

## RESEARCH

# A disease-specific Quality of Life questionnaire for primary aldosteronism

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## Abstract

**Objective:** To develop a primary aldosteronism (PA) disease-specific Health-Related Quality of Life (HRQoL) questionnaire.

**Methods:** We included newly diagnosed patients with PA ( $n = 26$ ), and patients with PA after adrenalectomy ( $n = 25$ ) or treated with mineralocorticoid receptor antagonists ( $n = 25$ ). According to the guidelines for developing HRQoL questionnaires from the European Organization for Research and Treatment of Cancer (EORTC): Phase I: systematic literature review followed by focus group meetings with patients ( $n = 13$ ) resulting in a list of 94 HRQoL issues. Relevance of issues was rated by 18 other patients and by health care professionals ( $n = 15$ ), resulting in 30 remaining issues. Phase II: selected issues were converted into questions. Phase III: the provisional questionnaire was pre-tested by a third group of patients ( $n = 45$ ) who also completed the EORTC core Quality of Life questionnaire (QLQ-C30). Psychometric testing resulted in a final selection of questions with their scale structure.

**Results:** After the collection and selection of HRQoL issues a provisional questionnaire consisting of 30 items was formed. Of these items, 26 could be assigned to one of the four scales 'physical and mental fatigue', 'anxiety and stress', 'fluid balance' and 'other complaints' cumulatively accounting for 68% of variation in all items. All scales had good reliability and validity. There was a significant correlation of all four scales with the QLQ-C30 in most cases.

**Conclusions:** We developed the first PA-specific HRQoL questionnaire (PA-QoL) using standard, methodologically proven guidelines. After completion of the final validation (phase IV, international field testing), the questionnaire can be implemented into clinical practice.

## Key Words

- ▶ hyperaldosteronism
- ▶ quality of life
- ▶ patient health questionnaire
- ▶ development

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## Introduction

Primary aldosteronism (PA) is a common cause of hypertension, accounting for 5–15% of hypertensive patients (1, 2, 3). PA is categorized in an aldosterone-producing adenoma (APA) or bilateral adrenal hyperplasia (BAH). When unilateral aldosterone hypersecretion is suspected (APA), patients generally undergo adrenalectomy (ADX).

Patients with BAH receive a mineralocorticoid receptor antagonist (MRA).

Over the last decade, there has been increasing interest in the patient perspective with the assessment of HRQoL, commonly defined by the functional effect of an illness and/or its treatment upon a patient, as perceived by the patient (4).

HRQoL studies in patients with PA demonstrated a reduced HRQoL compared to reference values (5, 6) and improvement in HRQoL after ADX (5) and during MRA-based treatment (6). Furthermore, psychological symptoms impairing HRQoL in patients with PA have been reported: occurrence of anxiety disorders and stress is higher than in the general population (7) and in patients with essential hypertension (8). Also, depression, somatization, psychological distress and a lower level of well-being among patients with PA compared to healthy controls have been demonstrated (8). This can be explained by overstimulation of the abundantly present mineralocorticoid receptors in brain structures that are involved in fear and anxiety (9), as well as the multiple drugs needed to control blood pressure and hypokalemia (and thereby adverse reactions) to which patients affected by PA are frequently exposed to. Additionally, uncontrolled hypertension, especially when it cannot be explained by their physician and diagnostic delay can increase anxiety and stress.

For the assessment of HRQoL, it is usually recommended that a generic HRQoL questionnaire (assessing multiple domains of HRQoL) is combined with a disease-specific questionnaire (assessing HRQoL aspects relevant for a specific disease) (10). In previously conducted studies among patients with PA, HRQoL was assessed with a generic HRQoL questionnaire (for example the Short Form 36 General Health Survey (SF-36)). No disease-specific HRQoL questionnaire for patients with PA is currently available. Therefore, the aim of this study was to develop a PA-specific HRQoL questionnaire in order to be able to assess HRQoL and the effect of treatment in PA more accurately.

## Methods

### Patients and study design

The European Organization for Research and Treatment of Cancer (EORTC) is an independent, non-profit cancer research organization which coordinates and conducts international translational and clinical research to improve the standard of cancer treatment for patients. The EORTC Quality of Life Group is dedicated to the development of HRQoL questionnaires. We decided to develop our questionnaire according to their guidelines for developing HRQoL questionnaires as this is the most widely used and comprehensive guideline in the field (11, 12, 13). The questionnaire will not be part of the EORTC portfolio, since the latter is primarily designed for use

in cancer clinical trials, but the guideline can be used in other research settings and clinical practice as well (11).

Below we describe the different phases of development of a PA-specific HRQoL questionnaire. All patients included in this study were treated at the Radboud University Medical Center, had been diagnosed with PA (aldosterone after intravenous salt loading test  $>280$  pmol/L or 140–280 pmol/L and positive consensus of PA by an expert panel) (14) and had sufficient Dutch language skills (15). All patients have undergone subtyping by adrenal vein sampling (cut-off value for selectivity index  $\geq 3$  and for lateralization index  $\geq 4$ ). The Ethics Committee of the Radboud University Medical Center judged that no detailed review was warranted given the non-intrusive and non-experimental character of this study. We obtained written informed consent of all patients.

### Phase I

This phase was aimed at compiling an exhaustive list of HRQoL issues relevant to patients with PA. For this purpose, three sources were used. First, we performed an extensive systematic literature search identifying all studies investigating HRQoL, related constructs as well as physical and mental symptoms in patients with PA, published previously (16). The review resulted in a list of issues. Second, we organized two patient focus group meetings. We invited 43 patients, of whom 13 were willing to participate: one meeting for untreated patients (no previous ADX and no use of MRA;  $n=5$ ) and one for patients treated by ADX ( $n=4$ ) or with MRA ( $n=4$ ). Treated patients were included in order to explore treatment-related HRQoL issues. Furthermore, these patients could also contribute to the issues they experienced before treatment, by remembering the changes caused by the treatment and comparing pre- and post-treatment period. In these qualitative interviews, we collected all possible relevant issues for HRQoL in a semi-structured way. Additionally we asked the patients to describe their experience and to provide information freely. Finally, they were asked to add missing issues on a form, in view of privacy matters. We recorded these meetings and wrote out these recordings verbatim. From these, two authors (MV and AN) made a list of issues and combined this with the items from the list from the literature review.

We presented this combined list to health care professionals (HCP  $n=15$ ) with expertise in the field of PA from different countries and with different professional background and to another group of 18 patients (untreated  $n=6$ ; ADX  $n=6$ ; MRA  $n=6$ , invited  $n=39$ ). These were all

asked to indicate the relevance on a four-point Likert scale (1 = not at all; 2 = a little bit; 3 = moderately; 4 = extremely) and to select a maximum of 15 most important issues. An issue was selected for inclusion in Phase II on the condition that the mean score for both patients and HCPs was >2 and/or that >30% of the patients stated an issue as important. We also asked them to indicate missing issues, serving as our third source for HRQoL issues.

## Phase II

We converted the final list of HRQoL issues into Dutch questions (items) fitting the following response options 'not at all', 'a little', 'quite a bit' and 'very much'. When available, we used phrases that had been used previously in EORTC HRQoL modules (17).

The Dutch questions were then translated into English by applying an iterative forward-backward procedure (18).

## Phase III

We asked 45 patients not included in phase I (untreated  $n=15$ ; ADX  $n=15$ ; MRA  $n=15$ , invited  $n=54$ ) to fill out the provisional questionnaire together with the EORTC core cancer quality of life questionnaire (QLQ-C30) online in Castor EDC (Ciwit B.V. 2017) or, when preferred, on paper. The QLQ-C30 served as a generic HRQoL questionnaire for validation purposes and is a standard item in this phase according to the EORTC guideline (11). When specific HRQoL items are included in the provisional questionnaire, there should be an adequate correlation with the QLQ-C30. Most questions (1–26) referred to the patient's experience *during the last week*, some (27–30) to the patient's experience *during the last 4 weeks*. Additionally they underwent a debriefing in which they had to indicate whether they felt any questions to be confusing, intrusive, annoying or upsetting. Furthermore, we asked patients to indicate any missing questions. For every item we checked the following criteria for inclusion in the final questionnaire (response categories: 1 = not at all; 2 = a little; 3 = quite a bit; 4 = very much): 1. Mean score >1.5; 2. Prevalence ratio >30% (number of patients reporting score 2, 3 or 4); 3. Range >2 points; 4. Responses of scores in categories 1/2 and 3/4 >10%; 5. No significant concerns expressed by patients; 6. Response >95%. When >2 criteria were not met in the total group ( $n=45$ ) nor in the untreated group ( $n=15$ ), we did not include the item in the final questionnaire. Items with low correlation with the total of other items were excluded for the final factor

analysis (initial communalities <0.1, see data analysis). For the issues added in the last step of phase I (missing issues), patients had to indicate the relevance and importance according to phase I and issues were included in the final questionnaire on corresponding condition (mean score >2 and/or >30% of the patients stated an issue as important).

For a disease-specific HRQoL questionnaire, it is not possible to include healthy or essential hypertension controls for any comparison in this phase, because not all questions are applicable for them.

## Data analysis

We performed all analyses in SPSS version 22.0 for Windows (SPSS Inc.). We used descriptive statistics for phase I and phase III.

We tested the suitability of the data for exploratory factor analysis with the Kaiser–Meyer–Olkin (KMO) method, of which the value must be >0.5 (19). Initial communalities, estimates of the variance in each variable accounted for by all factors, were determined ( $R^2$  between one variable and all others in multiple regression analysis). We used exploratory factor analysis to explore the underlying constructs, explaining significant portions of variance. We based the number of constructs on the Kaiser–Guttman rule (the amount of the total variance explained by that factor must be >1) and the Cattell's scree plot. We analyzed the factor loadings to explain the meaning of each construct. This was done by oblique rotation, which assumes that the factors could be related, simplifying their interpretation. Only items with factor loadings of >0.325 were retained for further analysis.

We performed multi-trait scaling analysis to confirm the scale structure found in exploratory factor analysis. We examined for item-scale convergent validity, corrected for overlap. We tested item-scale discriminant validity by comparing the correlation of each item with its own scale versus the other scales. Convergent and discriminant validity are subtypes of construct validity. Demonstration of both convergent and discriminant validity implicates construct validity, that is the questions are actually measuring what they are supposed to measure.

We tested reliability by measuring Cronbach's alpha coefficient, of which the preferred level is >0.7 (20). Additionally convergent and discriminant validity were examined by calculating Spearman's correlations between the scales and the items and between the scales/items and those of the EORTC QLQ-C30 (21). For the inclusive criterion, we considered an item-scale correlation of >0.40 (corrected for overlap) as adequate.

The exclusive criterion was met on the condition that the correlation with the other scales was lower than the correlation with the corresponding scale. When both criteria were met, we concluded that there was scaling fulfillment (20).

For the summed (scale) scores, all answers of the items contributing to the scale were summed, with a higher score corresponding with a lower HRQoL. Finally, we compared all summed scale scores and the total score (also added up) of the three patient groups with the Kruskal–Wallis *H* test.

## Results

### Phase I

The patient characteristics are listed in Supplementary Table 1 (see section on [supplementary data](#) given at the end of this article). We included 15 studies in our systematic review (16), from which we extracted 66 HRQoL issues. Both focus group meetings resulted in an additional 28 issues. These 94 issues were rated by 18 patients (mean age 53 years) and 15 HCP (ten Dutch and five other European professionals; six internist-vascular specialists, six endocrinologists, two nurses and one surgeon). This resulted in the removal of 65 issues. As more than one patient indicated ‘edema’ as missing issue, we added this to the list.

### Phase II

The resulting 30 issues were converted into questions, resulting in the provisional PA-QoL. Whenever possible,

we selected questions from the EORTC item bank (<http://www.eortc.be/itemlibrary/>), a collection of all available EORTC (HRQoL) questionnaires. The issues for which no corresponding question was available in the EORTC item bank ( $n=11$ ) were converted into questions after consensus by MV, JD and OH, in line with the EORTC style.

### Phase III

All questionnaires were filled out completely. This was done online by 37 patients and on paper by eight patients. Patient characteristics are shown in Table 1. Response information is provided in Table 2. The time for completion of the provisional PA-QoL was  $\leq 10$  min for 24 patients, 11 to 15 min for 15 patients, 16 to 20 min for three patients and 21 to 25 min for three patients. Two patients indicated gynecomastia as a missing item. Therefore, this item was added as a single item only for men on spironolactone.

We excluded item 24 about impaired concentration based on our pre-defined criteria (mean score was  $<1.5$  for the total group as well as for the untreated patients, prevalence ratio  $<30\%$  and responses of scores ‘quite a bit’/‘very much’  $<10\%$ ). We treated the item regarding erectile dysfunction as a single item. The KMO value for the remaining 28 items was 0.78. All initial communalities were  $>0.1$ . Four factors were identified, cumulatively accounting for 68% of variation in all items (respectively 48, 9, 6 and 5%). The oblique rotation of the exploratory factor analysis resulted in factor loadings and correlations which are shown in Table 3. We adopted a cut-off value of 0.325 for factor loadings. In other words, only those

**Table 1** Characteristics of the patient groups involved in phase III.

Characteristics	Untreated ( $n = 15$ )	MRA-based treatment <sup>a</sup> ( $n = 15$ )	Adrenalectomy ( $n = 15$ )
Gender M/F	9/6	13/2	12/3
Age (years) <sup>b</sup>	50.7 $\pm$ 10.4	59.6 $\pm$ 10.3	55.7 $\pm$ 8.3
Subtype APA/BAH	9/5 <sup>c</sup>	0/15	15/0
Years since diagnosis of PA (SLT) <sup>b</sup>	<1	4.6 $\pm$ 1.4	3.8 $\pm$ 0.8
Years since diagnosis of hypertension <sup>b</sup>	11.3 $\pm$ 8.7	17.2 $\pm$ 12.3	15.6 $\pm$ 12.1
Years since (start of) treatment <sup>b</sup>	NA	3.9 $\pm$ 1.8	2.7 $\pm$ 0.4
Hypokalemia yes/no <sup>d</sup>	11/4	2/13	0/15
Systolic BP at first visit <sup>b</sup>	159 $\pm$ 23	162 $\pm$ 22	155 $\pm$ 18
Diastolic BP at first visit <sup>b</sup>	94 $\pm$ 12	92 $\pm$ 9.8	92 $\pm$ 14
Number of antihypertensive drugs at first visit <sup>e</sup>	2 (1–3)	2 (1–4)	2 (1–3)
Renin concentration (SLT at T = 0) (mU/L) <sup>e,f,g</sup>	4.9 (2.0–8.5)	3.4 (3.0–4.6)	5.1 (3.0–9.2)
Aldosterone concentration (SLT at T = 4 h) (nmol/L) <sup>e,f,h</sup>	0.48 (0.33–0.77)	0.38 (0.31–0.47)	0.49 <sup>i</sup> (0.32–1.09)

<sup>a</sup>Spironolacton ( $n = 6$ ), eplerenon ( $n = 8$ ), methyldopa ( $n = 1$ , due to pregnancy wish); <sup>b</sup>mean  $\pm$  SD; <sup>c</sup>unknown in one patient; <sup>d</sup>at time of filling out questionnaire; <sup>e</sup>median (25th–75th percentile); <sup>f</sup>pre-treatment; <sup>g</sup> $n = 11/13/10$  (missings due to renin activity measurements in other patients), if renin was  $<3$  ( $n = 11$ ) it was considered as renin = 3, reference values 6.2–65 mU/L; <sup>h</sup> $>0.28$  nmol/L confirms PA; <sup>i</sup> $n = 14$ .

APA, aldosterone-producing adenoma; BAH, bilateral adrenal hyperplasia; NA, not applicable; PA, primary aldosteronism; SLT, salt loading test.

**Table 2** Item descriptive statistics of 45 patients with (treated) primary aldosteronism (phase III).

Item	Question	Mean				Distribution of responses (n = 45) <sup>a</sup>			
		Total (n = 45)	Untreated (n = 15)	MRA (n = 15)	ADX (n = 15)	1	2	3	4
1	Have you had headaches?	1.38	1.53	1.40	1.20	30	13	2	0
2	Have you had muscle weakness?	1.38	1.40	1.60	1.23	33	8	3	1
3	Have you been physically limited?	1.73	2.00	1.87	1.33	24	13	4	4
4	Have you lacked energy?	1.89	2.20	1.93	1.53	17	19	6	3
5	Have you felt restless or agitated?	1.47	1.80	1.40	1.20	30	10	4	1
6	Have you felt physically exhausted?	1.56	1.93	1.53	1.20	27	14	1	3
7	Have you felt mentally exhausted?	1.47	1.67	1.40	1.33	31	10	1	3
8	Have you had to urinate frequently during the day?	1.69	1.87	1.60	1.60	22	15	8	0
9	Have you had to urinate frequently at night?	2.04	2.07	2.27	1.80	11	23	9	2
10	Have you felt overly thirsty?	1.58	1.93	1.47	1.33	28	8	9	0
11	Have you taken large quantities of fluid?	1.56	1.87	1.33	1.47	28	9	8	0
12	Have you woken up for long periods during the night?	1.71	1.53	1.93	1.67	23	14	6	2
13	Have you, as far as you know, snored?	2.09	2.33	1.87	2.07	15	16	9	5
14	Did you feel sleepy during the day?	1.84	2.20	1.67	1.67	16	22	6	2
15	Have you felt tired (not rested) when you woke up?	1.84	2.20	1.80	1.53	20	14	9	2
16	Have you worried about the consequences of your high blood pressure?	1.64	2.07	1.57	1.40	24	15	4	2
17	Have you been worried about the consequences of the elevated levels of aldosterone?	1.76	2.27	1.67	1.33	21	16	6	2
18	Have you worried about your treatment causing future health problems?	1.71	1.87	1.93	1.33	22	15	7	1
19	Have you been worried about the side effects of your treatment?	1.71	2.00	1.80	1.33	23	14	6	2
20	Have you felt frustrated about the number of pills that you are taking?	1.93	2.20	1.87	1.73	22	10	7	6
21	Have you felt impatient?	1.49	1.67	1.60	1.20	29	11	4	1
22	Did you feel irritable?	1.69	2.00	1.67	1.40	22	16	6	1
23	Did you get angry easily?	1.58	1.73	1.60	1.40	25	15	4	1
24 <sup>b</sup>	Have you had difficulty in concentrating on things, like reading a newspaper or watching television?	1.36	1.40	1.33	1.33	32	11	1	1
25	Have you had difficulty remembering things?	1.58	1.60	1.60	1.53	23	19	2	1
26	Have you had swelling in your legs or ankles?	1.47	1.80	1.40	1.20	31	9	3	2
27	Have you felt frustrated about a possible delay in the diagnosis of primary aldosteronism?	1.49	2.00	1.33	1.13	31	6	8	0
28	Have you felt worried about the diagnosis?	1.53	2.07	1.33	1.20	28	11	5	1
29	Have you had decreased libido?	1.82	1.93	1.80	1.73	25	8	7	5
30	<i>For men:</i> Did you have difficulty obtaining or maintaining an erection?	1.68	1.78	2.08	1.17	22	4	5	3
		n = 34	n = 9	n = 13	n = 12				

Patients were untreated, after adrenalectomy (ADX) or on mineralocorticoid receptor antagonists (MRA).  
<sup>a</sup>1 = not at all; 2 = a little; 3 = quite a bit; 4 = very much; <sup>b</sup>item deleted based on pre-defined criteria (see text).

items scoring higher than this threshold were retained for further analysis. As a result, we treated item 13 (snoring) and item 21 (impatient) also as single items.

The multi-trait analysis showed that scaling assumptions were not met for one item in three of the four scales (Table 4). The reliability of all scales was adequate. There was a significant correlation of all four scales with the EORTC QLQ-C30 in most cases (Table 5). We named the scales physical and mental fatigue, anxiety and stress, fluid balance and other complaints.

The summed scale scores were the highest for the untreated patients and the lowest for the patients after ADX

(Table 6). This was statistically significant for scale 2 and for the total of all 28 items (item concerning erectile dysfunction not included).

### Discussion

We have developed the first disease-specific HRQoL questionnaire for patients with PA, the PA-QoL, consisting of the 30 most relevant items, according to an established guideline (11). This preliminary analysis resulted in four scales, all with a fair reliability and validity. There is an adequate correlation with the validated EORTC





**Table 3** Rotated component matrix of all items with a factor loading >0.325.

Scale	Item	Description	Factor loadings	Correlations
Physical and mental fatigue	2	Muscle weakness	-0.391	-0.645
	3	Physically limited	-0.663	-0.806
	4	Lack energy	-0.531	-0.783
	6	Physically exhausted	-0.331	-0.706
	7	Mentally exhausted	-0.777	-0.882
	12	Woken up long period night	-0.633	-0.668
	14	Sleepy daytime	-0.582	-0.766
	15	Tired wake up	-0.326	-0.641
	25	Problems remember	-0.682	-0.663
	29	Decreased libido	-0.454	-0.602
Anxiety and stress	5	Restless agitated	-0.463	0.623
	16	Worries high blood pressure	-0.663	-0.714
	17	Worries aldosterone	-0.988	-0.947
	18	Worries future health	-0.704	-0.820
	19	Worries side effects	-0.864	-0.893
	20	Frustrated number pills	-0.432	-0.551
	27	Frustrated delay diagnosis	-0.645	-0.733
	28	Worries diagnosis	-0.842	-0.818
Fluid balance	9	Urinate night	0.426	0.566
	10	Thirsty	0.737	0.802
	11	Fluids	0.779	0.789
Other complaints	1	Headaches	0.654	0.724
	8	Urinate frequently day	0.663	0.674
	22	Irritable	0.757	0.893
	23	Angry	0.668	0.779
	26	Swelling legs ankles	0.448	0.510

QLQ-C30 questionnaire. Also, the PA-QoL demonstrates the differences in HRQoL between the different treatment groups, with the lowest QoL in untreated patients and the highest in patients treated by ADX, as established before (22).

This is the first study which assessed all specific and relevant QoL issues in patients with PA, with contributions of 76 patients with PA. The items included in the provisional questionnaire cover a range of difficulties that patients with PA are facing. Some of them can be directly assigned to the pathophysiological effects of high aldosterone, such as edema, disturbances of the fluid balance and muscle weakness (mediated by hypokalemia). The questionnaire

also consists of a number of mental issues, which is in line with earlier findings (16). Sleeping disorders and snoring might be due to the association of PA with obstructive sleep apnea (23). Finally, a decreased libido and erectile dysfunction might be the result of spironolactone, as this mineralocorticoid receptor antagonist has anti-androgen activity (24). All these issues are specific for PA and are not covered by a general HRQoL questionnaire. Remarkably the specific issues concerning depression and generalized anxiety were not included in phase III because relevance and importance was judged low by patients and HCPs, although these problems are known from the literature as associated with PA (16).

**Table 4** Results of multi-trait analysis.

Scale	N	Mean (s.d.)	Cronbach's $\alpha$	Item-scale convergent validity (inclusive criterion)		Item-scale divergent validity (exclusive criterion)		Scaling fulfillment
				Range (mean) of item-scale correlations <sup>a</sup>	Number of item-scale correlations <sup>b</sup>	Range (mean) of correlations with other scales <sup>c</sup>	Number of items higher correlations with other scale <sup>d</sup>	Number of items that meet criterion 1 and 2
1	10	16.8 (6.7)	0.93	0.39–0.79 (0.64)	9/10	0.23–0.76 (0.44)	0/10	9/10
2	8	13.2 (5.5)	0.92	0.51–0.87 (0.69)	8/8	0.29–0.74 (0.42)	1/8	7/8
3	3	5.2 (2.0)	0.77	0.55–0.67 (0.61)	3/3	0.31–0.52 (0.42)	0/3	3/3
4	5	7.8 (2.9)	0.84	0.37–0.81 (0.62)	4/5	0.25–0.77 (0.48)	0/5	4/5

<sup>a</sup>Correlation between items and hypothesized scale; <sup>b</sup>number of item-scale correlation that meet minimum standard for convergent validity ( $\geq 0.40$ );

<sup>c</sup>correlation between items and other scales; <sup>d</sup>number of items that have higher correlation with at least one of the other scales compared to that with the hypothesized scale.

**Table 5** Correlations between EORTC QIQ-C30 and scales/items PA-QoL.

EORTC QIQ-C30	Scales				Single items	
	1	2	3	4	Snoring	Impatience
	Physical and mental fatigue	Anxiety and stress	Fluid balance	Other complaints		
Physical functioning <sup>a</sup>	-0.77**	-0.37*	-0.47**	-0.67**	-0.18	-0.51**
Role functioning <sup>a</sup>	-0.77**	-0.52**	-0.45**	-0.61**	-0.16	-0.43**
Emotional functioning <sup>a</sup>	-0.78**	-0.53**	-0.48**	-0.73**	-0.29	-0.65**
Cognitive functioning <sup>a</sup>	-0.73**	-0.54**	-0.50**	-0.74**	-0.46**	-0.68**
Social functioning <sup>a</sup>	-0.72**	-0.63**	-0.38**	-0.52**	-0.33*	-0.66**
Global QoL/health status	-0.63**	-0.58**	-0.41**	-0.63**	-0.33*	-0.51**
Fatigue	0.87**	0.45**	0.45**	0.75**	0.21	0.50**
Nausea/vomiting	0.54**	0.30*	0.30*	0.39**	0.20	0.26
Pain	0.60**	0.29	0.13	0.36*	0.06	0.30*
Dyspnea	0.63**	0.26	0.27	0.52**	0.06	0.30*
Sleep/insomnia	0.60**	0.52**	0.33*	0.49**	0.32*	0.42**
Appetite loss	0.51**	0.34*	0.43**	0.52**	0.33*	0.58**
Constipation	0.07	-0.05	0.04	0.15	0.26	-0.06
Diarrhea	0.27	0.18	0.20	0.26	0.06	0.24
Financial difficulties	0.28	0.32*	0.30*	0.27	0.27	0.19

<0.40 weak correlation, 0.40–0.60 moderate and >0.60 high.

<sup>a</sup>A high score for these functional scales represents a high/healthy level of functioning; \**P* < 0.05; \*\**P* < 0.01.

This study confirms the impaired HRQoL of patients with PA, and its improvement by treatment. An impaired HRQoL has been reported by various studies before (5, 6, 16, 22, 25), as well as the improvement after therapy (5, 6, 16, 22, 26). In our study, patients after ADX have a better disease-specific HRQoL than patients on MRA-based treatment. This is in line with our earlier findings with a generic HRQoL questionnaire (22). However, in other studies, this relation between therapy for PA and QoL has been questioned (6, 25). This inconsistency might have been the result of a too small number of patients and/or the use of a general/less extensive HRQoL questionnaire. As the PA-QoL is disease-specific, it has presumably more potential to discriminate.

A few limitations should be taken into account. Firstly, the PA-QoL has not yet been tested in a large group of patients from different countries. Therefore, the created scales have a preliminary status. Furthermore, because the scaling is mainly a result of statistical procedures, the grouped items are not always consistent with how we would create scales intuitively. Therefore, the scale structure needs to be confirmed in a large international sample of patients with PA.

Also, one item regarding gynecomastia (*‘Have you had sore or enlarged nipples or breasts?’*) was added to the questionnaire specifically for men using spironolactone as a single item. The relevance will be checked in the final phase. Moreover, when interpreting the outcome of the PA-QoL, we have to take into account the possibility of a selection bias, which might have influenced the scores (in all groups) in phase III, assuming that patients with more complaints are more willing to participate for example. However, we do not think this has influenced the final questionnaire because a lot of issues were eliminated in the following phases due to the elaborated process of constructing this questionnaire. Finally, the gender distribution in phase III might have influenced our results. It has been suggested in previous studies that (untreated) female patients are more affected regarding mental QoL (25), anxiety and depression (7, 8, 27). In our study, more male patients were included, especially in the post-treatment groups. This is a result of a structural higher percentage of men compared to women with PA that are treated in our hospital.

**Table 6** Comparison of treatment groups for outcomes provisional PA HRQoL questionnaire.

Scale	Items (n)	Untreated (n = 15)	MRA-based treatment (n = 15)	Adrenalectomy (n = 15)	P value
1 Physical and mental fatigue	10	18 (14–22)	14 (12–21)	13 (11–17)	0.134
2 Anxiety and stress	8	9 (8–12)	13 (9–14)	9 (8–12)	0.006
3 Fluid balance	3	6 (4–7)	4 (4–6)	4 (3–5)	0.112
4 Other complaints	5	8 (6–12)	7 (5–10)	6 (5–9)	0.192
Total	28	38 (23–43)	42 (36–49)	49 (40–64)	0.027

Data expressed as medians (25th–75th percentile).

MRA, mineralocorticoid receptor antagonists.

The PA-QoL could be used in clinical as well as research setting to measure an important outcome, especially from the patients' perspective. This will allow a more optimal and precise mapping of the impaired HRQoL in patients with PA and make physicians more aware of the potential HRQoL issues that patients with PA are dealing with. Furthermore, it will help them to inform patients adequately and to identify those who most suffer from PA-related health problems, other than hypertension or hypokalemia.

The final phase of the development of the PA-QoL will assess responsiveness to change and cross-cultural validity. The larger number of patients will help to fully validate the scales. We expect that this will also result in a significant difference between treatment groups in all subscales. In our study for two out of three subscales, there was a non-significant trend, probably due to the limited number of patients in each group ( $n=15$ ).

In conclusion, we developed the first PA-specific HRQoL questionnaire using standard, methodologically proven guidelines. This questionnaire can be of value in the care for patients with PA. After final validation in phase IV is completed, it can be implemented into practice.

#### Supplementary data

This is linked to the online version of the paper at <https://doi.org/10.1530/EC-19-0026>.

#### Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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