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Prevalence of Cardiovascular Events in Patients with Autosomal Dominant Polycystic Kidney Disease

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Abstract

Background—This study evaluates the prevalence of cardiovascular events in autosomal dominant polycystic kidney disease (ADPKD) patients.

Methods—We distributed surveys to 1439 subjects from our ADPKD research database. 426 subjects with ADPKD completed and returned surveys. Seven of 426 surveys returned were children and were excluded from the study.

Results—ADPKD patients responding were female (63.2%), non-Hispanic (88.1%) and white (93.6%). The mean age of total group was 53.2 ± 13.7 years. 82.8% had a family history of ADPKD and 32.5% had reached end-stage renal disease (ESRD). With respect to cardiovascular risk factors 86.6% were hypertensive with a mean age at diagnosis of 36.9 ± 12.9 years and hypertension was significantly more prevalent in males. In addition, 19.6% of subjects were obese, 20.8% were smokers, 8.7% had diabetes, 45.7% had high cholesterol and 17.8% were sedentary. The most prevalent self reported cardiovascular events were arrhythmias (25.9%), evidence of peripheral vascular disease (16.5%), heart valve problems (14.4%), cardiac enlargement (9.5%), stroke or cerebral bleeding (7.5%), myocardial infarction (6%) and brain aneurysm (5.0%). The most commonly used antihypertensive medications were renin-angiotensin inhibitors used by 75% of ADPKD patients. Older ADPKD patients and those at ESRD had a significantly higher incidence of cardiovascular events.

Conclusion—These findings support the high prevalence of cardiovascular risk factors and events in ADPKD patients and thus increasing risk for mortality. Due to the prevalence of cardiovascular risk factors in the ADPKD population, early diagnosis and clinical intervention are recommended.

Keywords

Autosomal dominant polycystic kidney disease; cardiovascular events; risk factor; morbidity; mortality

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Disclosures None

Introduction

Approximately 6 million Americans have combined chronic cardiovascular and kidney disease resulting in an increasing epidemic of heart and kidney failure [1]. This morbid association represents unique challenges to the clinician. Approximately 600,000 Americans are affected with autosomal dominant polycystic kidney disease (ADPKD), with over 2000 patients starting dialysis every year [2]. Patients with ADPKD have an increased incidence of early onset hypertension, left ventricular hypertrophy (LVH) and cardiovascular abnormalities [3, 4]. The reported relative mortality rate in patients with ADPKD ranges between 1.6 fold (95% confidence interval, 1.3 to2.0) and 3.2 folds higher (95% confidence interval, 2 to 4.8) in comparison with the general population [5].

Cardiovascular complications are the most common cause of morbidity and mortality in patients with ADPKD [6]. Primary prevention therefore is important to reduce early morbidity and mortality. Thus there is a need for detection and treatment of cardiovascular risk factors in the ADPKD population. There is evidence that blockade of the renin-angiotensin-aldosterone system (RAAS) with better control of blood pressure has improved ADPKD patient and renal survival [7–9]. There also are results in hypertensive ADPKD patients which demonstrate that initial therapy with RAAS inhibition as compared to diuretics necessitates significantly few antihypertensive medications for comparable control of blood pressure [10].

The present study analyzed the cardiovascular events and risk factors in a large number of ADPKD patients according to gender, age, hypertension, cholesterol, smoking and end-stage renal disease (ESRD). This observational study was undertaken in an era in which the majority of patients were receiving RAAS inhibition.

Methods

Data source and study population

We developed a 6-page survey that was distributed to 1439 study subjects listed as having ADPKD in our database. The survey asked basic demographic questions and specific questions related to occurrence of cardiovascular disease in ADPKD patients, including occurrence of stroke, peripheral arterial disease, abdominal aortic aneurysm, angina pectoris, myocardial infarction, atrial or ventricular arrhythmias, left ventricular hypertrophy, and cardiac valvular abnormalities. The survey also collected information regarding the presence and treatment of cardiovascular risk factors, including hyperlipidemia, smoking, diabetes mellitus, hypertension, and medication use.

The survey was sent in a single mailing (January 2011) with instructions to return the survey using a provided return envelope. 426 subjects with ADPKD (30%) returned the survey completed. Out of 426 surveys returned, 7 were from patients under the age of 18 and were excluded from the analysis.

Statistical analysis

SAS version 9.3 PROC FREQ and PROC MEANS were used to obtain descriptive statistics for the surveys. The difference between the distribution of age categories for men and women was tested using a contingency table Chi-square test. For this test p < 0.05 was considered significant.

Proportions for demographics were calculated as a percentage of all respondents. Proportions for other tables were calculated as a percentage of those who responded to that question.

Because multiple outcomes were tested, p-values were adjusted using the Bonferroni method. Adjusted p-values less than 0.0036 (0.05 / 14) or unadjusted p-values of less than 0.05 were considered significant. This adjustment corrects for the probability of getting a significant p-value purely by chance.

Results

Descriptive analysis of ADPKD patients responding

ADPKD patients responding were female (63.2%), non-Hispanic (88.1%) and white (93.6%) (Table 1). The mean age of the total group was 53.2 ± 13.7 years. 82.8% had a family history of ADPKD and 32.5% had reached ESRD. Among respondents analysis of cardiovascular risk factors (Table 2) demonstrated that 86.6% had hypertension with mean age of diagnosis of 36.9 ± 12.9 years with significantly higher prevalence in males, 19.6% were obese, 20.8% were smokers, 8.7% had diabetes, 45.7% had high cholesterol and 17.8% were sedentary. The most prevalent self reported cardiovascular events (Table 3) were arrhythmias (25.9%) with mean age of diagnosis of 43.3 ± 16.4 years, evidence of peripheral vascular disease (16.5%) with mean age of diagnosis of 45 ± 13 years, heart valve problems (14.4%) with mean age of diagnosis of 41.2 ± 16.5 years, cardiac enlargement (9.5%) with mean age of diagnosis of 42.6 ± 13.9 years, stroke or cerebral bleeding (7.5%) with mean age of diagnosis of 50.8 ± 13.4 years, myocardial infarction (6%) with mean age of diagnosis of 53.4 ± 9.6 years and brain aneurysm (5.0%) with mean age of diagnosis of 43.4 \pm 13.7 years. Angiotensin converting enzyme inhibitors (ACEIs) and angiotensin receptor blockers (ARBs) were used in 75% of hypertensive ADPKD patients (Table 4). Statins and anti-platelet medications (aspirin) were used in 11 % and 22.5 % respectively.

Sub-groups analysis

Demographic parameters or cardiovascular risk factors were not significantly different between males and females (Table 5). The occurrence of reported heart attacks was significantly higher in males (11.4%) compared to females (3.1%) (Adjusted p-value of 0.0136) (Table 4).

ADPKD respondents over the age of 45 were significantly more likely to report hypertension and high cholesterol than those 45 or under (Table 6). Cardiovascular events were higher in older ADPKD respondents but did not reach significance (Table 6). ADPKD respondents with ESRD were significantly more likely to report diabetes, hypertension, and high cholesterol levels (Table 7). They also reported significantly higher incidence of stroke or cerebral bleeding, heart attack, and cardiac enlargement (Table 7).

Discussion

The most common extra-renal complications that contribute to morbidity and mortality in ADPKD patients are of cardiovascular nature [4]. Hypertension is the most frequent cardiovascular complication in ADPKD and contributes to both an increased incidence of cardiovascular mortality and faster progression to ESRD [6, 11, 12]. Hypertension develops early in the course of ADPKD [13] and occurs in 50 – 70 % of ADPKD patients with normal kidney function [14,15]. Previously we have reported a median age at diagnosis of hypertension in ADPKD of 32 years in males and 34 years in females [16]. The current results support the presence of early hypertension in ADPKD. Hypertension is a widespread feature of this disease and has been reported in up to 80% of ADPKD patients with ESRD on dialysis [17]. Thus, the main and most effective therapy in ADPKD remains control of hypertension primarily by including RAAS inhibition [7, 8]. The definitive answer of whether treatment with either ACEIs or / and ARBs results in decreased rate of renal disease progression in ADPKD awaits the results of the HALT-PKD study [18]. However, it is important to control hypertension in ADPKD patients since it is a specific risk factor for intra-cerebral hemorrhage and aneurysm ruptures [19].

Our study demonstrates a high prevalence of cardiovascular risk factors including hypertension, obesity, diabetes and hypercholesterolemia in ADPKD population. In a previous study 22% of ADPKD patients (age 35.9 ± 11.1 years) with normal kidney function also fulfilled the International Diabetes Federation criteria of metabolic syndrome [20].

LVH is a significant risk factor for cardiovascular morbidity and mortality and a common finding in hypertensive and even normotensive ADPKD patients [21–24]. However, a recent study in ADPKD patients with preserved renal function reported a prevalence of LVH of 3.9 % (25). Increased left ventricular mass index does occur in children and young adults with ADPKD [13, 26–28]. The early onset of hypertension in ADPKD may be associated with LVH in nearly 50% of ADPKD patients by their 40s [22]Increased LVMI has been found to be associated with poor renal and overall outcomes in ADPKD patients [12]. A significant correlation between hypertension and increased LVMI has been demonstrated in both children and adults with ADPKD [13, 26–28]. RAAS inhibition in hypertensive ADPKD patients has led to long-term reversal of LVH [29,30]. This finding was significantly greater in association with rigorous control of blood pressure (< 120/80 mmHg) in ADPKD patients [30].

Structural cardiac abnormalities are found more often in ADPKD patients than in non-ADPKD family members or normal controls [31]. A prospective echocardiographic study in ADPKD subjects found mitral valve prolapse in 26% and mitral regurgitation in 31% [27]. Tricuspid regurgitation and aortic regurgitation also were found in 15% and 8%,

respectively (29). In the current ADPKD study overall heart valve problems were found in 14.4% of patients.

The occurrence rate of coronary events, such as angina, myocardial infarction, and need for coronary revascularization in ADPKD patients with normal renal function has not been previously reported in the literature. Our survey reported that 3.3% of respondents had angina, 6% had suffered a heart attack and 5.9 % had undergone angioplasty, angioplasty and stent or cardiac valve surgery. The mean age of heart surgery was 50.7 ± 11.9 years. ADPKD patients with ESRD had less coronary events than matched ESRD patients of other causes [32, 33]. This has been attributed to less severe anemia in ADPKD patients [32, 33]. This is probably due to increased endogenous erythropoietin production in ADPKD patients [34].

Arterial aneurysms, particularly intracranial aneurysms, are more prevalent in ADPKD patients than in the general population (4.0–11.7% versus 1.0%) [35, 36]. Moreover, it has been suggested that ADPKD is a risk factor for coronary artery aneurysms [37]. Abdominal aortic aneurysm also appears to be more prevalent in ADPKD patients [38, 39]. The incidence of abdominal aortic aneurysm in our cohort was very low (0.8%). However, a tendency towards larger aortic diameters in ADPKD patients compared to a control population has previously been reported [39].

The other major vascular abnormality in ADPKD is intracranial aneurysms (ICA). The prevalence ranges from 5% in patients with no family history of ICA to 21% in those with a positive family history of ICA rupture [32, 35, 41]. The prevalence may be even higher in ADPKD patients on dialysis, as observed in our study. An occurrence rate of both asymptomatic and ruptured ICA of 33.3 % has been reported in ADPKD patients with ESRD [42]. Another study [43] found no difference in incidence of cerebrovascular accidents (CVAs) between ADPKD patients on dialysis and a non-PKD dialysis patient population. Only 25-50% of CVAs in ADPKD patients have been reported to result from ICA rupture [6, 44]. In our cohort the prevalence of brain aneurysms and stroke or intracerebral bleeding were respectively 5% and 7.5%. ICA rupture accounts for a 35-55% risk of combined morbidity and mortality [19, 45], thus, identification and screening of patients at risk for developing symptomatic ICA are recommended. Systematic screening of ICA with 3-dimensional magnetic resonance angiography (MRA) is recommended for ADPKD patients, particularly for adult patients (30 years), with a positive family history of hemorrhagic stroke or ICA, patients undergoing major surgery with potential hemodynamic instability and those at high risk occupations [46, 47]. MRA has been recommended every 5 years if initially negative and every 2-3 years if positive [46]. However, recent data support less requierement for screening for ICAs in ADPKD patients, and therefore widespread screening is not indicated [48].

Patients with non-PKD chronic kidney disease are at significantly increased cardiovascular events and risk factors [49]. However, ADPKD is unique by early occurrence of hypertension, heart valve problems and ICA. As expected, older ADPKD patients and those with ESRD are at higher risk for cardiovascular events. However, male gender may be losing its importance as a risk factor for cardiovascular events in ADPKD. The early and

effective treatment of hypertension in ADPKD is critical in the prevention of cardiovascular events in ADPKD.

Conclusion

There are intrinsic limitations to the survey based nature of this study and the reported frequencies may be underestimated. Nevertheless, these present findings confirm the high prevalence of cardiovascular risk factors and events in ADPKD patients which are associated with increased risk for mortality. Moreover, older ADPKD subjects and those with ESRD had an increased risk for cardiovascular events, and this increased morbidity and mortality. Due to the prevalence and early onset of cardiovascular risk factors in the ADPKD population, early diagnosis and intervention by aggressively treating blood pressure in ADPKD patients is important to prevent LVH, cardiovascular complications, and mortality.

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Appendix. Survey on Polycystic Kidney Disease (PKD)

confidential. Feel free to attach a sheet	of paper if needed to clarify an answer.
	Date:
1.Name:	
(Last) (Fi (Maiden)	rst) (Middle)
2. Date of birth: DD DD DDDD	
Month Day	Year
3. Gender: Male Female	5
4. Ethnic categories (select one):	
Hispanic or Latino	Unknown
Not Hispanic or Latino	Decline to report
Decline to report	
5. Racial categories (select one):	
American Indian/Alaska Nat	ive 🗆 White
🗆 Asian	More Than One Race
Native Hawaiian or other F	acific Islander 🗆 Unknown
 Black or African America 	Decline to report
6. Do you have other family mem	pers with PKD?
7. Have you had either hemodialy	sis or peritoneal dialysis treatment? 🛛 Yes 👘
If yes, what was your age at t	he time you had your first dialysis treatment? 🗆
8. Have you received a kidney tra	nsplant? 🗆 Yes 🗆 No
If yes, what was your age at t	he time of your first transplant?
	ht (feet)
10. Have you ever smoked Cigare	ttes or Cigars: Yes No
	□ Number of years you have smoked: □□
Do you still smoke? Yes	
11. Have you ever been told by a c	loctor that you have diabetes?
□ Yes □ No If yes	, what was your age at diagnosis? 🗆
If yes, Type: 🗆 1 🗆 2	🗆 Gestational 🛛 🗆 Unknown
	(only during pregnancy)
If yes, initial treatment: D	iet
If yes for medication(s), which	ch diabetes medication(s) do you take?

Current diabetes treatment (if different from above):
Diet Diet Medication Insulin injection
What was your age when you began your current treatment?

If yes for medication(s), which diabetes medication(s) do you currently take?

12. Have you ever been told by a doctor that you have high blood pressure

Yes
 No
 If yes, what was your age at diagnosis?
 Have you been on medication for high blood pressure ?
 Yes
 No
 If yes, which high blood pressure medication(s) do you take?

13. Have you ever been told by a doctor that you have high cholesterol?

🗆 Yes 🗆 N	lo If yes,	what was your age at diagnosis?
Current treatment	nt: 🗆 Diet	Medication

If yes for medication, which high cholesterol medication(s) do you take?

	Do you exer	rcise? 🗆 Y	es 🗆 No	If yes, how man	y hours/week?	סכ
	What type o	f exercise?	D Walking	Running	🗆 Biking	Other
15.	Have you e	ver been to	ld by a doctor t	hat you had a s	troke or bleedi	ng in your
	head?					
	🗆 Yes	🗆 No	If yes, what	was your age at	diagnosis? 🗆	
	What wa	s the proble	m (Mark all that	apply)?		
	🗆 Symp	toms of stro	ke lasting less t	han 24 hours	Stroke	
	Bleed	ling	Rupture	d Aneurysm		
16.	Have you e	ver been to	ld by a doctor t	hat you have a	brain aneurysn	n?
	🗆 Yes	🗆 No	If yes, what w	as your age at d	iagnosis? 🗆	
	If	yes, have y	ou ever had surg	gery for your and	eurysm? 🗆 Yes	🗆 No
17	Has anyona	in your fa	mily been told l	by a doctor that	they have a hr	ain anour
17.			urysm in their	•	they have a br	am aneur
	□ Yes		Unknown	nead :		
				2		
			r age at diagnos	is? 🗆		
	How are	they related	to you?			
	If yes, w	hat was thei	r age at diagnosi	is? 🗆		
	How are	they related	to you?			
	If yes, w	hat was thei	r age at diagnosi	is? 🗆		
	How are	they related	to you?			
			ulation proble	ms in your legs	2	
18	Have you e	ver had cire				
18.	Have you e				n this first occur	red? nn
18.	Have you e			as your age whe	n this first occur	red? 🗆
	🗆 Yes	🗆 No	If yes, what w			
	🗆 Yes	🗆 No	If yes, what w	as your age whe	abdominal aort	
19.	YesHave you etYes	□ No ver been to □ No	If yes, what w Id by a doctor t If yes, what	as your age whe hat you had an was your age at	abdominal aort diagnosis? 🗆	
19.	YesHave you etYes	 No ver been to No ver been to 	If yes, what w Id by a doctor t If yes, what Id by a doctor t	as your age whe hat you had an	abdominal aort diagnosis? == ngina?	
19. 20.	 Yes Have you et Yes Have you et Yes 	□ No ver been to □ No ver been to □ No	If yes, what we do by a doctor t If yes, what d by a doctor t If yes, what we	as your age when hat you had an was your age at hat you have A	abdominal aord diagnosis? == ngina? ugnosis? ==	
19. 20.	 Yes Have you et Yes Have you et Yes 	 No ver been to No ver been to No 	If yes, what we do by a doctor t If yes, what do by a doctor t If yes, what we do by a doctor t	as your age when hat you had an was your age at hat you have A as your age at dia	abdominal aord diagnosis? == ngina? ngnosis? == eart attack?	
19. 20. 21.	 Yes Have you et Yes Have you et Yes Have you et Yes Have you et 	 No ver been to No ver been to ver been to No 	If yes, what we id by a doctor t If yes, what id by a doctor t If yes, what we id by a doctor t If yes, what we	as your age whe hat you had an was your age at hat you have A is your age at dia hat you had a h	abdominal aori diagnosis? == ngina? nginasis? == eart attack? ngnosis? ==	tic aneury
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19.20.21.22.	 Yes Have you e Yes Have you e Yes Have you e Yes Have you e Yes 	 No ver been to No ver been to No ver been to ver been to 	If yes, what we do by a doctor t If yes, what If yes, what we do by a doctor t If yes, what we do by a doctor t If yes, what we	as your age whe hat you had an was your age at hat you have A is your age at dia hat you had a h is your age at dia hat you have ar	abdominal aord diagnosis? == ngina? eart attack? ngnosis? == h irregular hear iagnosis? ==	t beat?

24. Have you ever been told by a doctor that you have a heart valve problem?

D Yes No If yes, what was your age at diagnosis?

- 25. Have you ever had heart surgery? Yes 🗆 No If yes:
 Angioplasty
 Stents
 Angioplasty+Stent
 Coronary bypass If yes, what was your age at surgery?
- 26. If working, how many days have you missed work in the last year due to symptoms related to your PKD?
- 27. Have you ever been told by a doctor that you have osteoporosis?
 - □ Yes □ No If yes, what was your age at diagnosis? □□
- 28. Have you ever been told by a doctor that you have low vitamin D?

□ Yes □ No If yes, what was your age at diagnosis? □□

- 29. Have you ever been told by a doctor that you had kidney stones, or other problems from kidney stones?

 - 🗆 Yes 🗆 No If yes, what was your age at diagnosis? □□
 - □ In one kidney □ In both kidneys

How were the kidney stones treated?

Surgery

Medication D Other

If yes for medication, which medication(s) did you take for your kidney stones?

30. What were your symptoms of kidney stones (mark all that apply):

- Back Pain Flank/Side Pain
 Abdominal/Stomach Pain
- Bloody Urine
 A doctor found blood in my urine
- Kidney infection
 Abnormal laboratory or radiology results

31. Do you know what kind of stone you had?

□ Uric acid stone □ Calcium containing stone □ Unknown

32. Any other Medical Conditions not mentioned above?

33. Are you currently taking prescription or over the counter medication(s) or vitamins?

□ Yes □ No, not currently taking medication □ Decline to report

Please list all current medicine(s) and the first date on which you started taking this medicine:

Date Began Taking

Name of Medication Name of Medication

Medication

Medication

Date Began Taking

Table 1

Demographic characteristics of 419 survey respondents with ADPKD

Variable	Ν	% of Total (419)
Sex		
Male	145	34.6%
Female	265	63.2%
Not reported	9	2.2%
Ethnicity		
Hispanic or Latino	18	4.3%
Not Hispanic or Latino	369	88.1%
Unknown or not reported	32	7.6%
Race		
American Indian / Alaska Native	3	0.7%
Asian	3	0.7%
Black or African American	3	0.7%
White	381	93.6%
More than one race	12	3.0%
Unknown or not reported	17	4.1%
Family history of ADPKD	347	82.8%
No	37	8.8%
Unknown	35	8.4%
ESRD	136	32.5%
No	271	64.7%
Unknown	12	2.9%
Hemodialysis or peritoneal dialysis	83	19.8%
No	329	78.5%
Unknown	7	1.7%
Transplantation	117	27.9%
No	291	69.5%
Unknown	11	2.6%
Dialysis and transplantation	66	16%
Preemptive transplantation (never on dialysis)	51	12.2%
		$Mean \pm SD$
Age at time of survey	418	53.2 ± 13.7

ADPKD, autosomal dominant polycystic kidney disease; ESRD, End-stage renal disease.

Table 2

Incidence of cardiovascular risk factors in 419 survey respondents with ADPKD

Variable	Ν	Percent
Obesity (BMI 30)	77 / 392	19.6%
Ever smoked	156 / 412	37.9%
Current Smoker	32 / 154	20.8%
Diabetes	36 / 412	8.7%
Hypertension	356 / 411	86.6%
High cholesterol	188 / 411	45.7%
Sedentary	73 / 411	17.8%

ADPKD, autosomal dominant polycystic kidney disease; BMI, Body mass index

Table 3

Prevalence of cardiovascular events 419 survey respondents with ADPKD

Variable	N	%	Age at Diagnosis Mean ± SD
Stroke or cerebral bleeding	31 / 412	7.5%	50.8 ± 13.4
Brain aneurysm	20 / 397	5.0%	43.4 ± 13.7
Circulation problems in legs	66 / 400	16.5%	45 ± 13
Abdominal aortic aneurysm	3 / 397	0.8%	35.7 ± 26.8
Angina	13 / 399	3.3%	48.9 ± 15.9
Heart attack	24 / 399	6%	53.4 ± 9.6
Irregular heart beat (arrhythmia)	103 / 398	25.9%	43.3 ± 16.4
Enlarged heart	38 / 400	9.5%	42.6 ± 13.9
Heart valve problem	57 / 397	14.4%	41.2 ± 16.5
			5
Heart surgery	23 / 393	5.9%	50.7 ± 11.9
Angioplasty	4 / 23	17.4%	
Stents	8 / 23	34.8%	
Angioplasty + Stents	3 / 23	13%	
Coronary bypass	4 /23	17.4%	
Cardiac valve surgery	4/23	17.4%	

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Helal et al.

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	Hypertensive Al	DPKD Patients	Non hypertensive	Hypertensive ADPKD Patients Non hypertensive ADPKD Patients All ADPKD Patients	All ADPKD	Patients
Drug	Z	%	Z	%	Z	%
Diuretics	78 /333	23.4%	1 / 53	1.9%	79 / 387	20.4%
Sympathetic blocking agents	92 / 329	28%	2 / 54	3.7%	94 / 385	24.4%
Vasodilators	14/331	4.2%	1 / 53	1.9%	15/387	3.9%
Ca Channel blockers	87 / 330	26.4%	0 / 54	%0	87 / 386	22.5%
Angiotensin converting enzyme inhibitors	155/331	46.8%	6 / 54	11.1%	161 / 387	41.6%
Angiotensin receptor blockers	92 / 331	27.8%	2/51	3.9%	94 / 384	24.5%

Table 5

Analysis of cardiovascular risk factors and events by gender among 419 survey respondents with ADPKD

Variable	Male	Ie	Female	alte	p-value ¹	Adjusted p-value
Obesity (BMI 30)	33/140	23.6%	44/248	17.7%	0.1668	1
Current Smoker	09/6	15.0%	22/91	24.2%	0.1719	1
Diabetes	12/144	8.3%	23/264	8.7%	0.8961	1
Hypertension	127/143	88.8%	225/264	85.2%	0.3127	1
High cholesterol	68/143	47.6%	119/264	45.1%	0.6322	1
Stroke or cerebral bleeding	10/144	6.9%	20/144	7.6%	0.8154^{I}	1
Brain aneurysm	5/138	3.6%	14/255	5.5%	0.4101^{I}	1
Abdominal aortic aneurysm	2/138	1.5%	1/255	0.4%	0.2808^{2}	1
Angina	8/140	5.7%	5/255	2.0%	0.0726 ²	-
Heart attack	16/140	11.4%	8/255	3.1%	0.0010^{I}	0.0136
Irregular heart beat (arrhythmia)	32/139	23.0%	71/255	27.8%	0.2980^{I}	-
Enlarged heart	21/140	15.0%	17/256	6.6%	0.0069 ¹	0.0971
Heart valve problem	17/138	12.3%	40/255	15.7%	0.3655 ¹	-

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ADPKD, autosomal dominant polycystic kidney disease; BMI, Body mass index;

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Table 6

Analysis of cardiovascular risk factors and events by age (more or less 45 years) among 419 survey respondents with ADPKD

	42 Y	45 years	>45 years	ears	p-value ^I	Adjusted p-value
Obesity (BMI 30) 1	17/86	19.8%	60/305	19.7%	0.9843	1
Current Smoker 1	11/28	39.3%	21/125	16.8%	0.0082	0.1146
Diabetes	3/91	3.3%	33/320	10.3%	0.0367	0.5140
Hypertension 5	59/91	64.8%	296/319	92.8%	<.0001	<.0001
High cholesterol	14/91	15.4%	174/319	54.6%	<.0001	<.0001
Stroke or cerebral bleeding	1/92	1.1%	30/319	9.4%	0.0078^{I}	0.1089
Brain aneurysm	2/90	2.2%	18/306	5.9%	0.1634^{I}	Ч
Abdominal aortic aneurysm (0/92	0	3/304	1.0%	n/a	1
Angina	1/92	1.1%	12/306	3.9%	0.3141 ²	-
Heart attack	1/92	1.1%	23/306	7.5%	0.0231^{I}	0.3236
Irregular heart beat (arrhythmia) 1	14/91	15.4%	89/306	29.1%	0.0089 ¹	0.1239
Enlarged heart	2/91	2.2%	36/308	11.7%	0.0067 ^I	0.0943
Heart valve problem	8/91	8.8%	49/305	16.1%	0.0828 ¹	-

ADPKD, autosomal dominant polycystic kidney disease; BMI, Body mass index;

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Table 7

Analysis of cardiovascular risk factors and events by gender among 419 ADPKD survey respondents with and without ESRD.

Helal et al.

	ESRD	7	No ESKU		p-value ^I	Adjusted p-value
Obesity (BMI 30) 2	21/129	16.3%	56/257	21.8%	0.2012	1
Current Smoker	8/57	14.0%	24/94	25.5%	0.0938	1
Diabetes 2	22/135	16.3%	13/270	4.8%	0.0001	0.0015
Hypertension 11	132/135	97.8%	218/269	81.0%	<.0001	<.0001
High cholesterol 8	86/134	64.2%	97/270	35.9%	<.0001	<.0001
Stroke or cerebral bleeding 2	20/135	14.8%	10/271	6.7%	<.0001 ¹	0.0008
Brain aneurysm	12/131	9.2%	8/259	3.1%	0.0102^{I}	0.1434
Abdominal aortic aneurysm	2/131	1.5%	1/259	0.4%	0.2621 ²	Т
Angina	8/132	6.1%	5/260	1.9%	0.0387 ²	0.5417
Heart attack 1	16/133	12.0%	8/259	3.1%	0.0005^{I}	0.0066
Irregular heart beat (arrhythmia) 4	44/133	33.1%	57/258	22.1%	0.0187^{I}	0.2614
Enlarged heart 2	21/132	15.9%	16/261	6.1%	0.0017^{I}	0.0240
Heart valve problem 2	23/130	17.7%	32/260	12.3%	0.1498^{I}	1

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ADPKD, autosomal dominant polycystic kidney disease; BMI, Body mass index;