ID Design Press, Skopje, Republic of Macedonia Open Access Macedonian Journal of Medical Sciences. https://doi.org/10.3889/oamjms.2018.262 eISSN: 1857-9655 *Clinical Science*



A Study of Nerve Conduction Velocity in Diabetic Patients and its Relationship with Tendon Reflexes (T-Reflex)

Khadijeh Haji Naghi Tehrani^{*}

Department Neurology, Islamic Azad University, Tehran Medical Sciences Branch, Tehran, Iran

Abstract

Citation: Tehrani KHN. A Study of Nerve Conduction Velocity in Diabetic Patients and its Relationship with Tendon Reflexes (T-Reflex). Open Access Maced J Med Sci. https://doi.org/10.3889/oamjms.2018.262

Keywords: Diabetes; Neuropathy; Neurological disorder; Nerve conduction velocity; Tendon reflex

*Correspondence: Khadijeh Haji Naghi Tehrani. Department of Neurology, Islamic Azad University, Tehran Medical Sciences Branch, Tehran, Iran. E-mail: Dr_tehrani/0@yahoo.com

Received: 19-Apr-2018; Revised: 21-May-2018; Accepted: 25-May-2018; Online first: 17-Jun-2018

Funding: This research did not receive any financial support

Competing Interests: The authors have declared that no competing interests exist

BACKGROUND: Neuropathy is one of the most common complications of diabetes mellitus. Neuropathy can cause the sensory deficit, neurological disorder, limb ulcers, osteomyelitis, and amputation. Therefore, neurological examinations, determining the nerve conduction velocity and performing sensory and motor tests are important for timely diagnosis and treatment.

AIM: The present study aimed to investigate the nerve conduction velocity in diabetic patients and its relationship with tendon reflexes.

MATERIAL AND METHODS: The present study was observational-cross sectional research carried out on 77 diabetic patients who were admitted into the EMG/NCV Department of Shariati Hospital in the academic year 1996-1997. In all patients, the medical history of the patient (age, duration of diabetes, gender and age of onset of diabetes), neurological examination, nerve conduction velocity, heat test, vibration test, tendon reflexes, D.L and Amplitude were examined and recorded. Finally, the raw data obtained were entered into the IBM SPSS Statistics software, and the important relationships between these variables were analysed. Moreover, in the present study, the statistical significance level (P-value) was considered less than 0.05.

RESULTS: The present study was conducted on a population consisting of 48 women and 29 men with diabetes. The age range of participants was 14-70 years old with an average age of 50.506 ± 7.50 . The results of present study showed that the participants with clinical neuropathy (11.2 ± 7.2) had a significantly longer duration of diabetes than the normal group and those participants with sub-clinical neuropathy (P-value = 0.12). Statistical analyses indicated that increase in age, increase in the duration of diabetes and the gender of male significantly made the nerve conduction velocity abnormal. The analysis of the response to neural reflexes indicated that the ratio of neurological disorders in the five nerves of the ankle and knee was generally higher in the abnormal group (the patients with nerve conduction disorder) compared to the normal (the patients with normal nerve conduction) and in some cases, such as the ulnar motor nerve of ankle (P-value = 0.010), and the ulnar motor nerve of knee (P-value = 0.003) and the sural sensory nerve of knee (P-value = 0.003), increase in neurological disorders was significant.

CONCLUSION: Increase in age, increase in the duration of diabetes, and the male gender can significantly increase the risk of abnormal nerve conduction velocity.

Introduction

Diabetes mellitus is one of the most commonly known chronic diseases in the world. Today, in many countries, diabetes has become an epidemic disease [1]. According to World Health Organization (WHO), in 2000, there were about 170 million people with diabetes and this population will increase to 366 million in 2030, if this disease is not treated and prevented [2]. In Iran, the prevalence of diabetes in different regions is more than 5% [3]. Studies have shown that the diabetes is the main cause of blindness and amputation in some countries [4]. One of the most important complications of diabetes is neuropathy or neurological disorder. There are different types of diabetic neuropathy: polyneuropathy, diabetic amyotrophy, autonomic neuropathy, mononeuropathy multiplex, and diabetic ophthalmoplegia. mononeuropathy, Diabetic neuropathy is observed in different forms in patients and corneal sensorimotor neuropathy (about 75%) can be mentioned as its common form [5]. In various studies, the prevalence of neuropathy was reported to be between 1.5% and 100% [6]. Neuropathy has a high prevalence in both IDDM and NIDDM forms [7]. According to the studies in this field, one of three diabetic patients has diabetic neuropathy [8]. Diabetic neuropathy is diagnosed in 10% of diabetic patients at the time of diagnosis of diabetes. and rest of the patients would suffer neuropathy some vears after onset of the diabetes [9]. Some complications of diabetic neuropathy are severe pain, reduced sensation, increased diabetic foot ulcers and amputation [10]. One of the main problems of diabetic neuropathy is the lack of a reliable and agreed clinical scale for grading the severity of neuropathy, so that it can be used in clinical trials to examine the effect of different therapeutic approaches among patients [11]. The most important diagnostic criteria for diabetic neuropathy, which are confirmed by experts. are disturbances in nerve conduction velocity, increased threshold of sensory nerves, and disturbances in autonomic system function tests. One of the diagnostic methods for diabetic neuropathy is to conduct electro-diagnostic tests (nerve conduction velocity determination). According to a study by Dyck, Nerve Conduction Velocity (NCV) determination is not only the most sensitive test for the diagnosis of diabetic neuropathy, but also has some features such as being repeatable. Moreover, it is considered as a specific test for neurological disorder. According to him, the disadvantage of this test is that it does not provide direct information on signs and symptoms of neuropathy [12]. Nerve damages caused by diabetes can be categorized into two groups; myelin and axonal damages categories. The nerve conduction velocity mainly reflects the myelin changes, while the action potential amplitude indicates the axonal changes and the state of the nerve fibers. The action potential amplitude is an estimate of the number of neural fibers activated by electrical stimulation, and its reduction implies an axonal damage. According to studies conducted in this field by researchers and experts, it was found that the nerve conduction velocity is more variable than the action potential amplitude and is more affected by interventions [13].

Given contents mentioned above and in this regard, the present study aimed to investigate the nerve conduction velocity in diabetic patients and its relationship with tendon reflexes.

Material and Methods

The present research was an observationalcross-sectional study. It was conducted on 77 patients with type 2 of diabetic neuropathy who were under the diabetes clinic and admitted into EMG/NCV department of Shariati Hospital in the academic year 1996-1997. After explaining the research process and getting informed consent from patients, the patients voluntarily participated in the present study by medical ethics principals. Exclusion criteria were including: being older than 70 years old, the presence of thyroid disease, uremia, autoimmune diseases including Rheumatoid arthritis (RA), nutritional, and toxic disorders, collagen-vascular diseases (CVDs) are a heterogeneous group of autoimmune disorders and history of taking certain drugs. Then, from all the patients selected, serologic tests including thyroid tests, complete blood count (CBC) collagen vascular tests and Rheumatoid factor (RF) were taken, and the patients were excluded from the study if the results of tests were positive. In the next step, the medical history of the patients was recorded, and then, they were placed under physical and neurological examinations. A questionnaire used in present study includes the questions on gender, age, duration of diabetes, the age of onset of diabetes and type of treatment.

Neurological examination includes the examination of cranial nerves, motor system, sensory system (the perception of external sensations (exteroception), including touch, temperature. pressure, vibration, and itch) proprioception, vibration sense and heat sensation), and tendon reflexes. The last part of the study, is devoted to the measurement of the nerve conduction velocity in the median nerves (both sensory and motor branches), ulnar nerve (motor branch), peroneal nerve (motor branch) and the sural nerve (sensory branch) and the rates of NCV (M/S), D.L (s), and amplitude (mv) in the mentioned nerves were recorded and if they were insignificant, they were recorded as detectable in the united nations system. The values of these parameters were evaluated according to the tables presented in the valid electrophysiology books written by researchers such as OH [14] and Chu-Andrews [15] and considering the age of the patient. Finally, they were divided into two groups. The other main variables were the Ankle Jerk (AJ) Reflex and the Knee Jerk (KJ) Reflex. To score the reflexes and according to the age range of patients, just Ankle Jerk (AJ) Reflex could be adjusted.

For this purpose, if the Ankle Jerk (AJ) Reflex was absent and the patient was older than 50, he/she would get the score 1, instead of zero, and if it were reduced, he/she would get the score 2, instead of 1. To assess the vibration sense, the diapason (Hz) was used. First, it was placed on the medial malleolus of the right foot, and the test was considered abnormal when the vibration sense was confirmed by the patient. In the case the vibration sense was not confirmed by the patient, the diapason was immediately moved to the Tuberosity of the tibia. To investigate the heat sensation, two test tubes containing 20°C and 45°C water were used. To examine orthostatic hypotension, greater than 20 mmHg reduction in systolic pressure or 10 mmHg reduction in diastolic pressure while changing the position of the patient from lying down to standing was considered as a positive sign for orthostatic hypotension.

It should be noted that all tests mentioned above, except for vibration and heat test performed by the students at the Research Center of Endocrinology Clinic, were conducted by an expert. Then, the raw data obtained were entered into IBM SPSS Statistics. To analyse the data, T-test, Variance Fisher's Exact Test and Chi-squared test were used. The statistical significance level (P-value) was considered less than 0.05.

Results

In the present study, 77 patients participated of which 49 and 29 patients were female and male, respectively (the female-to-male ratio was 1.6 in the present study). The age range of patients was 14 to 70 years old, and their average age was 50.506 ± 7.50 years the average age of patients at the onset of diabetes was 40.50 ± 10.50 years. Descriptive statistics show that average duration of diabetes was 9.70 ± 6.90 years. Moreover, 41.2% of patients were treated with the pill (oral antidiabetic agents), and 44.2% of patients were treated with insulin. In this study, 66.2%, 11.7% and 2.6% of patients complained of paresthesia, weakness and pain, respectively. According to the examinations, it was found that in 36.5% of patient, heat sensation was impaired and in 32.8% of them, vibration sense was impaired. The patients who participated were divided into three groups based on clinical signs and symptoms:

The first group (clinical neuropathy) was made up of 54 patients with severe neurological symptoms.

The second group (sub-clinical neuropathy) had 20 patients with main disturbances in vibration and heat tests

Third group (normal): consisting of 3 patients with normal test results.

In this study, the duration of the disease was examined with three clinical groups (clinical neuropathy, subclinical neuropathy and normal) (Table 1). The results indicate that there are significant differences between parameters mentioned above (P-value < 0.05).

Table 1: The relationship between clinical symptoms and duration of the disease

		Duration of the	P-Value
		disease	
		Mean ± S.D. (year)	
Group	Clinical neuropathy	11.2 ± 7.2	0.012
	Sub-clinical neuropathy	8.9 ± 8.3	
	Normal	4.41 ± 4.48	

On the type of treatment, in both normal and abnormal groups, no significant difference was observed between parameters mentioned above (P- value = 0.1000). According to NCS results, the patients studied were divided into two groups: normal (normal nerve conduction) and abnormal (disturbance in nerve conduction). There were 7 patients in the normal group, and there were 70 patients in the abnormal group. In Table 2, the average age of the patient, the age of onset of the disease and the duration of the disease in both groups (normal and abnormal) are summarised. Statistical analysis showed that in both groups, there was a significant difference between the age of the patient and the duration of the disease (P-value < 0.05) (each group was examined separately).

Table 2: Relationship between the parameters studied in the two NCS groups

	In NCS group Mean ± S.D.		P-Value	
Age (year)	53.3 ± 11.3	42.8 ± 13.8	0.002	
Age of the onset of disease (year)	41.07 ± 9.21	35.60 ± 13.37	0.580	
Duration of disease (year)	12.27 ± 7.40	4.21 ± 3.74	0.001	

The statistical analyses performed in the present study showed a statistically significant difference between gender and nerve conduction velocity (P-value = 0.019). In the present study, tendon reflexes were divided into two groups: Kneejerk Reflexes (Kj) and Ankle jerk Reflexes (Aj). Each of these groups has been classified into three parts: Absent, Decreased, and Normal. In both normal and abnormal groups, the highest frequency belonged to the Normal part (81.8%), and the lowest frequency belonged to the absent group (1.3%). The researchers have investigated the relationship between nerve conduction velocity and tendon reflexes (Table 3). Statistical analyses showed that there were significant differences among the knee-jerk reflexes and the ulnar motor nerves, peroneal motor nerves and sural sensory nerves (each separately) (P-value < 0.05). It was also found that there is a statistically significant difference between the ankle ierk reflexes and ulnar motor nerves (P-value < 0.05).

 Table 3: Relationship between neurological disorders and tendon reflexes

			Nerves (%)				
			Median	Median	Ulnar	Peroneal	Sural
			motor	sensory	motor	motor	sensory
Tendon	Ankle jerk	Normal	63.5	63.5	22.2	58.7	41.3
reflexes	reflexes	Abnormal	92.9	92.9	57.1	71.4	71.4
		P-Value	0.050	0.050	0.010	0.560	0.080
	Knee jerk	Normal	61.9	61.9	20.6	58.7	41.3
	reflexes	Abnormal	100	100	64.3	71.4	71.4
		P-Value	0.003	0.003	0.002	0.560	0.080

Discussion

There is a direct relationship between the prevalence of neuropathy and the progression of diabetes mellitus [16]. The neuropathy is caused by

the presence of signs and symptoms of peripheral nerve disorders in diabetic patients. Nerve damage in diabetic patients has various features and the change in nerve conduction velocity is one of its symptoms. The position and characteristics of the nerve fibers. the severity of diabetes and demographic characteristics (such as age, duration of disease, and gender) are of the factors playing a key role in the severity of sensorimotor neuropathy. As mentioned earlier, such complications are common in diabetics, and somewhat predictable, but they have a wide range of changes and potency, which have been the subject of research by researchers. For example, Soivers et al (2004) have conducted a research in order to perform the clinical and electroneurographic study of peripheral nerve involvement in diabetic patients.

This study was performed on 103 diabetic patients with a mean age of 52.6 ± 14.00 years old who were randomly selected from patients admitting to endocrinology clinic of Shiraz University of Medical Sciences. It was found that 29.4% of patients had type 1 diabetes and 70.6% had type 2 diabetes. In their study, they stated that there is a direct relationship between the prevalence of neuropathy and the duration of the disease. The most commonly result was the reduced ankle reflexes and reduced vibration sense in the legs [17]. In another study by Andersen et al., (2012), motor dysfunctions in diabetic patients were examined. They argued that neuropathy is a frequent complication of diabetes, and motor system involvement is rarely seen in a clinical examination and can be diagnosed using guantitative techniques (isoquinoline dynamometer, type of diabetes, ankle jerk reflex and knee-jerk reflex). In fact, they believe that muscular weakness depends on the symptoms and severity of diabetic neuropathy in patients. Therefore, it can be said that diabetic neuropathy can reduce muscle strength [18].

The diversity of studies conducted in this field allows us to review the reports on specific cases in a case study. For example, Aaron (2016) has conducted a study on sensorimotor neuropathy in diabetic patients. In this study, it was reported that a 65-yearold woman with a 5-year history of diabetes and the symptoms such as pins-and-needles sensation and pain referred a physician and it was observed that in this case, burning and pinpricks sensation in the knee area, as well as the ability to detect vibration from a tuning fork 128 Hz, have decreased, and she has lost proprioception and sensation to the monofilament 1-g in her toes. Moreover, there was no knee-jerk reflex in her. All of these were due to diabetic sensory and motor neuropathy [19].

Based on the results of our study, it can be concluded that people with clinical neuropathy have a longer duration of diabetes. Moreover, increase in age, increase in the duration of diabetes, and the gender of the male can significantly make the nerve conduction velocity abnormal. The analysis of the response to neural reflexes indicated that the ratio of neurological disorders in nerves of the ankle and knee was generally higher in the abnormal group (the patients with nerve conduction disorder) compared to the normal (the patients with normal nerve conduction).

References

1. Aghamollaei T, Eftekhar H, Shojaeizadeh D, Mohammad K, Nakhjavani M, Pour FG. Behaviour, metabolic control and healthrelated quality of life in diabetic patients at Bandar Abbas diabetic clinic. Iranian Journal of Public Health. 2003; 32(3):54-9.

2. Aghamollaie T, Eftekhar H, Shojaeizadeh D, Mohammad K, Nakhjavani M, Ghafrani F. Effect of a health education programme on behavior, HbA1C and health-related quality of patient. Acta Med Iran. 2004; 43:89-94.

3. Larejani B, Zahedi F. Epidemiology of diabetes mellitus in Iran. Iranian Journal of Diabetes and Lipid Disorders. 2001; 1(1):1-8.

4. Rajab A, Mahmoodi M, Adili F. Assessment of effect of applying the transtheoretical model to physical activity on health indexes of diabetic type 2 patients. Medical Science Journal of Islamic Azad Univesity-Tehran Medical Branch. 2008; 18(1):21-7.

5. Bansal V, Kalita J, Misra U. Diabetic neuropathy. Postgraduate medical journal. 2006; 82(964):95-100. https://doi.org/10.1136/pgmj.2005.036137 PMCid:PMC2596705

6. Janghorbani M, Rezvanian H, Kachooei A, Ghorbani A, Chitsaz A, Izadi F, et al. Peripheral neuropathy in type 2 diabetes mellitus in Isfahan, Iran: prevalence and risk factors. Acta neurologica scandinavica. 2006; 114(6):384-91. <u>https://doi.org/10.1111/j.1600-0404.2006.00716.x</u> PMid:17083338

7. Maser RE, Steenkiste AR, Dorman JS, Nielsen VK, Bass EB, Manjoo Q, et al. Epidemiological correlates of diabetic neuropathy: report from Pittsburgh Epidemiology of Diabetes Complications Study. Diabetes. 1989; 38(11):1456-61. https://doi.org/10.2337/diab.38.11.1456.PMid:2620781

https://doi.org/10.2337/diab.38.11.1456 PMid:2620781

8. Ziegler D, Nowak H, Kempler P, Vargha P, Low P. Treatment of symptomatic diabetic polyneuropathy with the antioxidant α-lipoic acid: a meta-analysis. Diabetic Medicine. 2004; 21(2):114-21. https://doi.org/10.1111/j.1464-5491.2004.01109.x PMid:14984445

9. Feldman EL, Russell JW, Sullivan KA, Golovoy D. New insights into the pathogenesis of diabetic neuropathy. Current opinion in neurology. 1999; 12(5):553-63. <u>https://doi.org/10.1097/00019052-199910000-00009</u> PMid:10590892

10. Booya F, Bandarian F, Larijani B, Pajouhi M, Nooraei M, Lotfi J. Potential risk factors for diabetic neuropathy: a case control study. BMC neurology. 2005; 5(1):24. <u>https://doi.org/10.1186/1471-2377-5-24</u> PMid:16336693 PMCid:PMC1343576

11. Tkac I, Bril V. Glycemic control is related to the electrophysiologic severity of diabetic peripheral sensorimotor polyneuropathy. Diabetes care. 1998; 21(10):1749-52. https://doi.org/10.2337/diacare.21.10.1749 PMid:9773742

12. Dyck PJ. Evaluative procedures to detect, characterize, and assess the severity of diabetic neuropathy. Diabetic Medicine. 1991; 8(S2). <u>https://doi.org/10.1111/j.1464-5491.1991.tb02156.x</u> PMid:1825958

13. Aminoff M, Albers J. Electrophysiological techniques in the evaluation of patients with suspected neurotoxic disorder. Electrodiagnosis in clinical neurology. 2005; 782.

14. Oh SJ. Clinical electromyography: nerve conduction studies: Lippincott Williams & Wilkins, 2003.

15. Chu-Andrews J, Johnson RJ. Electrodiagnosis: an anatomical and clinical approach: Lippincott, 1986.

16. Tesfaye S, Selvarajah D. Advances in the epidemiology, pathogenesis and management of diabetic peripheral neuropathy. Diabetes/metabolism research and reviews. 2012; 28(S1):8-14. https://doi.org/10.1002/dmrr.2239 PMid:22271716

17. Soveid M, Ghavanini MR, Shirdel E, Omrani G. Clinical and electroneurographic evaluation of neuropathy among diabetic patients in Shiraz. Iranian Journal of Diabetes and Lipid Disorders. 2004; 3(1):53-6.

18. Andersen H. Motor dysfunction in diabetes. Diabetes/metabolism research and reviews. 2012; 28(S1):89-92. https://doi.org/10.1002/dmrr.2257 PMid:22271730

19. Vinik AI. Diabetic sensory and motor neuropathy. New England Journal of Medicine. 2016; 374(15):1455-64. https://doi.org/10.1056/NEJMcp1503948 PMid:27074068