

Prevalence of Diabetic Peripheral Neuropathy among Type I Diabetes – An Observational Study

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Abstract---

Objective:

The objective of this study is to find out the prevalence of diabetic peripheral neuropathy (DPN) among diabetics in Chennai.

Background of the Study: *DPN is the most common chronic and long term complications of diabetes mellitus and is considered as leading cause of mortality and morbidity rate, causing huge economic burden to the government. It is one of the serious microvascular complications of DM which has been diagnosed in 20-50% of diabetic population.*

Methodology: *This is an cross sectional study with 100 patients selected by simple random sampling method. Those who were aged more than 30 years and diagnosed positive for type I diabetes atleast 1year duration with random blood sugar >200 mg/dl as per Indian council of medical research guidelines 2005. Those who are willing to participate, who had regular follow up visits for atleast 6 months were included in this study. The exclusion criteria are who have type 1 diabetes, gestational DM and maturity onset diabetes of the young, significant musculoskeletal disorder in lower limb, who underwent amputation of whole foot, below knee and above knee amputation, peripheral neuropathy due to any other cause. The prevalence of DPN among diabetics in chennai will be found out. Pre and post test done using DN4, S-LAANS, MNSI questionnaire.*

Result: *The study was conducted in type 1 diabetes patient following results were obtained The prevalence was found using DN4, MNSI and S-LANSS questionnaire. The mean of DN4 was found to be 4.0800 and its significant with $p < 0.05$. The mean of MNSI is about 7.62000 with significance $p < 0.05$ and mean value of S-LANSS is 9.0600 with significance of $p < 0.05$. The overall prevalence of diabetic peripheral neuropathy was found to be 30%.*

Keywords--- *Diabetes, Peripheral Neuropathy, DN4, S-LAANS, MNSI.*

I. INTRODUCTION

Diabetes Mellitus is a metabolic disorder of multiple aetiology. It is characterized by chronic hyperglycemia with disturbance of carbohydrates, fat and protein metabolism which results from defects in insulin secretion, insulin action or both. The effects leads to long term damage, dysfunction and failure of various organs^[1]. The international diabetes federation has prescribed that the number of people with diabetes in southeast Asia region which was 71.4 million in 2011 and will increase 12.9 million by 2030^[2] and 60% of the world's are diabetic population resides in

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Asia^[3]. India has one of the highest prevalence of type 2 diabetes Mellitus in the world^[4]. Diabetes is the 5th leading cause for mortality which accounts for 3.8 million deaths annually^[5]. In 2030 diabetes will be 7th leading cause for death as predicted by world health organization^[6]. Long term elevation in blood glucose level leads to micro and macro vascular complications such as heart disease, stroke, blindness and kidney disease^[7]. The microvascular complications includes diabetic retinopathy, diabetic nephropathy and diabetic neuropathy. The macro vascular complications includes stroke, transient ischemic attack, coronary heart disease and peripheral vascular disease^[8]. By far, the most common is Diabetic Peripheral Neuropathy(DPN), which accounts for 75% of diabetic neuropathy^[9]. DPN is the most common chronic and long term complications of diabetes mellitus and which is considered as leading huge economic burden to the government^[10]. DPN is one of the serious micro vascular complications of both type 1 and type 2 diabetes mellitus which has been diagnosed in 20-50% of diabetic population^[9].

Approximately 15% of people with diabetes develop foot ulcers once in their lifetimes in which 60-70% are neuropathic in origin^[11]. This increase is due to chronicity of disease and poor glycemic control/. 15-25% of diabetic neuropathy are painful^[12]. A significant proportion of patients with peripheral neuropathy have no underlying cause. Multiple studies have implicated the metabolic syndrome as a potential cause^[13]. DPN is a progressive degeneration of peripheral nerves from distal to proximal leads to wide range of neuropathic symptoms such as numbness, burning, prickling, tingling, sharp pain or cramps, extreme sensitivity to touch, allodynia and loss of balance and coordination^[14]. The primary symptoms of diabetic peripheral neuropathy is abnormal or loss of sensation in toes, which extends to involve feet and legs in stocking distribution^[15]. The patient with DPN has limitations in activity of daily living. Because of;'] decreased sensation or weakness and pain particularly in hands and feet^[16].

High risk individual are those above 45 years, hypertensive people, obesity, family history of diabetes, dislipidemia. Adverse effects of peripheral neuropathy are due to poor foot hygiene, improper foot wear and frequent barefoot walking. In such circumstance, complications of foot infection and gangrene are common cause of hospital admission^[1].

Painful diabetic peripheral neuropathy has a significant impact on patient's quality of life, as it is accompanied by depression, anxiety and sleep disturbance^[17]. On the other hand this increase morbidity and mortality, where society undergoes extreme burden due to health care cost^[18]. After macro vascular disease and diabetic neuropathy, DPN is ranked third among the complications of DM which cause great life time expenditure of resources. But early detection and efficient glucose control leads to prevention of clinical neuropathy and its associated complications^[19].

The findings of S-LANSS score confirmed that S-LANSS scale as a valid and reliable self – report instrument for identifying neuropathic pain and is also acceptable for use in postal survey research^[20]. An analysis for MNSI confirm that MNSI is a simple, non-invasive and valid measure of distal symmetrical peripheral neuropathy when compared with gold standard diagnostic testing^[21]. A research on DN4 questionnaire concludes Turkish version of DN4 is reliable, valid, short and quick screening tool in identification of neuropathic pain patients^[22].

II. METHODOLOGY

This cross sectional study was carried out in ACS medical college -velapanchavadi over the period of 3 months using simple random sampling method. A total of 100 patients were randomly selected. Those who were aged more than 30 years, diagnosed positive for type I diabetes atleast 1 year duration with random blood sugar more than 200 mg/dl or fasting blood sugar more than 126 mg/dl as per Indian Council of medical research guidelines 2005, those who had regular follow up visits for atleast 6 months and who are willing to participate are included in this study. The exclusion criteria are those who having type 1 diabetes, gestational DM, maturity onset diabetes of the young, significant musculoskeletal disorder in lower limb, who underwent amputation of whole foot, below knee, above knee, peripheral neuropathy due to any other known cause.

The samples were fully explained about the study and the questionnaire to be filled. they were asked to fill the consent form in acceptance of participation in the study, which is duly signed by the samples.

III. PROCEDURE

Written informed consent was obtained from all study participants. Information regarding anthropometric measurement, blood pressure was collected. Additional details about investigations, complications was obtained from patients records. Data was collected from 100 study subjects. Peripheral neuropathy was assessed using LANSS, MNSI and DN4 questionnaire.

MNSI (michigan neuropathy screening instrument) has two components. First component is subjective and consist of peripheral neuropathy symptoms rated 15 self- administered questions. A higher score (total score 13) indicates higher neuropathic symptoms. The second component of the MNSI is a brief physical examination involving a) inspection of the feet b) semi- quantitative assessment of vibration sensation at the dorsum of great toe c) grading of ankle reflexes and d) semmes-weinstein monofilament testing. Patients are considered neuropathic if they screen positive ($>_2.5$ on a 10 point scale) on the second component of MNSI.

LANSS (the leeds assessment of neuropathic symptoms and signs) pain scale used to determine whether the nerves carrying pain signals are working or not. This also determines quality of pain, skin appearance, temperaturw and sensitivity. It also includes test for examining allodynia and pin prick threshold. Total score ranges from 0 to 24 where a score $>_12$ indicated presence of neuropathic pain.

DN4 (neuropathic pain diagnostic questionnaire) consist of four question 1 and 2 which were based on the interview of the patient, 3 and 4 based on standardized clinical examination. Question 1 have five Items related to pain, question 2 have four items related to parasthesia or dysesthesia, question 3 has four items related to sensory deficit and question 4 has four items related to evoked pain

IV. DATA ANALYSIS

DN4 (DouleurNeuropathique – 4):

One-Sample Statistics				
	N	Mean	Std. Deviation	Std. Error Mean
DN4	100	4.0800	2.40236	.24024

One-Sample Test						
	Test Value = 0					
	t	df	Sig. (2-tailed)	Mean Difference	95% Confidence Interval of the Difference	
					Lower	Upper
DN4	16.983	99	.000	4.08000	3.6033	4.5567

Test Distribution is Normal ($p < 0.05$)

MNSI (Michigan Neuropathy Screening Instrumentation):

One-Sample Statistics				
	N	Mean	Std. Deviation	Std. Error Mean
MNSI	100	7.6200	4.64938	.46494

One-Sample Test						
	Test Value = 0					
	t	df	Sig. (2-tailed)	Mean Difference	95% Confidence Interval of the Difference	
					Lower	Upper
MNSI	16.389	99	.000	7.62000	6.6975	8.5425

Test Distribution is Normal ($p < 0.05$)

S-LANSS (Leeds Assessment of Neuropathic Symptoms and Signs):

One-Sample Statistics				
	N	Mean	Std. Deviation	Std. Error Mean
LANSS	100	9.0600	6.53633	.65363

One-Sample Test						
	Test Value = 0					
	t	df	Sig. (2-tailed)	Mean Difference	95% Confidence Interval of the Difference	
					Lower	Upper
LANSS	13.861	99	.000	9.06000	7.7630	10.3570

Test Distribution is Normal ($p < 0.05$)

V. RESULT

The study was conducted in type 1 diabetes patient following results were obtained: The prevalence was found using DN4, MNSI and S-LANSS questionnaire. The mean of DN4 was found to be 4.0800 and its significant with $p < 0.05$. The mean of MNSI is about 7.62000 with significance $p < 0.05$ and mean value of S-LANSS is 9.0600 with significance of $p < 0.05$. The overall prevalence of diabetic peripheral neuropathy was found to be 30%.

VI. CONCLUSION

This study concluded that prevalence of DPN among diabetic population is about 30%. This study also highlighted severity of pain and difficulties in ADL in diabetic patients. This study formed basis for future researches

VII. DISCUSSION

Maryam Khan Kundi et al., reported that the study showed the severity of DPN to be mild with a frequency of 11.1 % moderate with a frequency of 22.0% and senesce with a frequency of 6.5 % respectively and diabetic

peripheral neuropathy is present in majority of the diabetic population of Hayatabad Peshawar^[23]. Ahlam A. Amour et al reported that DPN is widely prevalent in our setting occurring in more than half of the patients attending the diabetic clinic with more than a half experiencing the severe form^[24]. Department of non-communicable diseases et al., reported that DPN was present among 21.1 % of T2 Dm subjects in Bangladesh both men & women irrespective of urban or rural origin suffer from neuropathy which is almost similar to a study conducted among urban diabetic population by Kjersti M or Kridi in Bangladesh women are more suffering from DPN compare to men for their working pattern like almost all the women respondent were housemate^[25]. Twanakisozi et al., reported that 29.4% of recently diagnosed patient with diabetes had DPN and there is an urgent need to routinely screen for diabetes among the elderly population and examine newly diagnosed diabetic patients irrespective of their clinical symptoms^[26].

Nahlakhawaja et al reported that the overall prevalence of DPN based on MNSI was 39.5% and peripheral neuropathy is highly prevalent among Jordanian patients with type 2 diabetes mellitus^[27]. Gill HK et al., reported that the prevalence of DPN was 29.2% and detected a high frequency of DPN in newly diagnosed patients with T2DM^[28]. Javadkiani MD et al., reported that the prevalence of DPN in our studied

population was relatively high prevalence (45.7%)^[29]. Mamta Jaiswal et al., reported that the prevalence of DPN was 79 in youth with 71D and 22% in youth with T2D^[30]. S Jambart et al ; reported that the current study was designed to evaluate the prevalence of painful DPN among patients with type 1 or type 2 diabetes attending outpatient clinic in the middle east region in this setting, 53.7% of diabetic patient met the criteria for painful DPN^[31]. Prasad katulanda et al., reported that the prevalence of DPN according to the DNS and the TCSS scores was 48.1 % and 24.0% respectively^[32]

Aktarreyhanoghi D et al., reported that the MNA1 – TR is a valid and reliable method for evaluating peripheral neuropathy in Turkish speaking societies^[33]. Yuka matsuki et al., reported that the reliability and validation testing revealed that the Japanese version of questionnaire for discriminating between neuropathic and nociceptive pain in a Japanese sample^[34]. Rabiakoc et al., reported that the Turkish version of S – LANSS is a reliable and valid differential diagnostic measure of neuropathic pain in chronic pain patients^[35].

DPN is a common complication of DM. It is about prevalence of DPN among Chennai in A.C.S medical college. The prevalence of DPN by DN4, S-LANSS, and MNSI is about 30%. Descriptive statistics were calculated and is presented as mean and standard deviation. The scale shows a good significance of $p < 0.05$. The study is among referral hospital setting so the results are not generalized to the whole Chennai. The other limitations of study are small sample and study duration. Further researches were required for better knowing the influence of DPN in diabetic population.

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