

**FREQUENCY OF DIABETIC PERIPHERAL NEUROPATHY IN PATIENTS WITH NEWLY DIAGNOSED TYPE 2 DIABETES MELLITUS.**

Emaan Fatima Qureshi<sup>1</sup>, Muzaffar Ali Shaikh<sup>2</sup>, Madiha Shah<sup>3</sup>, Muhammad Hammad Jawaid<sup>4</sup>, Maarij Hassaan Qureshi<sup>5</sup>, Razia Hanif Memon.<sup>6</sup>

**ABSTRACT:**

**Introduction:** Diabetes Mellitus is a globally prevalent non-communicable disease involving hyperglycaemia and altered metabolic control, with high associated morbidities and mortalities. Diabetic neuropathies are commonly occurring complications among diabetic patients, out of which up to 50% may be asymptomatic. **Objective:** To determine the frequency of diabetic peripheral neuropathy (DPN) among patients newly diagnosed with Diabetes mellitus Type 2 **Design:** This was a hospital-based, cross sectional descriptive study to determine the frequency of DPN among patients newly diagnosed with Diabetes mellitus Type 2 **Place of Study:** Department of Medicine (OPD and inpatient), Liaquat University Hospital, Hyderabad/Jamshoro. **Duration of Study:** Six month period from 01-06-2019 to 01-12-2019 **Methodology:** Non-probability consecutive sampling was utilized for sample collection. Inclusion criteria were: Patients of both genders, between age 30 years and 60 years, newly diagnosed with Diabetes Mellitus Type 2. i.e. the fasting serum glucose level above 126 mg/dl along with HbA1c levels above 6.5% was considered as Diabetes mellitus., who were agreeable to take part in the study after informed consent. Exclusion criteria were pregnant women, patients with chronic lumbar pain, neurological findings indicative of myelopathic involvement or relevant radiculopathies, as well as patients with altered consciousness or who were inarticulate and unable to collaborate with neurological evaluation. The collected data was entered in SPSS (for version 20) and analysed accordingly. **Results:** 150 patients were recruited in this study. There were 60.7% male and 39.3% female patients. The mean age of the patients was 44.8±7.70 years. DPN frequency among patients newly diagnosed with Diabetes mellitus Type 2 was 39.3%. **Conclusion:** Our figures emphasized the necessity for rigorous programs aiming at early identification of DPN and patient education in diabetic patients with associated comorbidities such as altered lipid metabolism, cardiac dysfunction or eye involvement.

**Key Words:** Diabetic peripheral neuropathy, Type 2 Diabetes mellitus, Diabetic neuropathies

1. FCPS –II Resident, Medical Unit 3, LUMHS Jamshoro.
2. Professor and Chairman Medicine Department LUMHS Jamshoro.
3. Assistant Professor, Department Of Medical Unit 3, Liaquat University of Medical Sciences.
4. Lecturer Bilawal Medical College Jamshoro.
5. House Officer Medical Unit 1 LUMHS Jamshoro.
6. Senior Registrar Medical Unit 3. LUMHS Jamshoro.

**Corresponding Author:** Madiha Shah, Assistant Professor, Department of Medical unit 3, Liaquat University of Medical Sciences. Email; [madihashah@gmail.com](mailto:madihashah@gmail.com)

**How to cite this article:** Qureshi EF<sup>1</sup>, Shaikh MA<sup>2</sup>, Shah M<sup>3</sup>, Jawaid MH<sup>4</sup>, Qureshi MH<sup>5</sup>, Memon RH<sup>6</sup>. **FREQUENCY OF DIABETIC PERIPHERAL NEUROPATHY IN PATIENTS WITH NEWLY DIAGNOSED TYPE 2 DIABETES MELLITUS. JPUMHS;2020;10(03);18-21.**

<http://doi.org/10.46536/jpumhs/2020/10.02.2>

**INTRODUCTION**

Diabetes Mellitus is a globally prevalent non-communicable disease involving hyperglycemia and altered metabolic control, with high associated morbidities and mortalities. Currently, 415 million people are affected worldwide by Diabetes mellitus (DM) and it is expected that this number will increase to an astonishing 642 million by 2040. 1 in every 11 adults worldwide is diabetic and 50% of these adults with Diabetes mellitus are undiagnosed. As of 2015, there are currently over 7 million patients affected by Diabetes mellitus in Pakistan. Approximately 86,364 patients died of Diabetes mellitus alone in 2015 and this number is increasing<sup>1</sup>. Diabetic peripheral neuropathy (DPN) varies significantly in terms of its prevalence as per different studies, ranging from 8% to 59%<sup>2</sup>. The reported prevalence for DPN in patients newly diagnosed

with Diabetes mellitus Type 2 is 16.8%<sup>3</sup>. Certain chronic complications particularly neuropathies are the most commonly occurring in diabetic patients, out of which up to 50% may be asymptomatic. A study in the American Diabetes Association reported the DPN prevalence to be 22% in certain patients diagnosed with Type 2 DM<sup>4</sup>. DPN may manifest as existence of ulceration, absent ankle reflexes, impaired dorsal column sensation, and positive monofilament test. It aggravates potential risk of acquiring infection & foot ulcers alongside risk of non-traumatic amputations, potentially long term morbidity & disability<sup>5</sup> which is a point of concern for public health and healthcare agencies and need for advocating timely screening for DM plus associated complications and communal disease burden<sup>6</sup>. Timely diagnosis, increased patient

awareness and regular monitoring of serum glucose screening are absolutely vital <sup>7</sup>.

Broadly speaking, three components are vital in DPN management: Glycemic control, Foot care & pain relief. The American Diabetes Association issued a statement in 2005 recommending that firstly, DM patients with symptomatic DPN should have target glycemic control stabilized <sup>8</sup>. A 2012 systematic review revealed that this led to significant positive changes in proxy measures of DPN<sup>9</sup>. Additionally, patients ought to regularly examine their feet for evidence of break in the skin, any developing callus, and initial signs of inflammation around digits and nails. The physician also has a duty to carry out detailed evaluation and examination at each visit for DPN. Finally, symptomatic DPN patients should be managed in a methodical, one by one approach<sup>8</sup>. Before any pharmacological intervention, alternate causes of the symptoms need to be excluded. Medication for painful DPN includes certain antidepressants (e.g., amitriptyline, among others), anticonvulsants (e.g., pregabalin, sodium valproate) and capsaicin cream <sup>10,11</sup>. Other possible options include lidocaine patch, alpha-lipoic acid, isosorbide dinitrate topical spray and transcutaneous electrical nerve stimulation, and finally, surgical decompression.

Despite the large number of studies on Diabetes mellitus, to date, the local literature regarding DPN among newly diagnosed DM type 2 patients is scarce. Thus, this study is aimed to explore the magnitude of DPN so that the patients can be properly screened, identified early, and managed accordingly.

**MATERIAL & METHODS**

This was a hospital-based, cross sectional descriptive study to determine the frequency of diabetic peripheral neuropathy (DPN) among patients newly diagnosed with Diabetes mellitus

Type 2 over a six month period from 01-06-2019 to 01-12-2019 in Department of Medicine (OPD and inpatient), Liaquat University Hospital, Hyderabad / Jamshoro. Non-probability consecutive sampling was utilized for sample collection. Following informed consent, the data was noted on pre-determined proforma. Inclusion criteria were: Patients of both genders, between age 30 years and 60 years newly diagnosed with Type 2 Diabetes mellitus i.e. the serum fasting sugar level above 126 mg/dL along with HbA1c levels above 6.5% was considered as Diabetes mellitus. Who were agreeable to participate in the study after informed consent? Exclusion criteria were pregnant women, patients with chronic lumbar pain, neurological findings suggestive of myopathic involvement or relevant myelo/radiculopathies, as well as subjects with altered consciousness, Glasgow Coma Scale scoreless than 15 or who were inarticulate and unable to collaborate with neurological evaluation. The composed dataset was entered in SPSS (for version 20.0) and analysed accordingly. The frequency and percentage was calculated for gender, residence (urban or rural), hypertension, smoking, obesity, diabetic retinopathy, hyperlipidemia, diabetic nephropathy and DPN. The mean and standard deviation (SD) was obtained for all relevant quantitative data i.e. age, fasting blood sugar plusemoglobin A1C (HbA1c). Stratification regarding age, gender, urban or rural hypertension: smoking, obesity, diabetic retinopathy, hyperlipidemia and diabetic nephropathy was done to see the effect on outcome and to adjust for effect modification. The post stratification Chi-square test was utilized for categorical variables at 95% confidence interval and the p-value ≤0.05 was considered as statistically significant.

TABLE 1: FREQUENCY OF DIABETIC PERIPHERAL NEUROPATHY (DPN) AMONG NEWLY DIAGNOSED TYPE 2 DIABETES MELLITUS BY AGE, GENDER, OBESITY, HYPERTENSION, DIABETIC RETINOPATHY AND HYPERLIPIDEMIA STATUS

VARIABLE PARAMETER	DIABETIC PERIPHERAL NEUROPATHY		Total	P-value
	Yes	No		
<b>AGE IN YEARS</b>				
≤ 40	18(36%)	32(64%)	50	0.0005
41-50	15(25%)	45(75%)	60	
>50	26(65%)	14(35%)	40	
<b>GENDER</b>				
Male	51(56%)	40(44%)	91	0.0005
Female	8(13.6%)	51(86.4%)	59	
<b>OBESITY</b>				
<27.5 (kg/m <sup>2</sup> )	22(30.6%)	50(69.4%)	72	0.034
≥27.5 (kg/m <sup>2</sup> )	37(47.4%)	41(52.6%)	78	
<b>HYPERTENSION</b>				
Yes	52(63.4%)	30(36.6%)	82	0.0005
No	7(10.3%)	61(89.7%)	68	
<b>DIABETIC RETINOPATHY</b>				
Yes	32(65.3%)	17(34.7%)	49	0.0005
No	27(26.7%)	74(73.3%)	101	
<b>SMOKING</b>				
Yes	25(56.8%)	19(43.2%)	44	0.005
No	34(32.1%)	72(67.9%)	106	
<b>HYPERLIPIDEMIA</b>				
Yes	47(38.2%)	76(61.8%)	76	0.548
No	12(44.4%)	15(55.6%)	15	

## RESULTS

150 patients who had been recently diagnosed with Type 2 DM took part in this study.

The DPN frequency among study sample was 39.3% (n=59). The mean patient age was 44.8±7.7 years. The mean fasting blood sugar was 140.0±9.4 and mean HbA1c was 7.05±0.3. There were 60.7% majority of male (n=91) and 39.3% female (n=59) patients. Rural and urban cases were 50.7% (n=76) and 49.3% (n=74), respectively. 82 patients (54.7%) had hypertension, 49 (32.7%) had diabetic retinopathy, 44 (29.3%) were smokers each, and 58 (38.7%) had diabetic nephropathy from among total patients. The association of diabetic peripheral neuropathy and age was shown to be significant (p=0.0005). DPN was found to occur more in males which was significant (p=0.0005). Frequency of DPN was also associated with obesity which was significant (p=0.034), urban versus rural residence (p=0.0005), as well as presence of hypertension (p=0.0005), as demonstrated in Table 1. Frequency of DPN was also significant in patients with DM (p=0.0005) plus smokers (p=0.005). Diabetic retinopathy was not significantly associated with patients who had hyperlipidemia (p=0.548).

## DISCUSSION

Diabetes mellitus (DM) is an ongoing public health burden with numerous obstacles and an increasing occurrence. Progressive urban lifestyles have led to sedentary habits, high calorie foods, and psychosocial stresses<sup>12</sup>. Additionally, Pakistan urban areas have high exposure to contaminants and pollution which enhance risk. In developed Caucasian countries, Diabetes tends to affect older age group above 65 years; but developing countries have cases between 45 to 64 years<sup>13</sup>. The prevalence of DM Type 2 in patients between ages 30 to 50 years is also higher in developing countries<sup>14</sup>. Furthermore, prospective Asian studies opine regarding obesity, which is directly related to the frequency of hypertension, DM Type 2 and high lipid levels<sup>15</sup>. In our study out of 150 cases, 52% were obese and 20.67% were hypertensive. A University of Glasgow study suggested that males were more liable to receive type 2 DM diagnosis in comparison to female<sup>16</sup>. Our results are in concordance of this study. Out of 150 patients, there were 60.7% males and 39.3% were females. Frequency of DPN among newly diagnosed type 2 DM was 39.3%. We detected a high frequency of DPN in newly diagnosed DM patients. DPN was associated with advancing age and symptom periods, before DM was diagnosed as we found that rate of DPN was raised in patients above 50 years old. The results agree with Middle Eastern countries research, in which DPN prevalence was 45%, 31.9%, 25.6% and 29.2% in Saudi Arabia, Iran, United Arab Emirates (UAE) and India, respectively<sup>17,18</sup>. Our data showed that DPN was greater in patients with retinopathy complications associated with DM i.e. 65.3%. Previous studies demonstrated that DM patients

having additional micro- and macrovascular complications were more likely to have DPN<sup>19,20</sup>. This can be due to overlapping pathogenic pathways secondary to toxic hyperglycemia causing growing thickness of endo-neural blood vessels, advanced glycation end products accumulation (AGEs), polyol pathway activation and oxidative stress<sup>21,22</sup>. Other authors reported that patients with lipid disorders were 2.2 times more likely to have DPN<sup>19,23</sup>. Current data backs the principle that metabolic syndrome plus high BMI potentiate DPN risk. In contrast to this, our study reports that it was not statistically significant with patients who had hyperlipidemia.

## CONCLUSION

Our results reinforced the need for rigorous agenda spursing prompt DPN detection and rapid application of patient education particularly those with comorbidities and risk factors such as dyslipidemia along with cardiac and eye complications in chronic DM patients. Moreover, initial actions to prevent DPN by putting into practice lifestyle changes such as healthy diets and physical fitness ideally ought to be implemented to delay such an incapacitating complication.

## CONFLICT OF INTEREST

There is no conflict of interest to be declared.

## REFERENCES

- Schreiber AK, Nones CFM, Reis RC, Chichorro JG, Cunha JM. Diabetic neuropathic pain: physiopathology and treatment. *World J Diabetes*. 2015;6(3):432-44.
- Deli G, Bosnyak E, Pusch G, Komoly S, Feher G. Diabetic neuropathies: diagnosis and management. *Neuroendocrinology*. 2013;98(4):267-80.
- Ali LM, Naeem SN, Hafeez BA, Jai P. Frequency of peripheral neuropathy in newly diagnosed patients of diabetes mellitus: a clinical and electrophysiological basis. *Karachi: Pak J Neuro Sci*. 2014;9(4):31-5.
- Jaiswal M, Divers J, Dabelea D, Isom S, Bell RA, Martin CL, et al. Prevalence of and risk factors for diabetic peripheral neuropathy in youth with type 1 and type 2 diabetes: search for diabetes in youth study. *Diabetes Care*. 2017;40(9):1226-32.
- Khawaja N, Abu-Shennar J, Saleh M, Dahbour SS, Khader YS, Ajlouni KM. The prevalence and risk factors of peripheral neuropathy among patients with type 2 diabetes mellitus; the case of Jordan. *Diabetol Metab Syndr*. 2018;10(1):8.
- Khalil SA, Megallaa MH, Rohoma KH, Guindy MA, Zaki A, Hassanein M, et al. Prevalence of chronic diabetic complications in newly diagnosed versus known type 2 diabetic subjects in a sample of Alexandria population, Egypt. *Curr Diabetes Rev*. 2018;24:19.
- Kisozi T, Mutebi E, Kisekka M, Lhatoo S, Sajatovic M, Kaddumukasa M, et al. Prevalence, severity and factors associated with peripheral neuropathy among newly diagnosed diabetic patients attending Mulago hospital: a cross-

- sectional study. *Afr Health Sci.*2017;17(2):463-73.
8. Boulton AJ, Vinik AI, Arezzo JC, Bril V, Feldman EL, Freeman R, et al. Diabetic neuropathies: a statement by the American Diabetes Association. *Diabetes Care.*2005; 28:956-62.
  9. Callaghan BC, Little AA, Feldman EL, Hughes RA. Enhanced glucose control for preventing and treating diabetic neuropathy. *Cochrane Database Syst Rev.* 2012; :CD007543.
  10. Griebeler ML, Morey-Vargas OL, Brito JP, Tsapas A, Wang Z, Leon BG, et al. Pharmacologic interventions for painful diabetic neuropathy: An umbrella systematic review and comparative effectiveness network meta-analysis. *Ann Intern Med.* 2014;161:639-49.
  11. Dy SM, Bennett WL, Sharma R, et al. Preventing complications and treating symptoms of diabetic peripheral neuropathy. *Comparative Effectiveness Review No. 187. AHRQ Publication No. 17-EHC005-EF.* Rockville, MD: Agency for Healthcare Research and Quality; March 2017. <https://effectivehealthcare.ahrq.gov/ehc/products/612/2436/diabetic-neuropathy-report-170324.pdf> (Accessed on April 06, 2017).
  12. Hu FB. Globalization of Diabetes: The role of diet, lifestyle, and genes. *Diabetes Care.* 2011;34:1249-57.
  13. Cockram C. The epidemiology of diabetes mellitus in the Asia-Pacific region. *Hong Kong Med J.* 2000;6:43–52.
  14. Oh SW, Shin S-A, Yun YH, Yoo T, Huh B-Y. Cut-off point of BMI and obesity-related comorbidities and mortality in middle-aged Koreans. *Obesity Res* 2004; 12: 2031–40.
  15. <http://www.healthline.com/health/recognizing-diabetes-symptoms-men#Overview1>
  16. Al-Geffari M. Comparison of different screening tests for diagnosis of diabetic peripheral neuropathy in Primary Health Care setting. *Int J Health Sci.* 2012;6:127–34.
  17. Tabatabaei-Malazy O, Mohajeri-Tehrani MR, Madani SP, Heshmat R, Larijani B. The prevalence of diabetic peripheral neuropathy and related factors. *Iran J Public Health.* 2011;40:55–62.
  18. Won JC, Kwon HS, Kim CH, Lee JH, Park TS, Ko KS, et al. Prevalence and clinical characteristics of diabetic peripheral neuropathy in hospital patients with type 2 diabetes in Korea. *Diabet Med.* 2012;29:e290–6.
  19. Rosson GD, Dellon AL. Vascular risk factors and diabetic neuropathy. *N Engl J Med.* 2005;352:1925–7.
  20. Tesfaye S, Selvarajah D. Advances in the epidemiology, pathogenesis and management of diabetic peripheral neuropathy. *Diabetes Metab Res Rev.* 2012;28(1):8–14.
  21. Ziegler D, Ametov A, Barinov A, Dyck PJ, Gurieva I, Low PA, et al. Oral treatment with  $\alpha$ -lipoic acid improves symptomatic diabetic polyneuropathy the SYDNEY 2 trial. *Diabetes Care.* 2006;29:2365–70.
  22. Booya F, Bandarian F, Larijani B, Pajouhi M, Nooraei M, Lotfi J. Potential risk factors for diabetic neuropathy: a case control study. *BMC Neurol.* 2005;5:24–8.
  23. Al-Mahroos F, Al-Roomi K. Diabetic neuropathy, foot ulceration, peripheral vascular disease and potential risk factors among patients with diabetes in Bahrain: a nationwide primary care diabetes clinic-based study. *Ann Saudi Med.* 2007;27:25–31.