The effect of skin temperature on vibratory sensitivity in polyneuropathy

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SUMMARY In normal subjects, a rise in skin temperature causes a decrease in vibratory perception thresholds. In this study, vibratory thresholds on the foot were measured before and after local warming of the skin in patients with diabetic or uremic neuropathy. On warming, the thresholds increased in nine of 11 diabetic patients while they decreased in 10 of 13 uremic patients. In two-thirds of the patients, the response was outside the range of normal short term variation.

This study is based on two observations. First, vibratory perception thresholds decrease if the skin is warmed. Second, demyelinated nerve fibres are sensitive to small rises in temperature. An increase of only 0-5°C can block conduction in a demyelinated internode. The present investigation was undertaken to test the hypothesis that demyelinating peripheral diseases cause an abnormal response to skin warming, that is an increase in vibratory thresholds. The vibratory sensitivity was tested in 11 patients with diabetic neuropathy, which is, in part, a demyelinating neuropathy. For comparison, measurements were also made in 13 patients with uremic neuropathy, which is an axonal neuropathy.

Patients and methods

The diabetic group comprised three women and eight men with a mean age of 46 years (range 33–66 years). They all had type 1 diabetes mellitus with a duration of 18 to 46 years. The uremic group comprised six women and seven men with a mean age of 53 years (range 37–71 years). The diagnoses were chronic glomerulonephritis (6), chronic pyelonephritis (4), and polycystic kidney disease (3). They were treated with continuous ambulatory peritoneal dialysis (6) or haemodialysis (7). All patients had a creatinine clearance rate less than 5 ml/min.

Vibratory thresholds were determined with a handheld vibrometer (Somedic AB), where the movement of the stimulating probe (diameter 13 mm) was recorded continuously by means of an accelerometer. The movement ampli-
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When the skin temperature was increased from the normal range of 28–32°C to 36–37°C, vibratory thresholds increased in nine of the 11 diabetic patients and decreased in 10 of the 13 uraemic patients (p = 0.006), see figure. The mean change was +49.7 ± 59.8% in the diabetic group and −24.2 ± 23.5% in the uraemic group (p < 0.001).

In the control group of 20 patients, short term changes ranged from 0 to 17% with the mean of the absolute values being 7.8 ± 6.2%. This accords with previous studies. A 95% confidence interval then gives the following distribution: no significant effect of warming in 1/3 of the patients; a significant increase in 1/3, all diabetics; and a significant decrease in 1/3, all uraemics.

The effect of warming could not be related to (1) the degree of renal impairment, (2) the duration of diabetes, (3) the type of dialysis treatment, (4) the nerve conduction velocities, nor (5) the vibratory thresholds prior to warming.

Discussion

Previous studies have shown that vibratory perception thresholds decrease after a rise in skin temperature. The response depends on the stimulus frequency, and it is probably also influenced by the size of the stimulating probe, the stimulus-site location, the use of a rigid annulus around the stimulating probe, and the size of the area that is warmed. The major finding of this study was that, on warming the skin, the vibratory thresholds increased in the diabetic patients and decreased in the uraemic patients.

Axonal degeneration is the most common abnormality in nerve biopsies from uraemic patients. In diabetics, on the other hand, there is both axonal loss and primary segmental demyelination. It is possible that the remaining nerve fibres in uraemic neuropathy respond normally to warming. In contrast, demyelinated nerve fibres may contribute to the vibratory perception at normal temperatures, but when the nerve is warmed, conduction is blocked in these fibres and the vibratory threshold increases.

An alternative explanation could involve the normal response of vibratory thresholds on warming. The cause of this phenomenon is unknown. It is probably not related to an increased local blood flow or an altered skin conductivity. An investigation by Inman and Peruzzi may be relevant. In recordings from isolated Pacinian corpuscles, they observed repetitive firing after a single stimulus at temperatures above 31.5°C. Their explanation was that at room temperature, the durations of the receptor potential and the nerve action potential are about equal. This means that the first node is refractory during the entire course of the receptor potential. As the temperature is raised, the refractory period of the axon decreases, while the duration of the receptor potential remains constant. At a certain temperature, the refractory period will be sufficiently short to allow re-excitation. Close to the vibratory perception threshold, the nerve fibres are still not entrained. This means that warming could increase the number of impulses transmitted centrally, which would increase the probability of detection.

There was a substantial overlap between the two groups, which at present makes the method less useful clinically. It is possible that a better separation can be obtained by changing the size of the stimulating probe, by varying the stimulus frequency, or by warming the whole leg.

This work was supported by the Foundations of the Karolinska Institute and by the Vivian L. Smith Foundation for Restorative Neurology. I thank Ulf Lindblom and Christopher Korch for carefully reading the manuscript.

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doi: 10.1136/jnnp.48.2.176

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