



# Prevalence and Risk Factors of Erectile Dysfunction in Patients with Type 2 Diabetes Mellitus in Erbil-Iraq

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#### **Abstract**

**Background and Objectives:** Erectile dysfunction is a highly prevalent disease. It affects multiple aspects of health and can have a serious adverse effect on both the patients and their partners. Diabetes mellitus is a metabolic disorder associated with many chronic complications including erectile dysfunction. The aim of this study was to find out the prevalence and risk factors of erectile dysfunction in patients with type 2 diabetes mellitus.

**Patients and Methods:** A cross-sectional study had been done on 100 adult male patients with type 2 diabetes mellitus, attending endocrinology outpatient at Erbil teaching hospital between June and December 2021, and another 50 age-matched non-diabetic controls. Each one of them underwent detailed history taking, clinical examination, and relevant biochemical study.

**Results:** Sixty-three diabetic patients (63%) had erectile dysfunction compared to 6 nondiabetic subjects (12%). Mild, mild-to-moderate, moderate, and severe erectile dysfunction among these patients were 17 (27%), 21 (33.3%), 16 (25.4%), and 9 (14.3%), respectively. Among diabetics, erectile dysfunction was significantly associated with age, obesity, glycated hemoglobin level, duration of diabetes, presence of hypertension, dyslipidemia, and neuropathy (p= 0.001, 0.005, <0.001, 0.038, 0.02, 0.017 and 0.025 respectively)

**Conclusion:** Erectile dysfunction was significantly more prevalent in patients with type 2 diabetes than in non-diabetic patients. Being older than 50 years old, obesity, glycated hemoglobin level higher than 9, diabetes more than 10 years duration, presence of hypertension, dyslipidemia, and neuropathy were significantly in favor of a higher prevalence of erectile dysfunction in this group.

Key words: Diabetes; Erbil-Iraq; Erectile dysfunction; Prevalence; Risk factors.

## Introduction

Erectile dysfunction (ED) is a sustained inability to have and keep an erection enough to allow the sexual activity to be satisfactory. Available studies reveal that ED is highly prevalent worldwide; it affects multiple aspects of wellbeing and can have a serious adverse effect on both the patients and their partners. Based on the variation in the study methods, cultural differences, and the description of ED, large variations in the prevalence of ED

are being identified.<sup>6</sup> The prevalence of ED is anywhere from 35% to 75% in several cross-sectional publications.<sup>7-8</sup> Notable findings have been published in two landmark articles, the Massachusetts Male Ageing Study (MMAS) from the United States (USA) and the European Male Ageing Study (EMAS) from Europe.<sup>3, 9</sup> About 52% had an occurrence of mild to moderate ED in males aged 40-70 years, and this was significantly

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associated with age, health, and emotional condition, as stated by the MMAS. 10-11 Whereas. based on different subgroups, **EMAS** multicenter the population-based research for male patients 40-79 years of age recorded an incidence of 6-64 percent and an overall prevalence of 30 percent.<sup>9, 11</sup> Diabetes mellitus (DM), on the other hand, is a metabolic disorder associated with many chronic complications including sexual disorders.<sup>12</sup> Current data shows that DM seems to reach a pandemic level globally. As of 2019, nearly 463 million adults (20-79 years) were having diabetes; by 2045 this will go up to 700 million.<sup>13</sup> Furthermore, the main burden and the

## **Patients and Methods**

This observational cross-sectional study had been done at the endocrinology outpatient at Erbil teaching hospital which is located in the Kurdistan Region at the north of Iraq. Recruitment was for six months between June and December 2021. We employed a convenience sampling method. Men attending the endocrinology outpatient at Erbil teaching hospital during the study period were recruited if met the inclusion criteria. The patients included in this study must meet the following inclusion criteria: (1) male patients; (2) diagnosed with type 2 DM (as per the Diabetes Association American definitions); <sup>18</sup> (3) aged 20 years and more; and (4) being married. The exclusion criteria were: (1) pathological anomalies of the genitals that might affect erection (e.g., Peyronie disease); (2) type 1 diabetes; (3) a history of pelvic or gonadectomy operation; (4) organ transplantation history; (5) known or suspected chronic disabling disease such as chronic renal failure, chronic hepatic failure, chronic heart failure and chronic obstructive pulmonary disease; (6) a primary diagnosis (e.g., hyposexuality) of concomitant sexual dysfunction (such is primary hypogonadism); and (7) Drug abuse history over the past 12 months. In total, largest increase seems to be in the developing countries. About 80% of diabetic patients were discovered to be the residents of low- and middle-income countries. 13 Diabetes typically causes multiple variants of sexual dysfunction, mainly ED, premature ejaculation and diminished libido, occasionally delayed/prolonged ejaculation. 14-15 most common sexual disorder associated with DM is erectile dysfunction. 12, 16-17 The aim of this study is to find out the prevalence of ED in patients with type 2 DM and its relation to several contributing factors. More attention will be given to certain risk factors to assess significance in developing ED.

180 male participants had been invited to this study; however 150 participants were included in this study. Of them, 100 were suffering from Diabetes and considered as patients group and 50 age-matched disease-free participants who were named control group (non-diabetic). Diabetes period in this study is determined by the date of onset of the disease. The BMI was classified into: Normal=18-24.9 kg/m2, Overweight >25-29.5 kg/m2 and Obese > 30 kg/m<sup>2</sup>. Smoking was classified into never-smoker (including smoker), or current smoker. We recorded glycosylated hemoglobin A1c (HbA1c) levels assessed in the last 3 months. DM therapies were listed as oral anti-diabetic drug (OAD) or insulin (alone or in combination with OAD). The presence of hypertension was assessed by history and hypertension treatment was classified into 1/ Beta Blocker. 2/Angiotensin converting enzyme inhibitor (ACEi) or angiotensin receptor blocker (ARB). 3/Others. Ankle systolic blood pressure (BP) was measured and its ratio to the systolic BP of the arm was calculated according to a standardized method to find the ankle brachial pressure index (ABI) and was used to evaluate the presence of peripheral arterial disease (PAD). A value of 0.9 or less in either legs is a agreed to be diagnostic for PAD. Dyslipidemia was defined by the presence of any of the following: total cholesterol >4.5 mmol/L, low density lipoprotein >2.6 mmol/L, triglycerides >1.7 mmol/L, high density lipoprotein <1.1 mmol/L, or nonhigh-density lipoprotein >3.4 mmol/L. Total serum testosterone levels were considered to be normal (>10 nmol/L) or low (<10 nmol/L), depending on the ranges. 19 reference Thyroid agreed Stimulating Hormone TSH: was classified into Normal (0.4-4 IU/ml) or High (>4 IU/ml). Prolactin Hormone also was classified into either Normal (< 20 mcg/L) or High (> 20 mcg/L). Neuropathy (by neurological examination) and nephropathy testing (by for microalbuminuria or elevated levels for creatinine) were investigated by history, examination, and laboratory testing. Educational level of the participants was classified into Primary (elementary), Secondary (Intermediate and Preparatory) other (higher education, or graduation). Alcohol drinking classified into either yes (regardless of amount, frequency, or type) or No. Due to participants' very low responsiveness screening for depression was omitted from the study. ED was evaluated using a faceto-face interview approach implementing the five questions of the International Index of Erectile Function Questionnaire

### Results

We recruited 100 participants with type 2 diabetes Miletus ,with mean age of 51.6±11.4 years old, 54% were above 50 years, 32% were obese, more than 40% were with HbA1c above 9, more than 50% were with DM Duration above 10 years, 86% were receiving oral hypoglycemic agents, 53% with hypertension 19% receiving Beta Blocker medications,43%

(IIEFQ).20 IIEFQ-5 is scored on a Likert scale (0-5), in which a higher score suggests better sexual function. Sexual dysfunction severity was divided into five classes according to the total score (i.e., severe 5-7, moderate 8-11, mild to moderate 12-16, mild 17-21 and erectile dysfunction 21-25). Data was analyzed using SPSS (Statistical Package for Social Scientists) version 25.0 for Windows (Chicago, Illinois, Descriptive statistics in terms of mean and standard deviation (SD) was computed for the continuous variables. While descriptive statistics consisting of frequencies and percentages (%) was computed and level of significance was investigated using the Pearson Chi-square test to analyze the association between categorical data. A pvalue  $\leq 0.05$  was regarded as significant. All patients involved in our study were invited to join voluntarily. We obtained informed consent from our subjects based on a thorough clarification of the study's goals and procedures. This study was carried out in agreement with the ethical standards set out at the Helsinki Declaration and was accepted by Hawler Medical University's ethics committee. Respondents were further assured of confidentiality and anonymity. All participants were informed that they can refuse to participate and / or withdraw from this research.

were currently smokers,48% had dyslipidemia ,74% were of secondary education, 28% with peripheral arterial disease , 47% suffering from neuropathy, 39% from Nephropathy,6% were alcohol users, 9% with high prolactin level, 16% with low testosterone levels and 15% were with high TSH levels Table (1).

**Table** (1): Diabetic Participants' Distribution according to study Variables

Study Variable	Percentage					
Age	>50 years	(54)54%	≤50 Years old	(46)46%		
BMI	Normal	(33)33%	Overweight	(35)35%	Obese	(32)32%
HbA1C	<7	(39)39%	7-9	(20)20%	>9	(41)41%
DM Duration	>10 Years	(54)54%	≤10 Years	(46)46%		
DM Treatment	Oral Agents	(86)86%	Insulin	(14)14%		
Hypertension	Present	(53)53%	Absent	(47)47%		
Hypertension Treatment	Beta Blocker	(19)19%	ACEi or ARB	(17)17%	Other	(17)17%
Currently Smoker	Smoker	(43)43%	Non	(57)57%		
Dyslipidemia	Present	(48)48%	Absent	(52)52%		
ABI	Present	(28)28%	Absent	(72)72%		
Neuropathy	Present	(47)47%	Absent	(53)53%		
Nephropathy	Present	(39)39%	Absent	(61)61%		
Education	Primary	(17)17%	Secondary	(74)74%	Other*	(9)9%
Prolactin Hormone	High	(9)9%	Normal	(91)91%		
Testosterone Hormone	Low	(16)16%	Normal	(84)84%		
TSH	High	(15)15%	Normal	(85)85%		
Alcohol drink	Drinker	(6)6%	Nondrinker	(94)94%		

<sup>\*:</sup> other: higher education including undergraduate and postgraduate

We further recruited 50 diabetic free participants as control group. Their

demographic features are shown in Table (2).

**Table (2):** Control Participants' Distribution according to study Variables

Study Variable	Percentage					
Age	>50 years	(22)44%	≤50 Years old	(28)56%		
BMI	Normal	(27)54%	Overweight	(13)26%	Obese	(10)20%
Hypertension	Present	(18)36%	Absent	(32)64%		
Hypertension Treatment	Beta Blocker	(4)8%	ACEi or ARB	(8)16%	Other	(6)12%
Currently Smoker	Smoker	(30)60%	Non	(20)40%		
Dyslipidemia	Present	(12)24%	Absent	(38)76%		
ABI	Present	(6)12%	Absent	(44)88%		
Neuropathy	Present	(4)8%	Absent	(46)92%		
Nephropathy	Present	(2)4%	Absent	(48)96%		
Education	Primary	(15)30%	Secondary	(24)48%	Other	(11)22%
Prolactin Hormone	High	(2)4%	Normal	(48)96%		
Testosterone Hormone	Low	(4)8%	Normal	(46)92%		
TSH	High	(1)2%	Normal	(49)98%		
Alcohol drink	Drinker	(3)6%	Nondrinker	(47)94%		

The study showed that ED was significantly higher (p =<0.001 using Pearson Chi-square) among Diabetic

group 63% Vs 12% among control (Nondiabetic) groups Table (3)

**Table (3):** Erectile dysfunction distribution among study groups

Study Variable	ED	Not ED	P Value (Pearson Chi-square)
DM	(63) 63%	(37) 37%	<0.001
Not DM	(12) 12%	(88) 88%	<0.001

In this study diabetic participant were distributed according to ED severity into

23% mild, 60% moderate and 12% were severe Table (4).

**Table (4):** Diabetic Participants' Distribution according to ED severity

Erectile Dysfunction Severity	No.	%
Mild	17	27.0
Mild to Moderate	21	33.3
Moderate	16	25.4
Sever	9	14.3

There was a statistically significant association between increase in age and ED presence (p=0.001), being older than 50 was associated with higher ED prevalence. Also, a statistically significant association was found between ED and

BMI classification, Obese participants were with higher chance to develop ED (p=0.005). No statistical difference was found between ED presence and smoking, educational levels, or Alcohol use (p=0.71, 0.23 and 0.64 respectively) Table (5).

**Table (5):** Distribution according to Sociodemographic Variables among Diabetic Patients

Sociodemographic Variable		ED	No ED	P Value (Pearson Chi- square)	
	> 50 Years	(42) 77.80%	(12) 22.20%	0.001	
Age	≤ 50 Years	(21) 45.70%	(25) 54.30%	0.001	
	Obese	(26) 81.30%	(6) 18.80%		
BMI	Overweight	(23) 65.70%	(12) 34.30%	0.005	
	Normal	(14) 42.40%	(19) 57.60%		
Currently Smoker	Yes	(28) 65.10%	(15) 34.90%	0.703	
	No	(35) 61.40%	(22) 38.60%		
	Primary	(13) 76.50%	(4) 23.50%		
Education	Secondary	(43) 58.10%	(31) 41.90%	0.231	
	Other	(7) 77.80%	(2) 22.20%		
Alcohol user	Yes	(4) 66.7	(2) 33.3	0.847	
	No	(59) 62.8	(35) 37.2	0.047	

There was a strong association between ED and HbA1c levels, higher HbA1c (>9) was in favor of higher ED presence (p<0.001). Furthermore, the prevalence of ED was significantly associated with a longer duration of DM (p=0.038). Also, ED prevalence was significantly higher in diabetic patients with hypertension than in non-hypertensive patients (p=0.02). Plus, the presence of dyslipidemia in diabetic patients was again significantly associated

with higher ED than those who did not have dyslipidemia (p=0.017). ED prevalence was significantly higher in Diabetic patients with neuropathy than those with no neuropathy (p=0.02). |on the other hand, there was no significant relationship between the prevalence of ED and type of DM treatment, type of antihypertensive medication, PAD, or nephropathy among diabetic patients

(P=0.63, 0.13, 0.276 and 0.06 respectively) Table (6). **Table (6):** Distribution according to chronic illnesses and complication Among Diabetic Patients

Study Variable		ED	No ED	P Value (Pearson Chi- square)	
	>9	(35) 85.40%	(6) 14.60%	•	
HbA1C	7-9	(8) 40.00%	(12) 60.00%	< 0.001	
	<7	(20) 51.30%	(19) 48.70%		
DM Duration	>10 Years	(39)72.20%	(15) 27.80%	0.038	
DIVI Duration	≤ 10 Years	(24) 52.20%	(22) 47.80%	0.038	
DM Treatment	Insulin	(8) 57.10%	(6) 42.90%	0.624	
DM Treatment	Oral Hypo.	(55) 64.00%	(31) 36.00%	0.624	
Hypertension	Yes	(39) 73.60%	(14) 26.40%	0.019	
Trypertension	No	(24) 51.10%	(23) 48.90%		
Hypertension treatment	B Blocker	(14) 73.70%	(5) 26.30%	0.135	
	ACE or ARB	(12) 70.60%	(5) 29.40%		
	Other	(13) 76.50%	(4) 23.50%		
Dyelinidamia	Yes	(36) 75.00%	(12) 25.00%	0.017	
Dyslipidemia	No	(27) 51.90%	(25) 48.10%	0.017	
PAD	Yes	(20) 71.40%	(8) 28.60%	0.276	
	No	(43) 59.70%	(29) 40.30%	0.276	
Neuropathy	Yes	(35) 74.50%	(12) 25.50%	0.025	
	No	(28) 52.80%	(25) 47.20%	0.023	
Nephropathy	Yes	(29) 74.40%	(10) 25.60%	0.0599	
	No	(34) 55.70%	(27) 44.30%	0.0399	

No statistical difference was found in the prevalence of ED among diabetic patients and levels of prolactin, TSH or

Testosterone levels (p=0.44, 0.155 and 0.082 respectively) Table (7).

**Table (7):** Distribution according to hormones' level of Diabetic Patients

Hormone		ED	No ED	P Value (Pearson Chi-square)	
Prolactin	High	(5) 55.60%	(4) 44.40%	0.44	
Profactifi	Normal	mal (58) 63.70% (33) 36.30%		0.44	
TSH	High	(7) 46.70%	(8) 53.30%	0.155	
1311	Normal	(56) 65.90%	(29) 34.10%	0.133	
Tastastarona	Low	(7) 43.80%	(9) 56.30%	0.082	
Testosterone	Normal	(56) 66.70%	(28) 33.30%	0.082	

## **Discussion**

In order to keep marital peace and happiness, healthy sexual functioning is necessary. Diabetes mellitus can cause normal sexual function to be disrupted in both men and women as a result of diabetic-induced end organ damage and psychological stress. This study showed

that the prevalence of ED was significantly higher among diabetic group versus non-diabetic group (63% VS 12 %) with p <0. 001. This finding was in line with several studies from USA and Egypt. 21-23 This study showed that being older in age was significantly associated with ED (p

=0.001). A cross-sectional survey of 400 men confirmed Malaysian conclusion.<sup>24</sup> Due to alterations caused by atherosclerosis and the resulting decreased blood supply to the genitalia, aging posed a substantial risk for developing ED. Significantly higher prevalence of ED was detected among obese group (p= 0.005), which was consistent with Polish research that revealed that a BMI of over 30 kg/m2 was associated with a threefold increased probability of sexual dysfunction.<sup>25</sup> For obese patients, erectile dysfunction is caused by a number of issues that are common in people who have a lot of fat tissue, such as cardiovascular disease, diabetes, and dyslipidemia. This study revealed no association between ED and level of education (p=0.23) which was in contrast to two studies from Malaysia which showed men with secondary education were more prone to develop ED compared to those with tertiary education.<sup>24, 26</sup> The disparity in sample size and sampling procedure might explain the discrepancy, especially as most of the persons sampled in our study were the visitors to public hospitals and most were lower socioeconomic educational level. This study yielded in no significant difference between smoking status and ED (p=0.71) however this finding contradicted was by Massachusetts Male Aging Study, where cigarette smokers were found to have a 1.97 times higher risk to develop ED.<sup>27</sup> In Finland, 1130 men between the ages of 50 and 70 were tracked for ten years in a study comparable to the Massachusetts Male Health Study, the odds-ratio in this study was 1.4. however it did not meet statistical significance according to the author of the mentioned study.<sup>28</sup> This disparity might be explained by the varied methodologies utilized in those studies, as cross section studies in general may not be able to detect causal and temporal links between cause and effect. According to the Boston Area Community Health Survey, the risk of having ED became considerable

only after 20 pack-years. 29 This study showed no statistical association between ED and alcohol consumption (p=0.64), while a Chinese meta-analysis crosssectional study indicated that small to moderate alcohol intake (up to drinks/week) was found to be associated probability lower of dysfunction (OR = 0.71, P = 0.000). A non-linear association was discovered between the risk of ED and alcohol intake, meta-analysis.<sup>30</sup> dose-response in Possible explanation of this disagreement is that being alcoholic is a social stigma among Iraqi people and a lot of individuals hide the fact of drinking alcohol. Binge drinking has been shown to affect microvascular and macrovascular function, which might indicate early signs of risk.<sup>31</sup> cardiovascular Epidemiologic research shows that consuming alcoholic beverages at low levels (1-2 drinks per day) on a regular basis may reduce the incidence of unfavorable cardiovascular events.<sup>32</sup> This study found a clear positive relationship between ED and both HbA1c level, and DM duration (p=0.001,0.038, respectively), which was supported by studies from the United Kingdom (Five cross-sectional studies including 3299 patients).<sup>33</sup> Available data has revealed a link between erectile dysfunction and glycemic management, as well as the duration of diabetes.<sup>33, 34</sup> Diabetic males have a nearly threefold increased risk of developing ED when compared to nondiabetics in similar studies and they are also more likely to acquire ED 10 to 15 years before non-diabetics.<sup>3</sup> This study revealed a significant association between hypertension and ED on one side and dyslipidemia and ED on the other side (p=0.02)and 0.017 respectively). Comparable results were found within an Italian study of 555 men <sup>17</sup>. However, studies from China and USA showed no significant association between serum lipid and the risk of ED, pointing to the apparent conclusion that dyslipidemia does not have a significant role in the

probability of developing ED diabetics. <sup>34, 35</sup> The current study's findings, on the other hand, revealed no statistically significant differences in ED across various antihypertensive groups. This might be due to the fact that many patients were using more than one kind of antihypertensive medication, which could lead to confounding results if some medicines have a neutral/positive effect on the ED while others have unfavorable effects. Some earlier antihypertensive classes (beta blockers, diuretics) have a history of causing erectile dysfunction. whereas other medicines (ACEIs, calcium channel blockers) appear to be neutral. Furthermore, data shows that angiotensin receptor blockers may improve erectile function.<sup>36</sup> A significant increase in ED was reported among diabetic individuals with neuropathy (p=0.025), while no such significant increase was detected among diabetic participants with nephropathy (p=0.06) in this research. Similar findings were found in a Turkish, Japanese multicenter cross-sectional study, Romanian<sup>38</sup> and Indian studies.<sup>39</sup> This research found no link between the presence of peripheral artery disease and ED (p=0.27), however a study from the United States of 690 males found a favorable link <sup>40</sup>. This discrepancy might be related to differences in research duration and sample characteristics. Several atherosclerotic risk factors can lead to occlusive disorder in various arteries, eventually leading to some degree of vascular ED. Reduced neuronal or endothelial NO, as well as prolonged tissue ischemia; can result in decreased cavernosal smooth muscle relaxation. DM treatment showed no significant difference for ED prevalence (p=0.63). Same results were found with a Japanese 37 and a Chinese study. The level of serum prolactin had no effect on ED (p=0.44),

## **Study Limitation**

We are aware of several limitations of our study. First off, since this study is crossaccording to this study. In contrast to research from China, which found a deleterious impact on erectile function. 41 our study found no such effect. This might be disparity explained differences in research duration and sample size, especially as the current investigation focused solely on diabetes patients rather than the general population. The frequency of hyperprolactinemia was 9% in the study group, with most of them having just a mild (less than twice) rise in serum prolactin. This study showed no significant association between ED and testosterone levels (p=0.082). This finding contradicted other studies which showed lower levels of serum testosterone were associated significantly with higher prevalence of ED in men with Type 2 DM. A research done on 198 men with type 2diabetes had showed that the presence of ED was significantly associated with both of serum testosterone.<sup>42</sup> levels Different methodologies and inclusion criteria might be the cause of the disparity, as one of the exclusion criteria in our study was a documented history of diagnosed primary hypogonadism. Finally, there was no association between TSH levels and ED in this investigation (p=0.155), however this contradicted an Italian multicenter prospective analysis of 48 males at endocrinology and andrology clinics in university hospitals, which revealed hypothyroidism nearly tripled occurrence of ED.<sup>43</sup> In hypothyroid males, Krassas et al. found a considerably higher prevalence of ED than in controls (p 0.0001).<sup>44</sup> This discrepancy could be explained by the participants' different characteristics, and by the fact that this study investigated the presence of thyroid disorders in diabetic patients with ED, whereas most other studies looked into the presence of sexual disorders in those with thyroid dysfunction.

sectional in nature, we cannot be certain that ED and the study factors are related in

ED.

a causal way. Furthermore, because the sample was drawn from a single center, we were unable to generalize this information across the country. Finally, no information

**Conclusion and Recommendation** 

Erectile dysfunction was significantly more prevalent in patients with type 2 diabetes than non-diabetic patients. Being older than 50 years old, obesity, HBA1c level higher than 9, DM more than 10 years duration, presence of hypertension, dyslipidemia and neuropathy were significantly in favor of higher prevalence of ED in this group. Considering the

increased frequency of ED in type 2 diabetes patients, physicians should focus on early detection and treatment of ED in diabetic men. Furthermore, maintaining effective glycemic management, avoiding DM-related comorbidities, and managing obesity, hypertension and dyslipidemia may lower the chance of developing ED.

was gathered about the individuals' social

situation or partner, two factors that could

theoretically influence the prevalence of

#### **Conflicts of interest**

The author reports no conflicts of interest.

#### References

- 1. Xu, Y., Zhang, Y., Yang, Y. et al. Prevalence and correlates of erectile dysfunction in type 2 diabetic men: a population-based cross-sectional study in Chinese men. Int J Impot Res 2019; 31, 9–14.
- 2. Anonymous. NIH consensus conference. Impotence. NIH consensus development panel on impotence. JAMA. 1993;(270):83-90.
- 3. Feldman HA, Goldstein I, Hatzichristou DG, et al. Impotence and its medical and psychosocial correlates: results of the Massachusetts Male Aging Study. J Urol. 1994 1; 151(1): 54-61.
- 4. Fisher WA, Eardley I, McCabe M, et al. Erectile dysfunction (ED) is a shared sexual concern of couples I: couple conceptions of ED. J Sex Med. 2009;6(10):2746-60.
- 5. Salonia A, Castagna G, Saccà A, et al. Is erectile dysfunction a reliable proxy of general male health status? The case for the International Index of Erectile Function—Erectile function domain. The Journal of Sexual Medicine. 2012 1;9(10):2708-15.
- 6. Melman A, Gingell JC. The epidemiology and pathophysiology of

- erectile dysfunction. The Journal of urology. 1999 Jan;161(1):5-11.
- 7. Cander S, Coban S, Altuner S, et al. Prevalence and correlates of erectile dysfunction in type 2 diabetes mellitus: a cross-sectional single-center study among Turkish patients. Metabolic syndrome and related disorders. 2014 Aug 1;12(6):324-9.
- 8. Dan A, Chakraborty K, Mondal M, et al. Erectile dysfunction in patients with diabetes mellitus: Its magnitude, predictors and their bio-psycho-social interaction: A study from a developing country. Asian journal of psychiatry. 2014;7(1):58-65.
- 9. Corona G, Lee DM, Forti G, et al. Agerelated changes in general and sexual health in middle-aged and older men: results from the European Male Ageing Study (EMAS). The journal of sexual medicine. 2010 Apr 1;7(4):1362-80.
- 10. Giuliano FA, Leriche A, Jaudinot EO, de Gendre AS. Prevalence of erectile dysfunction among 7689 patients with diabetes or hypertension, or both. Urology. 2004 Dec 1;64(6):1196-201.
- 11. Nutalapati S, Ghagane SC, Nerli RB, Jali MV, Dixit NS. Association of erectile dysfunction and type II diabetes mellitus

- at a tertiary care centre of south India. Diabetes & Metabolic Syndrome: Clinical Research & Reviews. 2020 Jul 1;14(4):649-53.
- 12. Olamoyegun MA, Ala OA, Fagbemiro EY. Frequency and Determinants of Erectile Dysfunction among Type 2 Diabetes Patients in a Tertiary Health Institution in Nigeria. Journal of Advances in Medicine and Medical Research. 2020; 32(23):139-47.
- 13. Atlas D. International diabetes federation. IDF Diabetes Atlas, 7th edn. Brussels, Belgium: International Diabetes Federation. 2015: 10-17.
- 14. Sharifi F, Asghari M, Jaberi Y, Salehi O, Mirzamohammadi F. Independent predictors of erectile dysfunction in type 2 Diabetes mellitus: Is it true what they say about risk factors? ISRN Endocrinol. 2012;2012:502353.
- 15. Díaz-Díaz E., León M. C., Arzuaga, et al. Erectile Dysfunction: A Chronic Complication of the Diabetes Mellitus. In: Nunes, K. P., editor. Erectile Dysfunction -Disease Associated Mechanisms and Novel Insights into Therapy [Internet]. London: IntechOpen; 2012 [cited 2022 May 20]. Available from: https://www.intechopen.com/chapters/30219 doi: 10.5772/31005.
- 16. Johannes CB, Araujo AB, Feldman HA, et al. Incidence of erectile dysfunction in men 40 to 69 years old: longitudinal results from the Massachusetts male aging study. The Journal of urology. 2000 Feb;163(2):460-3.
- 17. Giugliano F, Maiorino M, Bellastella G, et al. Determinants of erectile dysfunction in type 2 diabetes. International Journal of Impotence Research. 2010 May;22(3):204-9.
- 18. Anonymous. American Diabetes Association. Diagnosis and classification of diabetes mellitus. Diabetes care. 2014 Jan 1;37(1):81-90.
- 19. Nieschlag E, Swerdloff R, Behre HM, et al. Investigation, treatment and monitoring of late-onset hypogonadism

- in males. The Aging Male. 2005 Jun 1;8(2):56-8.
- 20. Rosen RC, Cappelleri JC, Smith MD, Lipsky J, Pena BM. Development and evaluation of an abridged, 5-item version of the International Index of Erectile Function (IIEF-5) as a diagnostic tool for erectile dysfunction. International journal of impotence research. 1999 Dec;11(6):319-26.
- 21. Selvin E, Burnett AL, Platz EA. Prevalence and risk factors for erectile dysfunction in the US. Am J Med. 2007 Feb:120(2):151-7.
- 22. Rosen RC, Wing R, Schneider S, Gendrano N 3rd. Epidemiology of erectile dysfunction: the role of medical comorbidities and lifestyle factors. Urol Clin North Am. 2005 Nov;32(4):403-17.
- 23. El-Latif MA, Makhlouf AA, Moustafa YM, et al. Diagnostic value of nitric oxide, lipoprotein(a), and malondialdehyde levels in the peripheral venous and cavernous blood of diabetics with erectile dysfunction. Int J Impot Res. 2006 Nov-Dec;18(6):544-9.
- 24. Nordin RB, Soni T, Kaur A, Loh KP, Miranda S. Prevalence and predictors of erectile dysfunction in adult male outpatient clinic attendees in Johor, Malaysia. Singapore Med J. 2019 Jan;60(1):40-7.
- 25. Skrypnik D, Bogdański P, Musialik K. Obesity--significant risk factor for erectile dysfunction in men. Pol Merkur Lekarski. 2014 Feb;36(212):137-41.
- 26. Momtaz YA, Hamid TA, Ibrahim R, Akahbar SA. Racial and socioeconomic disparities in sexual activity among older married Malaysians. Archives of gerontology and geriatrics. 2014 Jan 1;58(1):51-5.
- 27. Kovac JR, Labbate C, Ramasamy R, Tang D, Lipshultz LI. Effects of cigarette smoking on erectile dysfunction. Andrologia. 2015 Dec;47(10):1087-92.
- 28. Shiri R, Hakama M, Häkkinen J, et al. Relationship between smoking and erectile dysfunction. International journal

- of impotence research. 2005 Mar;17(2):164-9.
- 29. Kupelian V, Link CL, McKinlay JB. Association between smoking, passive smoking, and erectile dysfunction: results from the Boston Area Community Health (BACH) Survey. European urology. 2007 Aug 1;52(2):416-22.
- 30. Wang XM, Bai YJ, Yang YB, et al. Alcohol intake and risk of erectile dysfunction: a dose-response meta-analysis of observational studies. Int J Impot Res. 2018 Nov;30(6):342-51.
- 31. Goslawski M, Piano MR, Bian JT, et al. Binge drinking impairs vascular function in young adults. Journal of the American College of Cardiology. 2013 Jul 16;62(3):201-7.
- 32. Leong DP, Smyth A, Teo KK, et al. Patterns of alcohol consumption and myocardial infarction risk: observations from 52 countries in the INTERHEART case—control study. Circulation. 2014 Jul 29;130(5):390-8.
- 33. Binmoammar TA, Hassounah S, Alsaad S, Rawaf S, Majeed A. The impact of poor glycaemic control on the prevalence of erectile dysfunction in men with type 2 diabetes mellitus: a systematic review. JRSM Open. 2016 Feb 12;7(3):2054270415622602.
- 34. Lu CC, Jiann BP, Sun CC, et al. Association of glycemic control with risk of erectile dysfunction in men with type 2 diabetes. J Sex Med. 2009 Jun;6(6):1719-28.
- 35. Wing RR, Rosen RC, Fava JL, et al. Effects of weight loss intervention on erectile function in older men with type 2 diabetes in the Look AHEAD trial. The journal of sexual medicine. 2010 Jan 1;7(1):156-65.
- 36. Doumas M, Douma S. The effect of antihypertensive drugs on erectile function: a proposed management algorithm. J Clin Hypertens (Greenwich). 2006 May;8(5):359-64.

- 37. Furukawa S, Sakai T, Niiya T, et al. Diabetic peripheral neuropathy and prevalence of erectile dysfunction in Japanese patients aged <65 years with type 2 diabetes mellitus: The Dogo Study. Int J Impot Res. 2017 Jan;29(1):30-34.
- 38. Mota M, Lichiardopol C, Mota E, Pănuş C, Pănuş A. Erectile dysfunction in diabetes mellitus. Rom J Intern Med. 2003;41(2):163-77.
- 39. Meena BL, Kochar DK, Agarwal TD, Choudhary R, Kochar A. Association between erectile dysfunction and cardiovascular risk in individuals with type-2 diabetes without overt cardiovascular disease. Int J Diabetes Dev Ctries. 2009 Oct;29(4):150-4.
- 40. Polonsky TS, Taillon LA, Sheth H, et al. The association between erectile dysfunction and peripheral arterial disease as determined by screening ankle-brachial index testing. Atherosclerosis. 2009 Dec;207(2):440-4.
- 41. Xu ZH, Pan D, Liu TY, et al. Effect of prolactin on penile erection: a cross-sectional study. Asian J Androl. 2019 Nov-Dec;21(6):587-91.
- 42. Kapoor D, Clarke S, Channer KS, Jones TH. Erectile dysfunction is associated with low bioactive testosterone levels and visceral adiposity in men with type 2 diabetes. Int J Androl. 2007 Dec;30(6):500-7.
- 43. Carani C, Isidori AM, Granata A, et al. Multicenter study on the prevalence of sexual symptoms in male hypo- and hyperthyroid patients. J Clin Endocrinol Metab. 2005 Dec;90(12):6472-9.
- 44. Krassas GE, Tziomalos K, Papadopoulou F, Pontikides N, Perros P. Erectile Dysfunction in Patients with Hyper- and Hypothyroidism: How Common and Should We Treat? J Clin Endocrinol Metab 2008; 93(5): 1815–9.