Determinants of Erectile Dysfunction in Hemodialysis Patients in CNHU-HKM, Cotonou

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Abstract

Objective: The purpose of this study is to identify determinants of erectile dysfunction in hemodialysis patients in CNHU-HKM, Cotonou.

Method: It was a cross-sectional, descriptive and analytical study conducted over a period of six months, from 1st March to 30th August 2017. Prospective data collection was carried out, along with a comprehensive census of hemodialysis patients meeting inclusion criteria. Sexual function was assessed through full version of International Index of Erectile Function (IIEF).

Results: 114 hemodialysis patients were included in the study, with mean age estimated at 51.05 years ± 11.66. Among the study population, 100 patients (87.7%) had hypertension as comorbidity, hypertensive Nephropathy (87.5%) and Diabetic Nephropathy (14%). The average duration on dialysis was 74.34 ± 55.74. Forty-three hemodialysis patients (37.7%) were with depression, 56 (49.1%) had lower testosterone, 38 (33.3%) high rate of LH and 63 (55.3%) high FSH. 98 (86%) of them experienced erectile dysfunction, 97(85.1%) sexual desire disorder and 80 (70.2%) orgasm disorder. Determinants of erectile dysfunction in hemodialysis patients included age (p<0.001; OR=1.168), hypertension (p<0.001; OR=10.111), senility in hemodialysis (p=0.008; OR=1.024), depression (p=0.022; OR=11.250) and testosteronemia (p=0.038; OR=0.740).

Conclusion: Sexual dysfunction prevalence is high in patients receiving dialysis. There is need to include care and support in these patient’s treatment strategies.

Keywords: Determinants; Hemodialysis; Erectile dysfunction

Introduction

Chronic Kidney Disease (CKD) is a heavily debilitating disease characterized by the loss of kidneys major functions [1]. It is a global public health issue, particularly in developing countries. Best estimates in sub-Saharan Africa suggest that 12 to 23% of adults develop CKD [2], with admission prevalence estimated at 7.5% [3]. Sexual dysfunction is one of the complications of CKD. It is a real problem confronting the patient on daily basis, though often under-estimated by healthcare staff. This is partly due to the insignificant number of studies conducted on this disease and the patient’s reluctance to discuss with the physician [4]. Male sexual dysfunction is defined by disorders in terms of erection, sexual desire, ejaculation and orgasm [5].

The frequency of these sexual disorders is amplified in CKD patients among whom it is estimated between 50% and 70%, depending on the stage of kidney disease [4,6,7]. Half of end-stage CKD patients suffer from sexual disorders with reduced libido in association with impaired spermatogenesis [8]. Among these sexual disorders, erectile dysfunction (ED) was the most studied condition. Its prevalence during CKD is 49-55% [1,9].

Major advancement in healthcare for CKD patients in recent decades have resulted in rise in their life expectancy. So, at an advanced stage, a substitution treatment is required: “transplant” or “dialysis.” The latter worsens pre-existing endocrine disorders and adversely affects the patient quality of life [8]. According to Nzibiti, in 75% of cases a patient’s sexual life is affected after having received dialysis [10]. Sexual dysfunction prevalence rises from 9% to 60-70% after dialysis [11]. Hence, erectile dysfunction (ED) estimated at 9% in CKD patients not on dialysis is close to 90% in dialysis patients [6]. Organic factors such as endocrine disorders affecting the hypothalamic-pituitary-gonad axis [10] and arterial disease specific of patients on dialysis [12] were revealed as well as risk factors: age, atherosclerosis, dialysis duration, and initial nephropathy [1]. The purpose of this study is to identify determinants of erectile dysfunction in hemodialysis patients in CNHU-HKM, Cotonou.

Materials and Methods

Type of study and study population

We carried out a cross-sectional, descriptive and analytical study from 1st March to 30th August 2017. The study population was made up of male hemodialysis patients seen in Nephrology-Hemodialysis Department of Hubert Koutoukou Maga Teaching Hospital (CNHU-HKM), Cotonou.

(a) Inclusion criteria

The followings were included in our study:

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• Patients who have been on dialysis for at least 6 months prior to the evaluation.
• Male hemodialysis patients, from 18 to 80 years.
• Patients having given their informed consent to participate in the study.

(b) Non-inclusion criteria

Patients non-included in our study were those:
• On dialysis catheter.
• With trauma or pathology related to the spine and pelvis.
• Unable to provide responses to questions, or having mental disorders.
• Who did not give their consent.

(c) Exclusion criteria

• Patients who did not carry out biological examinations.

Method and sampling technique

We resorted to non-probabilistic sampling method with comprehensive census of all hemodialysis patients meeting the inclusion criteria.

Variables and their operational aspects

(a) Dependent variable: The dependent variable was sexual dysfunction. Sexual function was assessed through full version of International Index of Erectile Function (IIEF). It was validated as a convenient and reliable diagnosis tool for sexual dysfunction. IIEF includes 15 questions addressing five areas: Erection, orgasm, sexual desire, satisfaction with sexual intercourse and overall satisfaction. Answers to each question represent the patient experience in the last four weeks using a five-point scale. The score of an area: Erection, orgasm, sexual desire, satisfaction with sexual intercourse and overall satisfaction is obtained by summing the scores of answer provided for each question in the area (Table 1) [13].

(b) Independent variables

(i) Socio-demographic variables
• Age: Expressed in number of completed years of age and clustered by age-group.
• Marital status: Married/concubinage, single, divorced, widowed.

(ii) Medical-surgical history related variables
• History of pelvic trauma (pelvis and/or perineum).

Table 1: IIEF interpretation.

<table>
<thead>
<tr>
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</tr>
</thead>
<tbody>
<tr>
<td>Desire</td>
<td>0-2</td>
<td>3-4</td>
<td>5-6</td>
<td>7-8</td>
<td>9-10</td>
</tr>
<tr>
<td>Erection</td>
<td>0-6</td>
<td>7-12</td>
<td>13-18</td>
<td>19-24</td>
<td>25-30</td>
</tr>
<tr>
<td>Orgasm</td>
<td>0-2</td>
<td>3-4</td>
<td>5-6</td>
<td>7-8</td>
<td>9-10</td>
</tr>
<tr>
<td>Satisfaction</td>
<td>0-3</td>
<td>4-6</td>
<td>7-9</td>
<td>10-12</td>
<td>13-15</td>
</tr>
<tr>
<td>with sexual intercourse</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Overall satisfaction</td>
<td>0-2</td>
<td>3-4</td>
<td>5-6</td>
<td>7-8</td>
<td>9-10</td>
</tr>
</tbody>
</table>

Interpretation

Areas: Severe dysf., Moderate dysf., Moderate to light dysf., Light dysf., No dysf.

History of abdominal-pelvic surgery
History of stroke
History of exposure to ionizing radiation.
Other conditions

(ii) Comorbidities related variables
• High blood pressure (HBP)
• Diabetes
• Peripheral neuropathy: Patients with dermal hypoesthesia, paraesthesia, burning feet and/or absent deep tendon reflexes were declared to have developed peripheral neuropathy.
• Congestive heart failure
• Prostatic hypertrophy
• Psychiatric pathology

(iii) Cardiovascular risk factors related variables
• Alcoholism: It was quantified in terms of number of grams of alcohol per day.
• Smoking: It was quantified in accordance with the number of sticks per day.
• Dyslipidemia: Matching one of the following anomalies:
  • LDL cholesterol levels ≥ 1.5 g/l or specific therapy ongoing
  • HDL cholesterol levels <0.40 g/l or specific therapy ongoing
  • Total cholesterol levels >2 g/l or specific therapy ongoing
  • Triglycerides levels >1.5 g/l or specific therapy ongoing

(iv) Initial nephropathy related variable
• Diabetic nephropathy
• Nephroangiosclerosis
• Chronic glomerulonephritis
• Polycystic kidney disease
• Indeterminate or others

(v) Hemodialysis parameters related variables
• Seniority in hemodialysis (expressed in months)
• Frequency of hemodialysis sessions per week

(vi) Variable relating to high risk treatment of sexual dysfunction
• β – blockbers
• Central anti-hypertensive
• Cimetidine
• Neuroleptic medication
• Antidepressant (MAOIs, anti-cholinergic drug)
• Diuretic
• Hypolipidemic drugs

(vii) Depression
It was evaluated on the basis of Patient Health Questionnaire.
(PHQ). It is an evaluation tool for depression diagnosis. It includes 9 questions directly from the nine signs and symptoms of depression according to DSM-IV. Items 1 to 9 are rated on 0-3 scale and item 10 (level of functioning) is rated on 0-4 scale. A score ranging from 1 to 4 suggests absence of depression, 5 to 9 mild depression, 10 to 14 moderate depression, 15 to 19 moderately severe depression, 20 to 27 severe depression [14].

Sensitivity and specificity of PHQ is higher than those of other questionnaires of the same caliber [15,16].

(viii) Physical examination related variables
- Nutritional status
  It was evaluated through calculation of Body Mass Index (BMI) on the basis of weight (expressed in kg) and height (expressed in meters). The classification was elaborated as follows [17]:
  - Malnutrition: BMI<18.5
  - Normal BMI: BMI between 18.5 and 25
  - Overweight: BMI between 25.1 and 30
  - Grade I overweight: BMI between 30.1 and 35
  - Grade II overweight: BMI between 35.1 and 40
  - Grade III overweight: BMI ≥ 40
  - Abdominal overweight: When waist size is greater than or equal to 102 cm [18].
- Secondary sexual features
- Gynecomastia
- Femoral pulse
- Femoral bruit
- External genitals examination

(ix) Hormonal parameters related variables
- Total testosterone
- Rate of LH
- Rate of FSH

Blood sample was taken from each patient included in the study for FSH, LH and testosterone competitive heterogeneous assays using a multiparameter device labelled Vidas® (Bio-Mérieux). The results of various exams were interpreted on the basis of usual values recorded in the case of an adult male considered as healthy:
- Total testosterone: 2, 5-8.5 ng/ml
- LH : 1-8 mUI/ml
- FSH: 1-5 mUI/ml

Results

This study, carried out as part of academic work, was in strict compliance with the principles of good clinical practices (GCP). Patients gave their free and informed written consent. Confidentiality was strictly maintained during data collection. The information obtained as part of this study was processed in complete anonymity. The results from the different analyses were transmitted to the physician for the benefit of patients.

Socio-demographic characteristics

(a) Age: The respondents mean age was 51.05 years ± 11.66 with extremes values of 24 and 75 years. 46-55 years age group was the most represented (33.33%).

(b) Marital status: Among the 114 patients included in the study: 93 (81.6%) were married. Table 2 outlines their distribution according to marital status.

Sexual dysfunction

(a) Sexually active patients: Over 80% of our patients (92) were still sexually active and 19.3% (22 patients) were no more.

(b) Prevalence of erectile dysfunction: Hemodialysis patients with erectile dysfunction were 98, representing 86%. Among the 98 patients suffering from erectile dysfunction, 31 (31.6%) had severe erectile dysfunction.

Determinants of erectile dysfunction

(a) Univariate analysis

(i) Age and erectile dysfunction: The prevalence of erectile dysfunction increases significantly with age (p<0.001). Table 3 outlines this correlation.

(ii) Comorbidities and erectile dysfunction: Among comorbidities, only hypertension is associated with the

<table>
<thead>
<tr>
<th>Category</th>
<th>Population</th>
<th>Frequency (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Single</td>
<td>12</td>
<td>10.5</td>
</tr>
<tr>
<td>Married/cohabitation</td>
<td>93</td>
<td>81.6</td>
</tr>
<tr>
<td>Divorced</td>
<td>2</td>
<td>1.8</td>
</tr>
<tr>
<td>Widower</td>
<td>7</td>
<td>6.1</td>
</tr>
<tr>
<td>Total</td>
<td>114</td>
<td>100</td>
</tr>
</tbody>
</table>

Table 2: Patients distribution according to marital status.
Table 3: Correlation between age and erectile dysfunction.

<table>
<thead>
<tr>
<th>Age group</th>
<th>≤ ED</th>
<th>≥ ED</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>≤ 25</td>
<td>3 (100.0%)</td>
<td>0 (0.0%)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>26-35</td>
<td>4 (57.14%)</td>
<td>3 (42.86%)</td>
<td></td>
</tr>
<tr>
<td>36-45</td>
<td>4 (15.38%)</td>
<td>22 (84.62%)</td>
<td></td>
</tr>
<tr>
<td>46-55</td>
<td>5 (13.16%)</td>
<td>33 (86.84%)</td>
<td></td>
</tr>
<tr>
<td>56-65</td>
<td>0 (0.00%)</td>
<td>27 (100.00%)</td>
<td></td>
</tr>
<tr>
<td>≥ 66</td>
<td>0 (0.00%)</td>
<td>13 (100.00%)</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>16 (14.04%)</td>
<td>98 (85.96%)</td>
<td></td>
</tr>
</tbody>
</table>

Table 4: Correlation between co-morbidities and erectile dysfunction.

<table>
<thead>
<tr>
<th>Comorbidities</th>
<th>≤ ED</th>
<th>≥ ED</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hypertension</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>9 (9.0%)</td>
<td>91 (91.0%)</td>
<td>0.001</td>
</tr>
<tr>
<td>Absent</td>
<td>7 (50.0%)</td>
<td>7 (50.0%)</td>
<td></td>
</tr>
<tr>
<td>Diabetes</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>14 (93.3%)</td>
<td>1 (6.66%)</td>
<td>0.338</td>
</tr>
<tr>
<td>Absent</td>
<td>84 (84.84%)</td>
<td>15 (15.15%)</td>
<td></td>
</tr>
<tr>
<td>Peripheral neuropathy</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>18 (100.00%)</td>
<td>0 (0.0%)</td>
<td>0.051</td>
</tr>
<tr>
<td>Absent</td>
<td>80 (83.3%)</td>
<td>16 (16.7%)</td>
<td></td>
</tr>
<tr>
<td>Heart disease</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>3 (100.00%)</td>
<td>0 (0.0%)</td>
<td>0.633</td>
</tr>
<tr>
<td>Absent</td>
<td>95 (85.6%)</td>
<td>16 (14.4%)</td>
<td></td>
</tr>
<tr>
<td>HBP</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Present</td>
<td>5 (100.00%)</td>
<td>0 (0.0%)</td>
<td>0.463</td>
</tr>
<tr>
<td>Absent</td>
<td>93 (85.3%)</td>
<td>16 (14.7%)</td>
<td></td>
</tr>
</tbody>
</table>

Table 5: Seniority in hemodialysis and erectile dysfunction.

<table>
<thead>
<tr>
<th>Seniority in hemodialysis (expressed in months)</th>
<th>≤ ED</th>
<th>≥ ED</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>≤ 12</td>
<td>4 (36.36%)</td>
<td>7 (61.06%)</td>
<td>0.0089</td>
</tr>
<tr>
<td>13-48</td>
<td>10 (27.03%)</td>
<td>27 (72.97%)</td>
<td></td>
</tr>
<tr>
<td>49-84</td>
<td>0 (0.00%)</td>
<td>22 (100%)</td>
<td></td>
</tr>
<tr>
<td>85-120</td>
<td>2 (7.69%)</td>
<td>24 (92.31%)</td>
<td></td>
</tr>
<tr>
<td>121-156</td>
<td>0 (0.00%)</td>
<td>4 (100%)</td>
<td></td>
</tr>
<tr>
<td>157-192</td>
<td>0 (0.00%)</td>
<td>7 (100%)</td>
<td></td>
</tr>
<tr>
<td>≥ 193</td>
<td>0 (0.00%)</td>
<td>7 (100%)</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>16 (14.04%)</td>
<td>98 (85.96%)</td>
<td></td>
</tr>
</tbody>
</table>

Table 6: Correlation between depression and erectile dysfunction.

<table>
<thead>
<tr>
<th>Depression</th>
<th>≤ ED</th>
<th>≥ ED</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>No</td>
<td>15 (21.1%)</td>
<td>56 (78.9%)</td>
<td>0.003</td>
</tr>
<tr>
<td>Yes</td>
<td>1 (2.3%)</td>
<td>42 (97.7%)</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>16 (14.0%)</td>
<td>98 (86.0%)</td>
<td></td>
</tr>
</tbody>
</table>

Table 7: Correlation between hormonal profile and erectile dysfunction.

<table>
<thead>
<tr>
<th>Testosteronemia</th>
<th>T</th>
<th>P</th>
<th>95% IC</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.182</td>
<td>0.031</td>
<td>[0.09; 1.9]</td>
<td></td>
</tr>
<tr>
<td>Rate of LH</td>
<td>0.869</td>
<td>0.397</td>
<td>[-2.5; 6.1]</td>
</tr>
<tr>
<td>Rate of FSH</td>
<td>0.965</td>
<td>0.349</td>
<td>[-5.4; 14.4]</td>
</tr>
</tbody>
</table>

Table 8: Determinants of erectile dysfunction.

<table>
<thead>
<tr>
<th>Determinants</th>
<th>OR</th>
<th>95% IC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>&lt;0.001</td>
<td>1.168</td>
</tr>
<tr>
<td>Hypertension</td>
<td>&lt;0.001</td>
<td>10.111</td>
</tr>
<tr>
<td>Seniority in hemodialysis</td>
<td>0.008</td>
<td>1.024</td>
</tr>
<tr>
<td>Depression</td>
<td>0.022</td>
<td>11.250</td>
</tr>
<tr>
<td>Testosteronemia</td>
<td>0.038</td>
<td>0.74</td>
</tr>
</tbody>
</table>

Discussion

Socio-demographic characteristics

(a) Age: In our study, the patient’s mean age was 51.05 ± 11.66 years. It is close to the age reported by Aribi et al. in Tunisia estimated at 51.2 years [1]. Several other studies reported similar results. In these various studies, mean age ranged from 48 to 53 years [7,19-23]. Over half of our patients (57.01%) were aged between 45 and 65 years. This could be explained by the fact that the prevalence of end stage CKD increases with age, particularly in the case of men [24]. In our study, extreme values of age which were 24 and 75 years are close to those of Aribi et al., Messina et al. and Avakoudjo et al. who respectively recorded 25 and 78 years; 21 and 76; 27 and 78 years [1,7,9] as extreme values. This could be due to the fact that end stage CKD is not only limited to the elderly, as it can also occur at any age.

(b) Marital status: Most respondents were married (81.6%). This high rate is similar to the studies carried out by Messina et al. and Mumtaz et al. who reported 87.9 and 96.6% of married patients [7,21]. This finding is not surprising as patients included in these different studies are adults as shown by the mean age estimated at 51.18 years.

Sexual dysfunction

(a) Sexually active patients: Nearly 1/5 of our patients were no more sexually active. It is also the case of studies conducted by Aribi et al. and Tounakara et al. who respectively recorded 26% and 30% of sexually inactive patients [1,25]. This could be explained by the fact that patients receiving dialysis lose interest in sexual activities due not only to chronic fatigue induced by dialysis but also depression [11,26]. Moreover, a large number of the patients were in their sixties. Patient in this age range generally have lower sexual activities.

(b) Prevalence of erectile dysfunction: The prevalence of erectile dysfunction recorded in our study was 86%. This result is similar to those reported in other studies. In fact, in West Africa, Ka et al. in Senegal in 2014 reported 84.9% prevalence [22]. In Asia:...
However, this prevalence is significantly higher than data reported in studies conducted in North Africa. Therefore, Aribi et al. [1] in Tunisia in 2016 and Zbiti et al. in Morocco in 2010 reported respectively 65.21% and 43.3% prevalence [1,10]. This gap could be explained by the fact that sexuality is considered a taboo subject in these regions. However, the low ED prevalence reported by Zbiti et al. could be attributable to the young age of the study population (mean age = 45 ± 11.8 years) and the difference in methodology. In fact, Zbiti et al. did not use International Index of Erectile Function (IIEF) to assess erectile function but rather a subjective estimate of the patients.

Factors associated with erectile dysfunction

(a) Age: Our study revealed that the more aged the patient, the higher the risk of ED onset. Aribi et al., Kharbach et al., Messina et al., Costa et al., Muntaz et al., Ka et al., Nassir et al. and Nishida et al. reported the same correlation [1,4,7,20-22,27,29]. In fact, aging is a risk factor for occurrence of atherosclerosis. The latter is part of erectile dysfunction pathogenesis [30]. This could justify the high rate of erectile dysfunction observed in elderly hemodialysis patients.

(b) Hypertension: In our study, hypertension is a risk factor for onset of ED. Aribi et al. reported the same correlation [1]. In fact, erectile dysfunction and hypertension have some pathophysiological mechanisms in common, including atherosclerosis and reduced synthesis of nitric oxide. Moreover, some antihypertensive agents such as β-blockers, diuretics, central antihypertensive drugs used for these patients are source of erectile dysfunction [30]. This justifies the strong tie between hypertension and erectile dysfunction. However, Kharbach et al., Nassir et al. and Malekmakan et al. did not report any correlation between hypertension and erectile dysfunction [4,27,28].

(c) Seniority in hemodialysis: It emerged from our study that, the longer the hemodialysis period, the higher the risk of ED occurrence. Aribi et al., Kharbach et al. and Nishida et al. reported the same correlation [1,4,29]. One of the reasons could be onset of arterial disease which deteriorates progressively [6]. This assumption was confirmed by the results of kidney transplants in early stage of CKD which seem to improve erectile dysfunction [31]. Yet, Messina et al., Ajina et al. and Malekmakan et al. reported no correlation between hemodialysis duration and the onset of erectile dysfunction [7,8,28].

(d) Depression: In this study, depression was closely associated with erectile dysfunction and sexual desire disorder. The same association was reported by Aribi et al. [1] and Cozialiby et al. [1,32]. This could be explained by the fact that some neurotransmitters including: Serotonin, norepinephrine and dopamine are involved in not only controlling sexuality in the central stage, but also the onset of depression [33]. As depression is linked to poor transmission of these neurotransmitter systems, it results in sexual function impairment. Therefore, depressive syndrome may lead to sexual dysfunction in patients receiving hemodialysis [34].

In the same vein, sexual disorders induce changed perception of oneself, thereby contributing to onset of depressive symptoms [11]. It is therefore a vicious life-altering condition for the patients.

(e) Testosteronemia: This study revealed significant decrease in testosterone in patients with erectile dysfunction. This result may be explained by the fact that testosterone stimulates the activity of centers involved in urges for sexual activity [33]. Consequently, hypogonadism causes inactivity of these centers resulting in reduced libido. ED is also the result of elimination of androgens trophic and vasodilation effects on the penis. Thus, hypogonadism is a risk factor for erectile dysfunction. Zbiti et al. and Kharbach et al. [4,10] reached the same conclusion.

Unlike our study and that of Zbiti et al. and Kharbach et al. reported a correlation between the elevation of serum gonadotropins and sexual disorders [4,10].

Determinants of erectile dysfunction

In multivariate analysis: Patient’s age, hypertension, seniority in hemodialysis, depression and testosteronemia are still associated with onset of erectile dysfunction in patients receiving hemodialysis. In fact, our study revealed that testosterone is a protective factor against erectile dysfunction: The risk of erectile dysfunction onset increases with reduced testosterone. Furthermore, hemodialysis patients with depression and hypertension are respectively 11.25 and 10.11 times more exposed to the risk of developing erectile dysfunction.

Conclusion

CKD and hemodialysis have significant impact in several areas of patient’s life. Sexuality is one of the most affected aspects. It is obvious from this study that prevalence of sexual dysfunction is very high in hemodialysis patients. All aspects of sexual function are affected. Determinants of erectile dysfunction were: Age, hypertension, seniority in hemodialysis, depression and lower testosterone. Very few hemodialysis patients discussed their sexual problems with a health officer in view of receiving convenient treatment. Sexual dysfunction must therefore be considered as a major factor impacting on patient’s quality of life, and there is need to include care and support in treatment strategies. It’s important to advance this study in a larger sample size including women receiving hemodialysis.

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References


