Depression and anxiety in patients with suspected renal artery stenosis

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Abstract

Objective: To investigate the occurrence of depressive and anxiety disorders in patients remitted for endovascular treatment of renal artery stenosis (RAS), and whether symptoms of depression and anxiety can be reduced after the treatment of RAS with percutaneous transluminal renal angioplasty (PTRA).

Methods Patients: Selected for renal angiography with pressure measurement in the renal arteries due to suspected RAS were invited to participate in the study. 37 patients agreed to answer a questionnaire. Eighteen patients also agreed to a clinical psychiatric examination including the International Neuropsychiatric Interview (M.I.N.I. 6.0) to be diagnosed according to the DSM-IV.

Results: Twenty-one of the 37 patients completing the questionnaire were found to have a pressure gradient of ≥ 10 mmHg at angiography, indicating significant stenosis. Ten of the 18 examined patients met the criteria for current DSM-IV anxiety diagnosis and two patients had a current depression. Three of the 18 examined patients had previously made a suicide attempt.

Conclusion: The results of this study suggest that patients with suspected RAS have a high psychiatric comorbidity, as 56% of the patients undergoing psychiatric examination met the DSM-IV criteria for a current anxiety disorder.

Introduction

Depressive and anxiety disorders are among the most common public health problems in the world. The lifetime prevalence of major depressive disorders is about 16% [1] and the 12-month prevalence 6.6%, while the lifetime prevalence of anxiety disorders is about 29% [2]. The Lundby study reported a lifetime prevalence of anxiety (up to 60 years of age) of 9.9% in men and 19.7% in women [3].

Among the well-known biological mechanisms behind affective disorders are disturbances in the serotonergic system [4], the hypothalamic–pituitary–adrenal (HPA) axis [5], and the immune system [6]. It has also been suggested that the renin–angiotensin–aldosterone system (RAAS) may be affected in some patients with depression [7], and in some with anxiety.

The RAAS prevents low blood pressure (BP) by means of a coordinated effect of the hormones angiotensin and renin. Angiotensin II stimulates the release of aldosterone from the adrenal gland. Aldosterone is a steroid that acts on mineralocorticoid receptors (MR). It reduces the loss of sodium ions and water from the kidneys, which leads to increased extracellular fluid, a greater blood volume and an increase in BP [8]. Constriction of the renal artery (renal artery stenosis, RAS) lowers the intrarenal pressure, thereby stimulating the RAAS, causing persistent hypertension [9,10]. RAS can remain an undetected cause of hypertension for several years, apart from causing deterioration of renal function [11]. RAS leading to severe hypertension or renal dysfunction can be treated with percutaneous transluminal renal angioplasty (PTRA) [12] with stenting. However, indications for treatment are the subject of debate, as the results of large randomized studies have not been convincing [13,14].
Interestingly, the RAAS also affects brain function [15], and there is growing evidence that the RAAS may have a psychopathological influence [7]. MR are found in several regions of the brain, including the hippocampus, the amygdala and the nucleus accumbens [16].

In recent years, several studies have shown that rats treated with aldosterone exhibit symptoms of depression and anxiety [17,18]. Moreover, MR seem to be involved in the response to antidepressant therapy in rats. Studies have shown both up- [19,20] and downregulation [21] of MR after antidepressant therapy.

The findings of some clinical studies support the role of the RAAS in depression. For example, increased levels of aldosterone have been found in patients suffering from depression [22]. Furthermore, in a study by Sonino, et al., six of ten patients with primary aldosteronism (PA) suffered from an anxiety disorder [23]. In a follow-up study, 23 healthy controls were compared to 23 patients with essential hypertension (EH) and 23 with PA [24]. Twelve of the 23 patients with PA suffered from an anxiety disorder, compared to four of the 23 with EH. Patients with PA also scored higher when rated for stress than the EH patients, indicating that not only hypertension, but also the overproduction of aldosterone, causes anxiety.

Little research has been conducted on the potential comorbidity between RAS and psychiatric illness. This study was therefore carried out to investigate the following two hypotheses: the occurrence of depressive and anxiety disorders is higher in patients remitted for endovascular treatment of RAS, and that symptoms of depression or anxiety can be reduced after the treatment of RAS with PTRA.

**Experimental procedures**

**Patients**

The study was approved by the Lund University Medical Ethics Committee (No. 2013/310). All the participants gave their written informed consent to participate.

The patients were recruited from the Vascular Centre, Skåne University Hospital, Malmö, southern Sweden, between August 2013 and June 2015. Patients selected for renal angiography with pressure measurement in the renal arteries due to suspected RAS were invited to participate in the study. All patients had treatment-resistant hypertension, signs of kidney dysfunction, or both, and pathological findings at magnetic resonance angiography of the renal arteries, suggesting RAS. The patients included in the study were asked to fill in a questionnaire and self-rating scales. They were also offered a clinical psychiatric examination. They were also informed that their decision to participate, or not, in the study would not affect their medical or interventional treatment.

During the study period, 71 patients met the inclusion criteria, 12 of which were excluded from the study: ten were unreachable by phone or did not speak Swedish, and two were hospitalized due to severe concomitant somatic disease. The remaining 59 were identified as potential participants in the study (Figure 1). Of these 59 patients, 37 agreed to answer the questionnaire part, including an agreement for the principal investigator to screen the medical records. The self-rating scales Montgomery–Asberg Depression Rating Scale (MADRS-S) [25] and Hospital Anxiety and Depression scale (HAD) [26] were added to the questionnaire part after 12 patients had been included (twelve patients with completed questionnaires and seven patients with completed clinical examinations). Eighteen patients also consented to the psychiatric clinical examination, which also included the HAD for the first seven patients. A total of 25 MADRS-S scores and 29 HAD scores were collected at baseline. Data on the somatic morbidity of the 18 patients who underwent the psychiatric examination are given in Table 1.

![Selection of participants for the study](https://www.heighpubs.org/hda)

**Table 1:** Somatic morbidity in the 18 patients who completed the clinical psychiatric examination. RAS: Renal artery stenosis.

| Condition                        | All (N=18) | With RAS (N=10) | Without RAS (N=7) |
|----------------------------------|------------|-----------------|-------------------|
| Hypertension                     | 18         | 10              | 7                 |
| Cardiovascular disease           | 15         | 10              | 4                 |
| Renal dysfunction                | 5          | 3               | 2                 |
| Type 2 diabetes                  | 4          | 1               | 2                 |
| Atrophic kidney                  | 2          | 2               | 0                 |
| Chronic back pain                | 2          | 2               | 0                 |
| Osteoarthritis                   | 2          | 1               | 1                 |
| Dyslipidaemia                    | 2          | 2               | 0                 |
| Hyperuricaemia                   | 2          | 2               | 0                 |
| Benign prostatic hyperplasia     | 1          | 1               | 0                 |
| Atrial fibrillation              | 1          | 1               | 0                 |
| Rheumatoid arthritis, RA         | 1          | 1               | 0                 |
| Psoriasis                        | 1          | 1               | 0                 |
| Ulcerative colitis               | 1          | 1               | 0                 |
| Stroke sequelae                  | 1          | 1               | 0                 |
| Migraine                         | 1          | 0               | 1                 |
| Chronic pancreatitis             | 1          | 0               | 1                 |
| Chronic obstructive pulmonary disease | 1       | 0               | 1                 |
| Asthma                           | 1          | 0               | 1                 |
Both the questionnaire part of the study and the clinical examination were performed a few weeks before the angiography, thus it was not known whether the patients suffered from significant RAS or not. Due to logistic reasons, 3 of 18 patients were examined shortly after their angiography (two to nine days) instead of before, but the psychiatric examiner was blinded concerning the result.

The questionnaire

The questionnaire included questions regarding: years with hypertension, previous anxiety, an anxiety disorder within the last year, previous treatment of anxiety, anxiety treatment within the last year, previous depression, depression within the last year, previous treatment of depression, depression treatment within the last year, treatment with psychotherapy, previous treatment with psychotropic medications, treatment with psychotropic medications within the last year. If they answered yes on the questions regarding psychotropic medications, they were asked to list the name of the drugs. The questionnaires were returned to the research unit in stamped addressed envelopes.

Self-rating scales

The self-rating Montgomery–Asberg Depression Rating Scale (MADRS-S) [25] consists of nine items scored from 0 to 6 points, giving a maximum of 54 points. A score of 13-19 indicates low-grade depression, 20-34 moderate depression and 35-54 severe depression.

The Hospital Anxiety and Depression scale (HAD) [26] consists of 14 items; seven covering anxiety and seven depression. Each item has a score from 0 to 3 points, giving a maximum score of 21 points for HAD anxiety and 21 points for HAD depression. A score of 0 to 7 points indicates no disorder, 8 to 10 points possible anxiety or depression, and 11-21 points most likely a disorder, on either subscale.

Medical records

Additional medical information on baseline p-renin, baseline p-aldosterone, previous and current medication and somatic comorbidity was collected from all patients’ medical records after the questionnaire or the clinical psychiatric examination.

Clinical psychiatric examination

Patients were examined by a resident in psychiatry (LH). The examination consisted of a structured clinical interview regarding previous and current psychiatric disorders, treatment, previous medical conditions, and family history. The patients were diagnosed according to DSM-IV using the structured diagnostic tool International Neuropsychiatric Interview (M.I.N.I. 6.0). Patients were also rated according to the comprehensive psychopathological rating scale (CPRS) [27], from which the subscales Montgomery–Asberg Depression Rating Scale (MADRS) [28] and the Brief Scale of Anxiety (BSA) [29] were extracted. In the CPRS each item is rated from zero to three points, and the score was therefore doubled to correspond to MADRS. The first seven patients also filled in the HAD during the clinical psychiatric examination.

Angiography and endovascular intervention

The mean pressure gradient between the aorta and the renal artery was measured during renal angiography, and if found to be ≥ 10 mmHg the patient was diagnosed as having RAS and underwent PTRA, followed by stent placement [30-32].

Data collection at follow-up

The follow-up part started after data had been collected on twelve patients. The first seven patients in the clinical psychiatric examination were also included in the follow-up. Thirty-two patients participated in the follow up part, of these 18 went through the psychiatric clinical examination at baseline.

In this part all patients filled in MADRS-S and HAD at baseline, together with the questionnaire and agreed to be contacted again after six months (follow up). At follow up the patients were contacted again by post six months after the angiography and intervention, and asked fill in the self-rating MADRS-S and HAD self-rating scales. Of the 32 patients included in the follow up 25 returned the envelopes. Of the 18 patients who went through the psychiatric clinical examination 16 returned the envelopes. During the same week, blood samples were collected in the morning for analysis of plasma (p-) aldosterone and p-renin. Patients also participating in the clinical psychiatric examination were interviewed by phone concerning changes in their medication.

Laboratory analyses

The standard methods for the analysis of p-aldosterone and p-renin were changed on November 17, 2014. Before November 17, 2014, p-aldosterone was determined using a radioimmunoassay (Aldosterone Coat-A-Count, DPC, CA, USA). The normal range for p-aldosterone is 110-860 pmol/L, and the imprecision (coefficient of variation, CV) were 8.6% at 100 pmol/L and 5.2% at 750 pmol/L. Plasma-renin was determined with the DiaSorin Liaison method (Liaison Renin, DiaSorin, Saluggia, Italy). The CV for this method were 4.4% at 27 mIU/L and 3.7% at 102.

After November 17, 2014 p-aldosterone and p-renin were determined with the IDS-iSYS methods (Immunodiagnostics Systems Nordic a/s, Copenhagen, Denmark). The CV for the aldosterone method was/is 18% and 10% at 230 and 555 pmol/L. The normal range for renin using this method was 5-80 mIU/L, and the CV was 10% at both 14 and 92 mIU/L.

All renin and aldosterone values prior to angiography were collected from patient files. These blood samples were all
drawn prior to November 17, 2014 and therefore analysed with the same method as described above. In the follow up blood samples were collected both prior to and after November 17, 2014. Fourteen out of 25 samples were determined with the previous method. Since the two renin methods showed good correlation (R=0.94), all p-renin values from the RIA were recalculated to the IDS-iSys method. Aldosterone values were not recalculated, because the correlation between methods was deemed insufficient in the low normal concentration range (< 400 pmol/L).

Results

Characteristics

Twenty-one of the 37 patients completing the questionnaire were found to have a pressure gradient of ≥ 10 mmHg at angiography, indicating significant stenosis [33]. All 21 underwent PTRA and 18/21 also underwent successful renal artery stenting. The 15 patients in whom no significant pressure gradient was observed underwent diagnostic angiography only. In one patient the renal artery could not be reached during angiography due to severe atherosclerosis of the abdominal aorta. The presence of RAS could therefore not be determined. Ten of the 18 patients participating in the clinical psychiatric examination were found to have a pressure gradient of ≥ 10 mmHg, and were diagnosed as having significant stenosis [30]. All ten underwent PTRA, and 8/10 also underwent successful renal artery stenting.

The characteristics of the 37 patients who answered the questionnaire are given in Tables 2 and 3. Among these patients, 13 reported anxiety at least once during their life, and eight reported anxiety during the previous year. Eight patients reported previous depression, and four reported depression during the previous year.

Psychiatric rating scales

No significant difference was found in baseline MADRS-S scores between patients with RAS (N=15; MADRS-S 6.4 ± 4.9) and without RAS (N=9; 7.3 ± 6.8) in the questionnaire part of the study. Furthermore, no significant differences were found in total baseline HAD scores, baseline HAD anxiety, or baseline HAD depression scores between patients with RAS (N=17) and those without RAS (N=11). See Table III for HAD scores.

In the questionnaire part of the study, eleven patients reported anxiety or depression within the past year. Among these, the means MADRS-S scores were 12.5 ± 9.2 (N=6) at baseline, and 13.4 ± 7.5 (N=9) at follow-up (NS). The corresponding total HAD scores were 13.7 ± 6.2 (N=9) at baseline and 13.0 ± 5.5 (N=9) at follow-up (NS).

MADRS-S and HAD scores for patients who took part in the clinical psychiatric examination and were diagnosed as currently having anxiety or depression (N=10) were analysed separately. All patients currently suffering from depression also had a current anxiety disorder. The MADRS-S scores for this group were 13.0 ± 7.8 (N=7) at baseline and 13.1 ± 8.3 (N=9) at follow-up (NS). The corresponding total HAD scores were 12.7 ± 6.5 (N=10) at baseline and 12.3 ± 5.7 (N=9) at follow-up (NS). The HAD anxiety and depression scores were 8.0 ± 3.5 and 4.7 ± 3.5 (N=10) at baseline, and 6.8 ± 3.9 and 5.6 ± 3.2 (N=9) at follow-up (NS for both).

The 18 patients with RAS and successful PTRA and stenting were also analysed separately. Their HAD anxiety and depression scores were 3.9 ± 2.8 and 2.2 ± 1.8 at baseline (N=16), and 4.0 ± 1.8 and 2.7 ± 2.4 at follow-up (N=13) (NS for both).

Hormones and blood pressure

The mean value of p-renin prior to the intervention was 296 ± 680 mIU/L (N=13). The corresponding value for p-aldosterone was 372 ± 539 pmol/L (N=13). This means that renin, but not aldosterone, levels were increased compared to the normal range in all patients.

Table 2: Questionnaire results. Anxiety, depression and treatment according to the questionnaire in patients who only answered the questionnaire and in patients who also completed the clinical psychiatric examination. Statistics: Pearson’s chi square test.

| Questionnaire, only (N=19) | Clinical psychiatric examination (N=18) | P     |
|---------------------------|----------------------------------------|-------|
| Previous experience of anxiety Yes/no/uncertain | 5/13/1 | 8/8/2 | 0.33 |
| Anxiety the last year Yes/no/uncertain | 2/15/2 | 6/11/1 | 0.23 |
| Previous anxiety treatment Yes/no/uncertain | 1/18 | 5/13 | 0.063 |
| Anxiety treatment the last year Yes/no | 0/19 | 1/17 | 0.30 |
| Previous depression Yes/no/uncertain | 0/16/3 | 8/5/5 | 0.001 |
| Depression the last year Yes/no/uncertain | 0/18/1 | 4/10/4 | 0.018 |
| Previous depression treatment Yes/no | 0/18 | 8/10 | 0.001 |
| Depression treatment the last year Yes/no | 0/19 | 1/17 | 0.30 |
| Treated with psychotherapy Yes/no | 1/18 | 1/17 | 0.94 |
| Treated with psychotropic medication Yes/no | 4/15 | 6/12 | 0.40 |
| Current treatment with psychotropic medication Yes/no | 1/18 | 0/17 | 0.34 |
P-renin level in patients with RAS (N=6) (623 ± 934 mIU/L) was significantly higher than in patients without RAS (N=6) (10.4 ± 7.9 pmol/L) (p = 0.002), whereas no difference was found in p-aldosterone levels between the groups. See Table 3 for aldosterone levels.

The mean value of p-renin at the 6-month follow-up (N=25) was 132 ± 341 mIU/L. No significant difference was found when comparing p-renin before and after stenting in patients with RAS who were successfully stented (N=4, p = 0.068).

Mean BP at admission was 172 ± 24 / 83 ± 11 mmHg (N=36). No differences in BP were found between patients without RAS and those later diagnosed with RAS.

Correlation analysis

Values of p-aldosterone were available in patient records for 13 of 37 patients, of whom nine had MADRS-S scores. No significant correlation was found between p-aldosterone level and MADRS-S in this group. However, p-aldosterone was significantly correlated to the MADRS-S score in patients with RAS (N=5, p = 0.037).

No significant correlations were found between renin levels and HAD anxiety, HAD depression or MADRS-S scores at the six-month follow-up.

Clinical psychiatric examination

**DSM-IV diagnosis:** The characteristics of the examined patients (N=18) are given in Table 4. Ten of the examined patients (56%) met the criteria for an anxiety disorder. Two patients were diagnosed as currently suffering from depression. Eight of the patients did not meet the criteria for a current DSM-IV diagnosis.

**Suicidality:** Three of the eighteen patients (16.6%) had attempted suicide. One had RAS, one could not be investigated, and one did not have RAS. The suicide attempts were 23-47 years ago. Two of three attempts had occurred during a depressive episode.

As the questionnaire part of the study did not include Table 3: Characteristics of the 37 participants who completed the questionnaire. RAS: Renal artery stenosis, MADRS-S (Montgomery–Asberg Depression Rating Scale, Self-administered), HAD: Hospital Anxiety. Depression scale. BP: Blood pressure. Normal range of p-renin is 5-80 mIU/L and for p-aldosterone 110-860 pmol/L. Renin and aldosterone before angiography are presented. Mean ± 1 SD.

|                      | All (N=37) | With RAS (N=21) | Without RAS (N=15) |
|----------------------|------------|-----------------|--------------------|
| Age                  | 64.1 ± 12.0| 63.2 ± 13.0     | 64.9 ± 11.1        |
| Sex (M/F)            | 16/21      | 12/9            | 4/11               |
| MADRS-S              | 7.6 ± 6.9  (N=25) | 6.4 ± 4.9 (N=15) | 7.3 ± 6.8 (N=9)    |
| MADRS-S follow-up    | 8.8 ± 7.0  (N=25) | 6.9 ± 4.8 (N=16) | 10.4 ± 7.7 (N=8)   |
| HAD                  | 7.4 ± 6.2  (N=29) | 5.9 ± 3.9 (N=17) | 8.0 ± 7.0 (N=11)   |
| HAD follow-up        | 8.3 ± 5.6  (N=24) | 6.4 ± 3.3 (N=15) | 9.9 ± 6.4 (N=8)    |
| HAD Anxiety          | 4.6 ± 3.9  (N=29) | 3.8 ± 2.7 (N=17) | 4.8 ± 4.5 (N=11)   |
| HAD Anxiety follow-up| 4.8 ± 3.2  (N=24) | 3.9 ± 2.0 (N=15) | 5.3 ± 3.5 (N=8)    |
| HAD Depression       | 2.8 ± 2.8  (N=29) | 2.1 ± 1.8 (N=17) | 3.2 ± 3.1 (N=11)   |
| HAD Depression – follow-up | 3.5 ± 3.0  (N=24) | 2.5 ± 2.3 (N=15) | 4.6 ± 3.4 (N=8)    |
| BP at admission (mmHg)| 172 ± 24 / 83 ± 11 (N=36) | 169 ± 19 / 85 ± 9 (N=21) | 174 ± 30/82 ± 12 (N=14) |
| P-renin (mIU/L)      | 296 ± 680  (N=13) | 623 ± 934 (N=6)  | 10.4 ± 7.9 (N=6)   |
| P-aldosterone (pmol/L)| 372 ± 539 (N=13) | 511 ± 755 (N=6)  | 283 ± 276 (N=6)    |

Table 4: Clinical psychiatric examination. DSM-IV diagnosis with International Neuropsychiatric Interview. CPRS: Comprehensive Psychopathological Rating Scale. MADRS: Subscale of the CPRS for depression. Montgomery–Asberg Depression Rating Scale. BSA: Subscale of CPRS for anxiety. Brief Scale of Anxiety.

|                      | All (N=18) | With RAS (N=10) | Without RAS (N=7) |
|----------------------|------------|-----------------|-------------------|
| Age Mean ± SD        | 65.4 ± 12.1 (N=18) | 67.9 ± 10.3 (N=10) | 60.7 ± 14.5 (N=7) |
| Sex (M/F)            | 8/10       | 7/3             | 1/6               |
| Current psychiatric diagnosis | 10        | 4               | 5                |
| Any anxiety disorder | 10         | 4               | 5                |
| Generalized anxiety disorder | 6        | 2               | 3                |
| Social phobia        | 6          | 2               | 3                |
| Agoraphobia          | 3          | 2               | 0                |
| Panic disorder       | 2          | 0               | 1                |
| Obsessive-compulsive disorder | 2        | 1               | 0                |
| Depression           | 2          | 1               | 0                |
| Alcohol dependence   | 1          | 0               | 1                |
| Previous psychiatric diagnosis | 9        | 4               | 4                |
| Depression N=9       | 9          | 4               | 4                |
| Panic disorder N=3   | 3          | 0               | 2                |
| MADRS Mean ± SD      | 9.9 ± 8.4  (N=18) | 6.9 ± 6.8 (N=10) | 10.9 ± 4.2 (N=7)  |
| BSA Mean ± SD        | 5.7 ± 3.7  (N=18) | 4.6 ± 3.3 (N=10) | 5.9 ± 2.4 (N=7)   |
Correlation analysis: No correlations were found between p-renin or p-aldosterone levels and the CPRS subscales MADRS or BSA (N=6). Strong correlations were found between the self-rated MADRS-S score and the MADRS score resulting from the psychiatric examination (R=0.83, \( p=0.001 \)), and between the HAD anxiety score and the BSA score (R=0.80, \( p<0.001 \)).

Bias

Thirty-seven patients completed the questionnaire, and of these 18 were examined by the resident in psychiatry. Cross tabulation was used to investigate whether the 18 patients examined differed from the patients participating only in the questionnaire part of the study. Significantly more patients who participated in the clinical psychiatric examination reported a history of depression than among those who did not take part in the psychiatric examination: 8/18 versus 0/19 (\( p=0.001 \)). No significant difference was found in the reported history of anxiety between the patients who participated in the clinical psychiatric examination and those who did not take part in the psychiatric examination: 8/18 versus 5/19 (\( p=\text{NS} \)) (Table 2).

Discussion

In this study we found a high rate of anxiety disorders (56%) and previous suicide attempts (17%) in patients with suspected RAS, when evaluated by psychiatric examination. We therefore suggest that anxiety disorders are more common in patients with suspected RAS than in the general population. The lifetime prevalence of an anxiety disorder in the general population is between 10% and 29% [2,3]. The anxiety rate in patients with suspected RAS may also be higher than in patients with hypertension in general. In a study based on over two million patients in Sweden, investigating registered diagnoses over a five-year period, it was found that 7.1% of the patients with hypertension also had a diagnosis of anxiety, compared with 6.3% of normotensive patients [34]. Patients with PA and RAS both overproduce aldosterone. A higher ratio of anxiety has been found in patients with PA (12/23 = 52%) than in those with essential hypertension (4/23 = 17%) [24]. Few studies have been carried out to investigate psychiatric comorbidity in RAS patients. However, it was shown in a recent case report [35] that successful PTRA in a patient with RAS cured problems of anxiety, negativism and asthenia. Renin and aldosterone levels were normalized one month after the intervention, and the psychiatric symptoms had disappeared.

Depression, on the other hand, was not as frequent as anxiety in our study. Only 2 out of 18 (11%) patients who underwent the psychiatric examination met the DSM-IV criteria for current depression. In a study by Sandstrom, et al., the prevalence of depression in patients with hypertension was found to be 9.6%, compared to 6.9% in normotensive patients [34].

In spite of the high rate of anxiety disorders the patients in the present study had relatively low MADRS-S and HAD scores. The HAD anxiety score for patients who met the DSM-IV criteria for diagnosis of an anxiety disorder was 8.0 \( \pm 3.5 \), corresponding to a possible anxiety disorder [26]. The discrepancy between the M.I.N.I. interview score and the self-rating scores may reflect the better evaluation of anxiety in the M.I.N.I. interview than the self-rated HAD scores.

An important limitation of this study concerning the high rate of anxiety disorders and suicide attempts is that it could be argued that only patients with psychiatric illness were interested in taking part in the clinical psychiatric examination, and that the high occurrence of psychiatric morbidity might, therefore, be a result of selection bias. Therefore, cross tabulation was used. This showed that self-reported anxiety by those completing the questionnaire was not over-reported in the group of patients who underwent the psychiatric examination, compared to those who only participated in the questionnaire part of the study. However, patients participating in the psychiatric examination reported previous depression more frequently than patients who only completed the questionnaire. This suggests that patients with previous depression were more prone to participate in the clinical psychiatric examination.

Questions about previous suicide attempts were not included in the questionnaire in this study, and thus cross tabulation could not be used to investigate possible selection bias regarding suicide attempts. The suicide attempts had occurred 23-47 years ago, before the onset of hypertension. The occurrence of a previous suicide attempt in this study is estimated between 4.2% and 16.6%. This should be compared with a lifetime prevalence of 2.7% reported for suicide attempts in the general population [36]. Thus, it is possible that the rate of previous suicide attempts is increased in patients with suspected RAS.

Patients suffering from RAS exhibit over-activation of their RAAS [37]. In this study we found increased renin levels but not aldosterone levels, and that patients with RAS had significantly higher renin levels than those without RAS. The reason for this discrepancy may be effects of antihypertensive medication. It should also be mentioned that when the study was designed, it was assumed that renin and aldosterone levels were determined in all patients remitted for renal angiography, however, this was the case in only 13 of 37 patients. The fact that renin, but not aldosterone, levels were elevated in RAS patients is probably explained by the fact that patients both with and without a significant pressure gradient in the renal artery showed some degree of kidney dysfunction.
One of our hypotheses was that symptoms of depression or anxiety would be reduced after endovascular dilatation in RAS patients. However, when RAS patients who were successfully stented rated their symptoms of anxiety and depression before and 6 months after the intervention, no such effect was found.

The patients included in this study had a large number of somatic disorders, and medication may also have affected the results. It has generally been thought that treatment with beta-blockers may induce symptoms of depression. However, recent studies do not support this [38]. Another limitation is the fact that 10 patients could not be included in the study as they were impossible to reach by phone or did not speak Swedish. However, this limitation applies to all studies using questionnaires.

The mechanisms by which overactivation of the RAAS can have depressive and anxiogenic effects are unclear. MR are found in the hippocampus, amygdala and nucleus accumbens ([16]. Animal studies have shown that aldosterone can pass through the blood–brain barrier [39]. Rats treated with aldosterone exhibit symptoms of depression and anxiety [17,18], and we have previously found a positive correlation between depressive symptoms and aldosterone concentration in humans [40]. In the present study we also found a positive correlation between the MADRS-S score and aldosterone level in patients with RAS, although the number of patients was small.

In 90% of cases, RAS is due to atherosclerosis. Patients with general atherosclerosis have elevated levels of cytokines, and patients with RAS have been found to exhibit elevated inflammatory activity. Cytokines are elevated compared to healthy controls in all patients undergoing renal angiography – both those with, and those without, a pressure gradient between the aorta and the renal artery [31].

Increased levels of cytokines have also been reported in patients with depression [41], and there is evidence that inflammation is involved in depressive pathophysiology, although the extent is debated [42]. The role of neuroinflammation in anxiety disorders, on the other hand, has been less investigated, but the overproduction of cytokines, similar to that in depression, has also been reported in this setting [43].

Inflammation has also been linked to suicidal behaviour [44,45]. Although the possible higher incidence of previous suicide attempts observed in this study occurred before the onset of hypertension. Increased inflammatory activity in these patients might have been a risk factor for both suicide and somatic illness. This should be investigated in future studies.

The results of this study suggest that patients with suspected RAS have a high psychiatric comorbidity, as 56% of the patients undergoing psychiatric examination met the DSM-IV criteria for a current anxiety disorder. Previous studies have also found an increased prevalence of anxiety in patients with hypertension. In light of these findings, it may be important to examine the occurrence of anxiety disorders in patients with treatment-resistant hypertension.

We were not able to find any improvement in depressive or anxiety symptoms after PTRA, followed by stent placement. As the number of patients included in this study was small, and no control group was used, more extensive studies are warranted.

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