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Accuracy of Monofilament Testing to Diagnose Peripheral Neuropathy: A Systematic Review

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INTRODUCTION

Peripheral neuropathy causes loss of sensation and increases the risk of ulceration of the feet. Timely identification of loss of protective sensation may allow preventive intervention. Peripheral neuropathy is a complication in approximately 50% of patients with diabetes, and up to 50% of patients with peripheral neuropathy may not have symptoms.1-3

Several tests are used to detect peripheral neuropathy, including vibration perception, application of warmth and cold, and nerve conduction studies, which are assumed to be the reference standard.4 Electrodagnostic tests can be complex, expensive, and time consuming, which hampers their widespread use, especially in primary care, where for most patients peripheral neuropathy is diagnosed and treated.

Monofilament testing is an inexpensive, easy-to-use, and portable test for assessing the loss of protective sensation, and it is recommended by several practice guidelines to detect peripheral neuropathy in otherwise normal feet.1 5-6 Monofilaments, often called Semmes-Weinstein monofilaments, are calibrated, single-fiber nylon threads, identified by values ranging from 1.65 to 6.65, that generate a reproducible buckling stress. The higher the value of the monofilament, the stiffer and more difficult it is to bend. Three monofilaments commonly used to diagnose periph-
eral neuropathy are the 4.17, 5.07 and 6.10. Forces required to bend these monofilaments are 1, 10, and 75 g, respectively. The filament is placed on the patient’s skin (usually the feet), when there is considerable loss of sensation, the patient will not be able to detect the presence of the filament at buckling. The 5.07/10-g monofilament has been described as the best indicator to determine loss of protective sensation.7-12 The aim of this review was to perform a meta-analysis of studies evaluating monofilament testing with the 5.07/10-g monofilament in diagnosing peripheral neuropathy of the feet from any cause.

**METHODS**

We searched MEDLINE and EMBASE from database inception to June 2007 to identify diagnostic accuracy studies of peripheral neuropathy that used monofilament testing. Our search strategy focused on monofilaments, peripheral neuropathy, and diagnostic studies. The complete strategy is available in Supplemental Appendix 1, available at http://annfammed.org/cgi/content/full/7/6/555/DC1). We applied no language restrictions, and we supplemented our searches by manually reviewing the reference lists of eligible studies.

**Selection**

Two reviewers (A.W., J.D.) independently selected potentially relevant studies by titles and abstracts. We included articles when peripheral neuropathy of the feet was the target condition, monofilament testing with a 5.07/10-g monofilament was the index test, and nerve conduction study was used as reference standard. If the 2 reviewers disagreed, consensus was sought with the help of a third reviewer (H.W.). Of all possibly relevant articles, the full text was reviewed using the above-mentioned inclusion criteria.

**Quality Assessment**

The methodological quality of the studies was independently assessed by 2 reviewers (A.W., J.D.) using the Quality Assessment of Diagnostic Accuracy Studies (QUADAS) checklist (the QUADAS checklist can be found in Supplemental Appendix 2, available at http://annfammed.org/cgi/content/full/7/6/555/DC1). In case of disagreement consensus was reached with a third reviewer (H.W.).

**Data Synthesis and Analysis**

Sensitivity and specificity were calculated from 2 × 2 tables or retrieved from data available in the primary articles. The aim of the review was to perform a meta-analysis.

**RESULTS**

The study selection process is shown in Figure 1. The characteristics of the final assessed 3 diagnostic accuracy studies are presented in Table 1. The results of the meta-analysis are shown in Figure 2. The diagnostic accuracy of monofilament testing was calculated for each monofilament (Table 2). The 5.07/10-g monofilament had the highest sensitivity and specificity for the diagnosis of peripheral neuropathy of the feet from any cause.
studies are shown in Table 1. All studies appeared to be limited to patients with diabetes mellitus. Sensitivity of the monofilament test ranged from 0.41 to 0.93, and specificity ranged from 0.68 to 1.00. All studies showed methodological limitations that could have inflated sensitivity or specificity. A meta-analysis was not possible because of this clinical heterogeneity.

We believe our identification of studies has been complete, as we applied no language restriction and conducted a sensitive search. The study with the best characteristics (Lee et al\(^{14}\)) showed a possibly serious methodological flaw: it was unclear whether the interpretation of the monofilament test was influenced by knowledge of the results of the reference standard and vice versa. In addition, a study population of 37 patients is quite small.

Another problem is the lack of standardization of the monofilament test methods. Different methods are described varying from 1 testing site\(^{15,16}\) to 10 testing sites\(^{14}\) on 1 foot, and there is no evidence or consensus about the most appropriate threshold.

We found various published reference standards for peripheral polyneuropathy, including clinical examination, vibration perception thresholds with biothesiometer/vibrameter/tuning forks, warm/cold detection, and nerve conduction studies. We rejected clinical examination, vibration perception, and warm/cold detection as reference standards. We also rejected clinical examination, vibration perception, and warm/cold detection as reference standards. There are no obvious limitations in sensitivity or specificity, and thermal sense detection because it tests small-fiber neuropathy, whereas applying a monofilament and light touch, such as vibration and nerve conduction, tests for large-fiber
neuropathy. Only 4 studies assessed with QUADAS used nerve conduction as the reference standard, of which 3 were included in our final selection.

We also rejected studies if the monofilament test was performed on patients who had visible ulcers. In patients with current visible ulcers, the interpretation of the monofilament test and the nerve conduction studies may be influenced by this knowledge (observer or reviewer bias).

We conclude that despite of the frequent use of the (Semmes-Weinstein) monofilament test, little can be said about the test accuracy for detecting neuropathy in feet that do not have visible ulcers, because diagnostic studies with adequate methodology are lacking. Further research on monofilament testing should focus on optimal standard test application procedures (number and sites) and on defining a reproducible threshold.

As this test is already widely used and advocated in many clinical guidelines, especially for diabetic patients, standardization of the method for the monofilament test and studies to define the sensitivity of this method in clinical practice are important. Meanwhile, the sole use of a monofilament test to diagnose peripheral neuropathy is not recommended. The diagnosis of peripheral neuropathy can be made only after a careful clinical examination with more than 1 test, as recommended by the American Diabetes Association. Tests for this clinical examination are vibration perception (using a 128-Hz tuning fork), pressure sensation (using a 10-g monofilament at least at the distal hallucies), ankle reflexes, and pinprick. When in doubt, a nerve conduction test might be necessary to establish a firm diagnosis.

To read or post commentaries in response to this article, see it online at http://www.annfammed.org/cgi/content/full/7/6/555.

Key words: Peripheral neuropathy; peripheral nervous system diseases; monofilament testing; Semmes-Weinstein monofilament; review, systematic; primary health care; diabetic foot

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Disclaimer: This article does not serve as an endorsement for any particular manufacturer of (Semmes-Weinstein) monofilaments.

References


