

Comparison of Two-point Discrimination Perception in Stroke Patients with and without Diabetes Mellitus

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Abstract. [Purpose] The aim of this study was to compare two-point discrimination (TPD) perception in stroke patients with diabetes mellitus (DM) and without diabetes mellitus (non-DM). [Subjects] The subjects were 53 poststroke hemiparetic patients (21 stroke patients with DM; 32 stroke patients without DM). [Methods] TPD was measured on the tips of the first through fifth fingers on both the affected and unaffected sides. [Result] Comparison of TPD between fingers on the unaffected side and affected side fingers showed significantly poorer responses in all five fingers on the affected side. TPD was also significantly poorer in the DM group compared with the non-DM group in all five fingers on the affected side, but no differences were observed for the unaffected side. [Conclusion] These findings suggest that TPD was significantly poorer in the fingers on the affected side vs. the unaffected side in poststroke hemiparetic patients. DM caused a significantly poorer TPD in the fingers on the affected side in post-stroke patients but had no significant effect on the fingers on the unaffected side.

Key words: Stroke, Two-point discrimination, Diabetes mellitus

(This article was submitted Mar. 1, 2013, and was accepted Apr. 8, 2013)

INTRODUCTION

The prevalence of diabetes mellitus (DM) in Korea is estimated to be 7.3%, and this has increased about 5-fold over the past 30 years according to a report of the Korea National Health and Nutrition Examination Surveys¹⁾. DM represents a strong independent risk factor for stroke²⁾, as long-term diabetes results in a variety of subtle cerebral disorders that occur more frequently than is common³⁾. DM also affects conductive function in the central and peripheral somatosensory pathways⁴⁾.

The incidence of sensory deficit in stroke is high⁵⁾. Severe sensory loss after stroke can cause patients to avoid use of their affected fingers in manual activities, even though good voluntary muscle activity is present⁶⁾. Sensory functions are important for a patient's rehabilitation to restore impaired motor function. Therefore, Dellon⁷⁾ suggested use of an instrument to measure the sensory threshold of two-point discrimination (TPD).

TPD is defined as the smallest separation between two stimulations placed on the skin that can be discriminated as two separate points. TPD testing has been found to be particularly helpful in the assessment of injuries to nerves distributed to the hand sensor⁷⁾. TPD has been demonstrated to be a valid measurement of functional sensibility in the hand with good test⁸⁾. Previous studies have used this measurement to assess sensory capacity in adult men and women⁹⁾, traumatic brain injury patients¹⁰⁾, lumbosacral

radiculopathy patients¹¹⁾ and normal subjects¹²⁾. However, most TPD studies have been conducted on peripheral injury patients or normal subjects, and few have focused on central nervous system disorders. To our knowledge, studies on poststroke patients are lacking, and the influence of DM on TPD has yet to be studied.

Therefore, this study was designed to identify the extent of differences in TPD in fingers on the affected and unaffected sides in poststroke hemiparetic patients. A second aim was to estimate the influence of DM on finger sensory abilities following stroke.

SUBJECTS AND METHODS

Fifty-three poststroke patients were recruited as subjects from the stroke rehabilitation center of An-dong Hospital (Table 1). Patients were excluded if they had previous neurological injury or disease (for example, traumatic brain injury, peripheral neuropathy, and brachial plexus injury), language deficits or psychiatric history. Informed consent was obtained from each subject prior to inclusion in the study.

TPD was measured following the guidelines published by Moberg in 1990⁸⁾. Accuracy was increased by using Aesthesimeters (Sammons Preston, USA) with a precision of 1 mm in place of a paperclip. During testing, the examiner and the examinee were seated with their fingers stabilized against a firm support to avoid movement. A screen was used to prevent the subjects from visualizing their fingers. TPD sensitivity in each region was examined by lightly touching the subject's skin at either one point or

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Table 1. General characteristics of the subjects (n=53)

	Age (years)	Height (cm)	Weight (Kg)	Sex (F/M)	Duration months	Diabetes Mellitus (Y/N)
Subjects	63.25 ± 9.17 ^a	160.58 ± 6.94	59.79 ± 10.83	23 / 30	2.01 ± 2.25	21 / 32

^aMean ± SD**Table 2.** Comparison of the fingers on the affected side and unaffected side in stroke patients (n=53)

Variable	Unaffected side (mm)	Affected side (mm)
First finger*	4.72 ± 1.06	6.32 ± 2.99
Second finger*	4.92 ± 1.28	6.74 ± 3.23
Third finger*	5.66 ± 1.59	7.62 ± 3.81
Fourth finger*	6.23 ± 1.81	8.57 ± 4.20
Fifth finger*	7.08 ± 2.00	9.36 ± 4.36

Mean ± SD, *p<0.05

two points simultaneously. This type of sensory testing is appropriately conducted using pressure that depresses the skin no more than 1 mm. Testing commenced with 0 mm between the two points of the Aesthesimeters, and then the distance was gradually increased until the subject was able to perceive two points. Testing was then repeated on the next finger. The collected data were analyzed with SPSS ver. 12.0. All parameters for all subjects were tested by the Kolmogorov-Smirnov test to show that the data were normally distributed. An independent t-test was performed to test differences between the affected and unaffected sides and between DM and non-DM patients. The significance level was set at p<0.05.

RESULTS

Comparison of TPD between fingers on the unaffected and affected side fingers showed significantly poorer capabilities in all five fingers on the affected side (p<0.05). Comparison of TPD between the DM and non-DM groups showed a significantly poorer response in all fingers on the affected side in the DM group (p<0.05) However, no significant differences were noted between the DM and non-DM groups for any of the fingers on the unaffected side (p>0.05) (Table 2).

DISCUSSION

The results of the study showed that the TPD of the stroke patients was significantly poorer on the affected than on the unaffected side. The stroke patients had suffered a cerebral vascular accident on the right or left brain hemisphere. Sensory processing involves many somatosensory pathways and many areas of the brain. Therefore, sensory impairment can result from a lesion located anywhere from the brainstem to the cortex¹³.

Table 3. Comparison of TPD in stroke patients with and without diabetes mellitus (n=53)

Finger		DM group (n=21)	Non-DM group (n=32)
First	Unaffected	4.90 ± 1.04 ^a	4.59 ± 1.07 ^b
	Affected*	7.76 ± 3.85	5.38 ± 1.77
Second	Unaffected	5.17 ± 1.24	4.78 ± 1.31
	Affected*	8.48 ± 3.93	5.59 ± 2.03
Third	Unaffected	5.90 ± 1.48	5.50 ± 1.67
	Affected*	9.38 ± 4.97	6.47 ± 2.23
Fourth	Unaffected	6.52 ± 1.66	6.03 ± 1.91
	Affected*	10.62 ± 5.63	7.22 ± 2.11
Fifth	Unaffected	7.33 ± 2.11	6.91 ± 1.94
	Affected*	11.14 ± 5.66	8.19 ± 2.76

^aMean ± SD; *p<0.05; ^b millimeters (mm)

Kim and Choi¹⁴) stated that discriminative sensory disturbances, which often occur bilaterally in some modalities, are common in patients with unilateral stroke, even in those with intact sensory function on routine examination. The study of Van Heest et al.¹⁵) showed that 90% of children with spastic hemiplegic damage had a TPD deficit. Similarly, the present study showed significantly poorer TPD in the fingers on the affected side compared with those on the unaffected side in poststroke patients.

The present study investigated the influence of DM on finger sensory perception in poststroke hemiparetic patients. Louraki et al.¹³) determined that DM can affect both the peripheral and autonomic nervous system. However, our results showed that DM had a significant negative influence on the fingers on the affected side in poststroke hemiparetic patients. Therefore, DM may influence the poststroke clinical evaluation, especially in the initial phase, while the area of the cerebral injury is increasing¹⁶). Alterations in cerebral blood supply and metabolic derangements probably play a role, as they do in the pathogenesis of DM³). Therefore, the DM group was more damaged in both primary somatosensory and ipsilateral supplementary perception when compared with the non-DM group.

In conclusion, our results suggest that TPD is significantly poorer in the fingers on the affected side than in those on the unaffected side in poststroke hemiparesis. Effects of DM were also apparent as poorer TPD in the fingers on the affected side in poststroke patients, although no differences were observed for TPD in the fingers on the unaffected side (Table 3).

The study had a number of limitations, particularly due to the relatively small sample size and the predominantly

cross-sectional methodology used for it. Future research would benefit from a larger sample size and a longitudinal design.

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