alanine transaminase levels, height-adjusted total kidney volume, height-adjusted total liver volume, height-adjusted total spleen volume, number of renal cysts, number of hemorrhagic renal cysts, number of

liver cysts, number of pancreatic cysts, pleural effusion, severe headache, and *PKD* genotype. The variables that had significance (ie, P < .05) were used in a multivariable linear regression analysis to assess their effect after adjusting for the other variables. The analysis was performed using GraphPad Prism software, Version 10.2.0392 (GraphPad Software).



**FIG 3.** A 73-year-old woman with ADPKD reporting a remote history of severe headache lasting 2 weeks. *A* and *B*, coronal, and *C*–*F*, sequential, axial T2-weighted single-shot FSE images show multiple large sacral spinal meningeal diverticula (*red arrows*). Also, note renal cysts (*yellow arrows*) and liver with multiple cysts (*green arrow*).

#### RESULTS

Abdominal MRI results were available in 285 subjects with ADPKD (median age, 47 [IQR, 3–56] years; women 54%, white 84%) and a control group without ADPKD controlled for age and sex (Fig 1 and Online Supplemental Data).

As expected, the ADPKD group had higher diastolic blood pressure, blood urea nitrogen and creatinine levels, a lower estimated glomerular filtration rate, and higher height-adjusted total kidney volume, height-adjusted total liver volume, height-adjusted total spleen volume, number of liver cysts, and number of kidney cysts (Online Supplemental Data). Information on race was available for more patients with ADPKD compared with control patients, reflecting the willingness of subjects with ADPKD to provide this information when enrolling in the Rogosin PKD Repository. This availability resulted a higher number of subjects with unknown race in the control population (Online Supplemental Data).

Indications for MRI in the control subjects included inflammatory bowel disease follow-up (n = 99), indeterminate lesion (n = 82), pain (n = 51), cancer follow-up (n = 23), liver disease (n = 15), pancreatitis (n = 7), hematuria (n = 4), small bowel obstruction followup (n = 2), endometriosis (n = 1), and uterine leiomyomas (n = 1).

#### Prevalence of Spinal Meningeal Diverticula in ADPKD

Spinal meningeal diverticula were identified in 51% of subjects with ADPKD compared with 23% of controls (Online Supplemental Data; P < .001). In the ADPKD group, 64% of all women had spinal meningeal diverticula compared with 36% of men (P < .001). In the control subjects, 31% of women had spinal meningeal diverticula compared with 14% of men (P = .001) (Online Supplemental Data).

In the ADPKD group, subjects with cysts had a mean of 3.6 (SD, 2.9) cysts per subject, which was 50% greater than the mean number in controls, (2.4 [SD, 1.9], P < .001). The median spinal meningeal diverticulum volume was 400 (IQR, 210-740) mm<sup>3</sup> in subjects with ADPKD with spinal meningeal diverticula compared with 250 (IQR, 180–440) mm<sup>3</sup> in controls (P < .001). The mean spinal meningeal diverticulum diameter was 6.6 (SD, 3.4) mm in subjects with ADPKD compared with 6.1 (SD, 2.6) mm in the control subjects (P = .06). Spinal meningeal diverticula were more prevalent in the sacrum compared with lumbar or thoracic spine locations (P < .001) (Online Supplemental Data). Large cysts (>10 mm) were more prevalent in the ADPKD group than the control group (35/285 [12.3%] versus 6/285 [2.1%]) (P < .001) (Online Supplemental Data).

The mean diameter of spinal meningeal diverticula was largest in the sacrum, (7.3 [SD, 4.1) mm, compared with the lumbar (5.8 [SD, 2.0] mm) and thoracic (5.4 [SD, 1.8] mm) spine, (P < .001) (Fig 3 and Online Supplemental Data).

As expected, the ADPKD group had a higher prevalence of pancreatic cysts, seminal megavesicles, and midline prostate cysts than the control group, as previously reported.<sup>5,6</sup> Serum albumin and aspartate transaminase levels were slightly higher in those with ADPKD compared with the control group; however, these values were within normal limits for both groups.

#### Interobserver Variability

Interobserver agreement for the presence of spinal meningeal diverticula was high, with a pair-wise Cohen  $\kappa = 0.74$ . Interobserver agreement for the number of spinal meningeal diverticula in each subject was excellent, with an intraclass correlation coefficient = 0.95.

#### Genotype

Among patients with ADPKD with either a *PKD1* (n = 187/245 = 76.3%) or *PKD2* (n = 58/245 [23.7%]) mutation,

Multivariate analysis for the presence of spinal meningeal diverticula in patients with  $\mbox{ADPKD}^{\rm a}$ 

Coefficients	95% CI (Asymptotic)	P Value
-0.18	-5.84-5.48	.94
-0.31	-0.481 to -0.14	<.001 <sup>b</sup>
0.91	-4.944-6.76	.76
0.003	-0.051-0.06	.92
-0.01	-0.107-0.09	.83
-0.24	-5.473-4.98	.92
0.004	-0.003-0.01	.31
-0.07	-0.161-0.02	.11
-0.00003	-0.0001-0.00004	.40
-0.0007	-0.001 to 0.0003	.17
0.00002	-0.0002 to 0.0002	.85
-0.0009	–0.0019 to 0.0000267	.06
0.093	-0.02503-0.2107	.12
	Coefficients -0.18 -0.31 0.91 0.003 -0.01 -0.24 0.004 -0.07 -0.00003 -0.0007 0.00002 -0.0009 0.093	Coefficients         95% CI (Asymptotic)           -0.18         -5.84-5.48           -0.31         -0.481 to -0.14           0.91         -4.944-6.76           0.003         -0.051-0.06           -0.01         -0.107-0.09           -0.24         -5.473-4.98           0.004         -0.003-0.01           -0.07         -0.161-0.02           -0.00003         -0.0001 to 0.0003           0.00002         -0.0001 to 0.0002           -0.0009         -0.0019 to 0.000267           0.093         -0.02503-0.2107

 $^{\rm a}$  Including parameters with P value < .05 on bivariate analysis (model P value =< .001).  $^{\rm b}$  Significant P value.

95 of 187 (51%) with *PKD1* mutations had spinal meningeal diverticula compared with 30 of 58 (52%) with *PKD2* mutations (Online Supplemental Data; P = .9). The prevalence of truncating mutations in *PKD1* was similar in those with and without spinal meningeal diverticula.

#### **Correlation with Laboratory and Imaging Parameters**

The presence of spinal meningeal diverticula on MRI in subjects with ADPKD was significantly correlated with female sex and severe headache by bivariate analysis (Online Supplemental Data). The presence of spinal meningeal diverticula negatively correlated with height, weight, body mass index, body surface area, blood urea nitrogen level, creatinine level, height-adjusted total kidney volume, height-adjusted total spleen volume, the number of renal cysts, and the number of hemorrhagic renal cysts. However, a multivariate analysis, including sex, height, weight, body mass index, body surface area, blood urea nitrogen level, creatinine level, height-adjusted total kidney volume, height-adjusted total spleen volume, the number of renal cysts, the number of hemorrhagic renal cysts, severe headache, and spinal meningeal diverticula, showed that sex was the only parameter to retain statistical significance (Table), though the number of hemorrhagic cysts was of borderline significance.

#### **Clinical Effects and Progression of Spinal Meningeal Diverticula**

There was more frequent reporting of severe headache in the ADPKD group with spinal meningeal diverticula (68/145 [47%]) compared with those without spinal meningeal diverticula (48/140 [34%], P = .03). Headache prevalence data were not systematically available for the control group. None of the 145 subjects with ADPKD with spinal meningeal diverticula reported any other symptoms attributable to these cysts. The Online Supplemental Data show the number of spinal meningeal diverticula diverticula observed for each of the 285 subjects with ADPKD, showing no relationship with patient age.

There were 50 subjects with ADPKD who had 10+ years of MRI scans, median follow-up = 11.2 years [IQR, 10.6–13.1]). The Online Supplemental Data show that the characteristics of these patients with ADPKD with 10+ years of follow-up are similar to those in the entire ADPKD population. One of these

subjects with ADPKD with follow-up initially had no spinal meningeal diverticula but developed a 5.8-mm spinal meningeal diverticulum. Subjects with ADPKD with follow-up also included 32 subjects with a spinal meningeal diverticulum on the initial MRI. For these 32 affected subjects with ADPKD with follow-up (Online Supplemental Data), the mean number of cysts on their first scan, 3.3 [SD, 2.7], was similar to the mean number of cysts on the most recent scan, 3.2 [SD, 2.6] (P = .4). However, the mean diameter of the largest cyst increased from 8.1 [SD, 3.2] mm on the first scan to 8.9 [SD, 3.7] mm (P = .01) on the most

recent scan (Fig 4). The Online Supplemental Data show the growth of spinal meningeal diverticula from the first scan compared with the most recent scan in all subjects with ADPKD with 10+ years follow-up.

#### DISCUSSION

These data from 285 subjects with ADPKD and matched control patients show that the prevalence of spinal meningeal diverticula was 51%, which was 2.2 times higher in those with ADPKD than the control group. In a multivariable model, women with ADPKD were more likely than men to have spinal meningeal diverticula. Furthermore, spinal meningeal diverticula were larger and more numerous in subjects with ADPKD, and the total spinal meningeal diverticula volume was 60% greater compared with controls.

In 1997, Schievink and Torres<sup>12</sup> reported spinal meningeal diverticula in 3 women with ADPKD being evaluated for headache, visual blurring aggravated by upright positioning, and low CSF pressure. Symptoms in 1 patient resolved after surgical repair of an 8-cm thoracic spinal meningeal diverticulum. Additional reports of spinal CSF leaks related to spinal meningeal diverticula in subjects with ADPKD also showed resolution of headache after surgical repair or an epidural blood patch.14,16 Although Kranz et al,<sup>32</sup> in 2013, reported no difference in the prevalence of spinal meningeal diverticula between patients with spontaneous intracranial hypotension and controls, asymptomatic spinal meningeal diverticula have also been reported in patients with ADPKD, all of whom were female.<sup>15,18</sup> Our data confirm the findings of Aşik et al,<sup>17</sup> who reported more and larger cysts in 50 subjects with ADPKD compared with 37 controls without ADPKD. In that study, the number and size of spinal meningeal diverticula correlated with a headache severity score. However, the ADPKD population in that study was not randomly selected, the prevalence of spinal meningeal diverticula was not reported, and the authors did not report a female predominance.<sup>17</sup>

Our data in control subjects without ADPKD, which showed that spinal meningeal diverticula were twice as prevalent in women compared with men, is in agreement with Burdan et al,<sup>33</sup> who evaluated spinal perineural cysts among Eastern European patients, and Langdown et al,<sup>34</sup> who reported that these cysts were seen in about 70% of women. Female prevalence was also



FIG 4. Coronal T2 single-shot FSE images spanning 10 years in a 44-year-old woman with ADPKD showing a right sacral spinal meningeal diverticulum (*red arrows*), which increased in short-axis diameter from 11 to 16 mm.

observed in a study of neurosurgical patients in which 102 (84%) of 122 patients who underwent percutaneous sealing of perineurial cysts were women.<sup>35</sup>

Spinal meningeal diverticula, also known as nerve root cysts or perineural cysts, are hypothesized to arise at sites of dural weakness and are more prevalent in connective tissue disorders, suggesting meningeal connective tissue weakness as a risk factor.<sup>19,20</sup> Other proposed mechanisms include nerve root inflammation, followed by fluid accumulation, arachnoidal proliferation, and communications between dilated sheaths and the subarachnoid space expanding under the influence of pulsatile hydrodynamic CSF forces.<sup>21</sup> The association of spinal meningeal diverticula with ADPKD supports the concept that ADPKD is a more generalized connective tissue disorder. The observation of more and larger sacral spinal meningeal diverticula, compared with the lumbar and thoracic regions, suggests that the increased hydrostatic pressure in the sacral region while the patient is upright may promote cyst formation. This concept is also supported by reports of spinal meningeal diverticula occurring with a type 2 CSF leak (ruptured spinal meningeal diverticulum) or type 3 CSF leak (an aberrant connection between the subarachnoid space and an adjacent paraspinal vein), often presenting as postural headache indicating spontaneous intracranial hypotension.<sup>28-30</sup>

The strengths of this study are the large, unselected population of subjects with ADPKD in the Rogosin PKD Repository with an age- and sex-matched control group and multiobserver analysis blinded to the patient data. We further strengthened the study by excluding all subjects with known connective tissue disorders, debilitating spinal deformities, and incomplete MRI sequences that may have confounded the analysis.

Limitations of this study include retrospective image analysis and failure of MRI sequences to cover the entire thoracic or cervical spine. This feature may have biased the study toward underreporting the true spinal meningeal diverticula prevalence. True blinding was also not possible due to ADPKD status being apparent on the images. None of the patients in this study underwent a procedure to measure CSF pressure or for cyst drainage or excision, which precluded histopathologic evaluation.

#### CONCLUSIONS

These findings indicate that spinal meningeal diverticula are prevalent in subjects with ADPKD, particularly in women, and should be included in the differential diagnosis of headache in ADPKD, especially when the headache is exacerbated or provoked by upright posture.

#### **ACKNOWLEDGMENTS**

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Disclosure forms provided by the authors are available with the full text and PDF of this article at www.ajnr.org.

#### REFERENCES

- Perumareddi P, Trelka DP. Autosomal dominant polycystic kidney disease. Prim Care 2020;47:673–89 CrossRef Medline
- Liu J, Fujikura K, Dev H, et al. Pericardial effusion on MRI in autosomal dominant polycystic kidney disease. J Clin Med 2022;11:1127 CrossRef Medline
- Liu J, Yin X, Dev H, et al. Pleural effusions on MRI in autosomal dominant polycystic kidney disease. J Clin Med 2023;12:386 CrossRef Medline
- Thimmappa ND, Blumenfeld JD, Cerilles MA, et al. Cisterna chyli in autosomal dominant polycystic kidney disease. J Magn Reson Imaging 2015;41:142–48 CrossRef Medline
- Reig B, Blumenfeld J, Donahue S, et al. Seminal megavesicle in autosomal dominant polycystic kidney disease. *Clin Imaging* 2015; 39:289–92 CrossRef Medline
- Zhang W, Stephens CJ, Blumenfeld JD, et al. Relationship of seminal megavesicles, prostate median cysts, and genotype in autosomal dominant polycystic kidney disease. J Magn Reson Imaging 2019; 49:894–903 CrossRef Medline
- Cornec-Le Gall E, Alam A, Perrone RD. Autosomal dominant polycystic kidney disease. *Lancet* 2019;393:919–35CrossRef Medline
- Allen A, Wiegmann TB, MacDougall ML. Arachnoid cyst in a patient with autosomal-dominant polycystic kidney disease. Am J Kidney Dis 1986;8:128–30 CrossRef Medline
- Howe G, Liddell J, Hunn A. Adult polycystic kidney disease and arachnoid cyst formation: Case report. J Clin Neurosci 1995;2:269– 70 CrossRef Medline

- Schievink WI, Huston J 3rd, Torres VE, et al. Intracranial cysts in autosomal dominant polycystic kidney disease. J Neurosurg 1995; 83:1004–07 CrossRef Medline
- 11. Krauer F, Ahmadli U, Kollias S, et al. Growth of arachnoid cysts in patients with autosomal dominant polycystic kidney disease: serial imaging and clinical relevance. *Clin Kidney J* 2012;5:405–11 CrossRef Medline
- Schievink WI, Torres VE. Spinal meningeal diverticula in autosomal dominant polycystic kidney disease. *Lancet* 1997;349:1223–24 CrossRef Medline
- Coche E, Persu A, Cosnard G, et al. Multiple thoracic paraspinal meningeal cysts in autosomal dominant polycystic kidney disease. *Am J Kidney Dis* 2003;41:E8 CrossRef Medline
- 14. Albayram S, Baş A, Ozer H, et al. Spontaneous intracranial hypotension syndrome in a patient with Marfan syndrome and autosomal dominant polycystic kidney disease. *Headache* 2008;48:632–36 CrossRef Medline
- Peces R, Peces C, Pérez-Dueñas V, et al. Sacral radicular cysts in autosomal dominant polycystic kidney disease. NDT Plus 2009; 2:360–61 CrossRef Medline
- 16. Schievink WI, Palestrant D, Maya MM, et al. Spontaneous spinal cerebrospinal fluid leak as a cause of coma after craniotomy for clipping of an unruptured intracranial aneurysm. *J Neurosurg* 2009;110:521–24 CrossRef Medline
- Aşık M, Tufan F, Akpınar TS, et al. Frequency of nerve root sleeve cysts in autosomal dominant polycystic kidney disease. *Balkan Med* J 2016;33:652–56 CrossRef Medline
- Niemczyk M, Gradzik M, Kulesza A, et al. Spinal meningeal cysts in autosomal dominant polycystic kidney disease. Nephrology (Carlton) 2018;23:95–96 CrossRef Medline
- Isono M, Hori S, Konishi Y, et al. Ehlers-Danlos syndrome associated with multiple spinal meningeal cysts-case report. Neurol Med Chir (Tokyo) 1999;39:380–83 CrossRef Medline
- Doi H, Sakurai S, Ida M, et al. A case of sacral meningeal cyst with Marfan syndrome [in Japanese]. No Shinkei Geka 1999;27:847–50 Medline
- Lucantoni C, Than KD, Wang AC, et al. Tarlov cysts: a controversial lesion of the sacral spine. Neurosurg Focus 2011;31:E14 CrossRef Medline
- 22. Acosta FL Jr, Quinones-Hinojosa A, Schmidt MH, et al. Diagnosis and management of sacral Tarlov cysts: case report and review of the literature. *Neurosurg Focus* 2003;15:E15 CrossRef Medline

- 23. Zhang T, Li Z, Gong W, et al. Percutaneous fibrin glue therapy for meningeal cysts of the sacral spine with or without aspiration of the cerebrospinal fluid. J Neurosurg Spine 2007;7:145–50 CrossRef Medline
- 24. Guo D, Shu K, Chen R, et al. Microsurgical treatment of symptomatic sacral perineurial cysts. Neurosurgery 2007;60:1059–65 CrossRef Medline
- 25. Murphy K, Oaklander AL, Elias G, et al. Treatment of 213 patients with symptomatic Tarlov cysts by CT-guided percutaneous injection of fibrin sealant. AJNR Am J Neuroradiol 2016;37:373–79 CrossRef Medline
- Peh WC, Evans NS. Tarlov cysts: another cause of sacral insufficiency fractures? Clin Radiol 1992;46:329–30 CrossRef Medline
- Rodrigues TP, Rodrigues MA, Suriano IC, et al. Idiopathic intracranial hypertension associated with symptomatic perineural cysts: presentation of 2 cases. World Neurosurg 2018;119:17–19 CrossRef Medline
- Schievink WI, Maya MM, Jean-Pierre S, et al. A classification system of spontaneous spinal CSF leaks. *Neurology* 2016;87:673–79 CrossRef Medline
- 29. Mamlouk MD, Shen PY, Jun P, et al. Spontaneous spinal CSF leaks stratified by age, body mass index, and spinal level. AJNR Am J Neuroradiol 2022;43:1068–72 CrossRef Medline
- Mamlouk MD, Shen PY, Dahlin BC. Modified synamic CT myelography for type 1 and 2 CSF leaks: a procedural approach. *AJNR Am J Neuroradiol* 2023;44:341–46 CrossRef Medline
- Pei Y, Obaji J, Dupuis A, et al. Unified criteria for ultrasonographic diagnosis of ADPKD. J Am Soc Nephrol 2009;20:205–12 CrossRef Medline
- 32. Kranz PG, Stinnett SS, Huang KT, et al. Spinal meningeal diverticula in spontaneous intracranial hypotension: analysis of prevalence and myelographic appearance. *AJNR Am J Neuroradiol* 2013;34: 1284–89 CrossRef Medline
- Burdan F, Mocarska A, Janczarek M, et al. Incidence of spinal perineurial (Tarlov) cysts among East-European patients. *PLoS One* 2013;8:e71514 CrossRef Medline
- Langdown AJ, Grundy JR, Birch NC. The clinical relevance of Tarlov cysts. J Spinal Disord Tech 2005;18:29–33 CrossRef Medline
- Murphy KJ, Nussbaum DA, Schnupp S, et al. Tarlov cysts: an overlooked clinical problem. Semin Musculoskelet Radiol 2011;15:163–67 CrossRef Medline