

#### **OPEN ACCESS**

EDITED BY Mamede De Carvalho University of Lisbon, Portugal

REVIEWED BY José Castro. Universidade de Lisboa, Portugal Elif Simin Issi. Afyonkarahisar Health Sciences University, Türkiye

\*CORRESPONDENCE Anne Kurver

⋈ A.Kurver@hagaziekenhuis.nl

†PRESENT ADDRESS Anne Kurver.

Department of Neurology, Haga Hospital, The Hague, Netherlands

RECEIVED 11 July 2025 ACCEPTED 23 September 2025 PUBLISHED 08 October 2025

Kurver A, Meulstee J, Bartels RHMA and Verhagen WIM (2025) Proposal for an efficient, minimal electrodiagnostic protocol for confirmation of clinically diagnosed ulnar nerve entrapment neuropathy at the elbow. Front. Neurol. 16:1664216. doi: 10.3389/fneur.2025.1664216

#### COPYRIGHT

© 2025 Kurver, Meulstee, Bartels and Verhagen. This is an open-access article. distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

# Proposal for an efficient, minimal electrodiagnostic protocol for confirmation of clinically diagnosed ulnar nerve entrapment neuropathy at the elbow

Anne Kurver<sup>1\*†</sup>, Jan Meulstee<sup>2</sup>, Ronald H. M. A. Bartels<sup>3</sup> and Wim I. M. Verhagen<sup>1</sup>

<sup>1</sup>Department of Neurology, Canisius Wilhelmina Hospital, Nijmegen, Netherlands, <sup>2</sup>Department of Clinical Neurophysiology, Canisius Wilhelmina Hospital, Nijmegen, Netherlands, <sup>3</sup>Department of Neurosurgery, RadboudUMC, Nijmegen, Netherlands

Introduction: The clinical diagnosis of ulnar nerve entrapment at the elbow can reliably be made based on typical clinical symptoms of UNE. In the present study, we constructed an efficient, minimal electrodiagnostic protocol for confirmation of clinically diagnosed UNE.

Methods: A prospective cross-sectional cohort observational study was conducted among patients with clinical suspicion of UNE. In all, 210 arms were included, each examined according to a standard neurophysiological protocol. Results: Nerve conduction studies (NCS) indicated abnormalities in 60.5% of the cases. Of these, 84.3% had abnormal NCS results for the abductor digiti quinti muscle (ADV). The first dorsal interosseous muscle (FDI) and sensory NCS were abnormal in 68.5 and 59.1% of these cases, respectively.

Conclusion: We recommend starting with NCS of the ADV for cases requiring only one abnormal test is needed to confirm the clinical diagnosis of UNE, followed by sensory NCS if the NCS of the ADV is normal.

KEYWORDS

UNE, ulnar nerve, electrodiagnostics, nerve conduction studies, protocol

#### Introduction

Ulnar nerve entrapment at the elbow (UNE) is the second most prevalent entrapment neuropathy (1, 2). The reported incidence of UNE ranges from 8.9 to 25.2 per 100,000 personyears (3, 4). The clinical diagnosis can be reliably established based on typical clinical symptoms of UNE: pain and numbness in the ulnar half of the hand, along with weakness and atrophy of the ulnar innervated muscles (5). In the Netherlands, however, most surgeons prefer electrodiagnostic confirmation prior to surgery (6). Patients would benefit from a diagnostic protocol that minimizes discomfort. In the present study, therefore, we analyzed patient data and constructed an efficient, minimal electrodiagnostic protocol for confirmation of clinically diagnosed UNE, based on a prospective electrodiagnostic study in patients with clinically defined UNE.

#### Methods

# Study design

We conducted a cross-sectional study among new patients (>18 years) who had been referred to our department for electrodiagnostic confirmation of the clinical diagnosis of UNE. We have used this database in previous studies (7). The only exclusion criteria was if the patient did not give consent to do all the NCS tests.

Criteria for the clinical diagnosis of UNE were numbness in the ulnar half of the fourth and fifth fingers, numbness in the ulnar half of the hand (palmar and dorsal) and/or weakness of the muscles in the hand innervated by the ulnar nerve (5). Patients were analyzed according to standard clinical procedures. For those with a bilateral clinical diagnose of UNE, both arms were included. The electrodiagnostic protocol was used to confirm the diagnosis.

We further analyzed the contribution of each electrodiagnostic criterion to the diagnosis of UNE.

#### Ethical approval

The regional medical ethical commission (CMO Nijmegen – Arnhem) and the hospital ethical commission (LTC) approved this research project. Anonymised data were collected from patients receiving standard care. Given that no experimental procedures were applied, both ethical commissions deemed that no informed consent was necessary.

# Neurophysiological protocol

We followed the criteria of the American Association of Electrodiagnostic Medicine (AAEM) for UNE (8). A detailed description of the neurophysiological protocol for UNE has been published previously (7).

Motor and sensory NCS of the ulnar nerve were performed. At four sites the ulnar nerve was stimulated: wrist (W), below the elbow (BE), above the elbow (AE) and at the bicipital sulcus. A tape-measure was used to measure conduction distances, with an accuracy of 5 mm (9). The conduction distance across the elbow was predetermined at 8 cm (9). Based on previous research, this method and this distance was used to get best balance between effects of measurement-error in short distances and 'dilution' of the slowing due to long distances (9). The elbow was kept flexed at 90 degrees during the examination (10). Stimulation was controlled to be supramaximal. Surface electrodes were used to record compound muscle action potentials (CMAP). Sensory nerve action potentials (SNAP) were recorded antidromically using ring electrodes positioned around the fifth finger, with an electrode distance of 4 cm, but less in smaller fingers (range 2–4 cm) CMAP and SNAP amplitudes were measured from negative to positive

Abbreviations: UNE, ulnar nerve entrapment at the elbow; NCS, nerve conduction studies; NCV, nerve conduction velocity; ADV, abductor digiti quinti muscle; FDI, first dorsal interosseous muscle; CMAP, compound muscle action potential; SNAP, sensory nerve action potential; EDX, Electrodiagnostic studies; AAEM, American Association of Electrodiagnostic Medicine; AECU, AAEM EDX confirmed ulnar neuropathy.

top in mV and  $\mu$ V, respectively. All latencies were measured from stimulus to onset deflection from baseline. Special care was taken to find the optimal positions of the "active" recording electrode above the hypothenar and first dorsal interosseal space (FDI) by shifting its position during stimulation, in order to achieve a maximal amplitude and an initial deflection that was as sharp as possible. During the whole procedure, CMAP configuration was observed in order to minimize or, if necessary, correct an altered position of the hand. The conduction velocity of all segments was computed. The arm was warmed by warming pads prior to all tests (11). Target temperature was 34 °C, with a lower limit of 30 °C. Skin temperature was monitored by an infrared thermometer device. Cooled regions were warmed again.

Based on our previous findings (7), we did not include our needle EMG data. As stated in this previous paper, we used this neurophysiological approach exclusively for the confirmation of clinical UNE in a high prior-odds diagnostic setting. EMG has limited value for diagnosis UNE if NCS are normal, but is necessary for evaluating the differential diagnosis, this is however outside the scope of this paper.

We categorized and analyzed the parameters presented in Table 1.

# Statistical analyses

All statistical analyses were performed using SPSS version 29 (*Statistical Package for Social Sciences, Chicago, IL, USA*). We used descriptive statistics to analyze the demographic characteristics and results of motor and sensible NCS. Furthermore, we have used binomial tests to compare the results of AAEM EDX confirmed ulnar neuropathy (AECU) group with the whole group.

#### Results

The sample included a total of 210 arms: 96 (45.7%) right; 114 (54.3%) left (see Table 2). The study population consisted of 91 men (43.3%) and 119 women (56.7.3%). The mean age of the patients was  $54.5 \pm 14.5$  years. The NCS results were abnormal for 127 (60.5%) of the arms analyzed.

An overview of the NCS abnormalities found is presented in Table 3. Abnormalities in the ADV were present in 84.3% of the cases

TABLE 1 Electrodiagnostic criteria for UNE.

М		

Stimulation at the wrist: absent CMAP of the ADV and/or FDI

Over the elbow (AE-to-BE) segment motor-NCV 10 m/s slower compared to lower arm segment (BE-to-wrist [W])

Decrease in ADV and/or FDI CMAP negative peak amplitude from BE to AE more than 20%.

#### Sensory

Absent distal SNAP (wrist)

Over the elbow (AE-to-BE) segment sensory-NCV 10 m/s slower compared to lower arm segment (BE-to-wrist [W])

ADV, abductor digiti quinti muscle; FDI, first dorsal interosseous muscle; CMAP, compound muscle action potential; SNAP, sensory nerve action potentials. Adapted from Kurver et al. (6).

TABLE 2 Demographic characteristics.

Demographic characteristics	N = 210	
Male-to-female ratio	91:119 (43.3%:56.7%)	
Age (years)		
Mean ± standard deviation	53.7 ± 14.5	
Range	19-91	
Affected side (R/L)	96/114 (45.7%/54.3%)	
UNE diagnosis according to AAEM criteria	127 (60.5%)	

of AAEM EDX confirmed ulnar neuropathy (AECU), with FDI and sensory abnormalities found less frequently (68.5 and 59.1%, respectively). The distribution of NCS abnormalities is presented in Figure 1.

The contribution of each criterion within the group of patients with NCS abnormalities is displayed in Table 4. Slowing of the motor nerve conduction velocity (NCV) was the most frequent NCS abnormality (89.9%; see Table 4), followed by slowing of the sensory NCV (44.1%; see Table 4). The other criteria were abnormal in only a small number of cases.

TABLE 3 Overview of NCS abnormalities.

NCS abnormalities	Frequency	% of total group (N = 210)	% of AECU group (n = 127)	<i>p</i> -value
ADV abnormalities	107	51.0%	84.3%	< 0.001
FDI abnormalities	87	41.4%	68.5%	<0.001
Sensory abnormalities	75	35.7%	59.1%	< 0.001
No NCS abnormalities	83	39.5%	0	n.a.

NCS, nerve conduction studies; ADV, abductor digiti quinti muscle; FDI, first dorsal interosseous; AECU, AAEM EDX confirmed ulnar neuropathy.

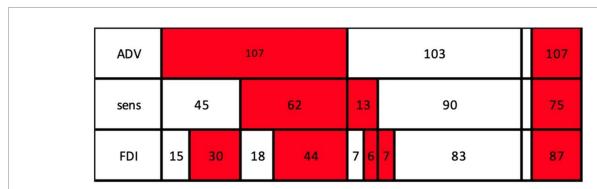


FIGURE 1

Distribution of NCS abnormalities. Red = abnormal NCS. White = normal NCS. ADV, abductor digiti quinti muscle NCS; FDI, first dorsal interosseous muscle NCS; sens, sensory NCS.

TABLE 4 Contribution of each criterium.

Motor				
		Frequency	Percent	
Absent wrist site CMAP of the:	ADV	3	2.4%	
	FDI	3	2.4%	
	ADV or FDI	4	3.1%	
An AE-to-BE segment greater than 10 m/s slower than the BE-to-wrist (W) segment of the:	ADV	103	81.1%	
	FDI	83	65.4%	
	ADV or FDI	114	89.8%	
Decrease in CMAP negative peak amplitude from BE to AE greater than 20% of the:	ADV	1	0.8%	
	FDI	1	0.8%	
	ADV or FDI	2	1.6%	

Sensory		
	Frequency	Percent
Absent distal SNAP	9	7.1%
An AE-to-BE segment greater than 10 m/s slower than the BE-to-wrist (W) segment of the sensory nerve velocity	56	44.1%

 $ADV, abductor\ digiti\ quinti\ muscle; FDI,\ first\ dorsal\ interosseous\ muscle; CMAP,\ compound\ muscle\ action\ potential.$ 

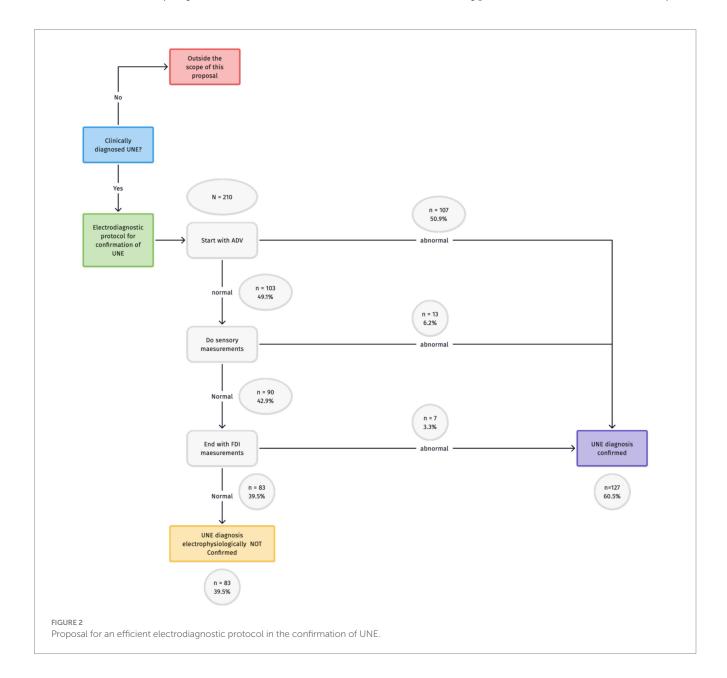
In the AAEM guideline another criterium is mentioned for the diagnosis of UNE: "Absolute motor nerve conduction velocity (NCV) from above elbow (AE) to below elbow (BE) of less than 50 m/s" (7). We have analyzed this criterium for the ADV muscle and we found only three patients with an absolute NCV in this segment less than 50 m/s in the group without the UNE diagnosis. All these three patients had only a slightly decreased NCV of 47 m/s.

#### Discussion

As reported in our previous studies, electromyographic testing for the presence of denervation in the absence of conduction abnormalities does not contribute to the electrodiagnostic confirmation of clinically diagnosed UNE (7). In the current study, our objective was to further optimize the electrodiagnostic protocol for confirmation of clinically diagnosed UNE. All results of the current study are presented in Figure 1, and all possible combinations can be derived. For cases requiring only one abnormal test for confirmation, NCS of the ADV was most promising (107 patients). For cases in which NCS of the ADV is normal, NCS of FDI or sensory NCS can follow. Both FDI and sensory NCS were abnormal in 13 patients (when ADV was normal).

For cases requiring at least two abnormal nerve conduction tests, the combination of NCS of the ADV and the FDI yielded the most abnormal test results (74 of the 107 patients), in contrast to the combination of ADV and sensory tests, which yielded abnormal results for only 62 of the 107 patients in this group. The combination of NCS of the FDI and sensory test identified only 50 patients with abnormal tests. If three abnormal tests are required for diagnosis, only 44 patients would meet this criterium.

The results of our investigation are presented in Figure 2. Given that the most abnormalities were found in NCS of the ADV, this should be the starting point. If these results are normal, sensory NCS



should follow. If the results of both of these tests are normal, the final step should be NCS of the FDI.

The choice of sensory NCS as the second step is not based solely on the numbers, given that sensory and FDI NCS are equally likely to yield abnormal results. In our experience, however, arriving at supramaximal stimulation in sensory NCS requires less stimulation strength than is the case for motor NCS. Patients might perceive this as less painful. However, the difference between pain perception in motor and sensory NCS is, to our knowledge, never investigated. Furthermore, given that the numbers are equal and the clinical symptoms are mainly sensory, we consider it appropriate to choose sensory NCS as a second step.

We must nevertheless note that, for cases requiring two abnormal tests, a combination of a motor and a sensory test is likely to be preferable to two motor tests, as the clinical symptoms are mainly sensory. Based on our data, however, the combination of two motor tests (ADV and FDI) yielded more abnormal results than did the combination of ADV and sensory NCS (74 vs. 62, respectively; see Figure 1).

In light of our preference for a minimal electrodiagnostic protocol for confirmation, one abnormal test should be sufficient; we propose this minimal electrodiagnostic protocol in a setting with well-defined clinically examined patients with the clinical diagnosis of UNE. If the clinical symptoms have a broad differential diagnosis (i.e., radiculopathy, plexopathy, polyneuropathy) a more extensive protocol including also electromyography is mandatory (12).

This study has several limitations that should be acknowledged. Firstly, we did not have specific exclusion criteria such as diabetes mellitus, prior surgery, or radiculopathy. The absence of these exclusions may have introduced confounding factors that could influence the electrodiagnostic findings and the generalisability of our results.

Secondly, the order of the diagnostic protocol was determined by the authors, with sensory NCS as the second step. However, this decision was not based on robust scientific evidence beyond the rationale provided above, which may limit the reproducibility of our approach in other settings.

Furthermore, it is noteworthy that 39.5% of patients in our cohort demonstrated normal findings on electrodiagnostic testing, despite a clinical diagnosis of ulnar neuropathy.

For such cases, we suggest using ultrasound of the ulnar nerve as the next step (12–14). If both of these tests are normal, another diagnosis should be considered. If the clinical symptoms are still consistent with ulnar neuropathy and possible further differential diagnosis have been excluded with more extensive neurophysiological studies including electromyography, the diagnosis can still be made, given the possibility of false negatives in both NCS and ultrasound. Within the context of CTS, it has been demonstrated that patients with negative NCS and ultrasound tests can still benefit from CTS treatment (15). The same could be true for patients with ulnar neuropathy, although further research is necessary to confirm this.

# Conclusion

We propose starting with NCS of the ADV for cases requiring only one abnormal test to confirm the clinical diagnosis of UNE. If these results are normal, sensory NCS should follow. For cases requiring at least two abnormal tests, the combination of NCS of the ADV and FDI would be most useful. If no abnormalities are found

in NCS, an ultrasound of the ulnar nerve should be the next step. If both tests are normal, another diagnosis should be considered and more extensive electrophysiological studies including electromyography must be done. If the clinical symptoms are still consistent with ulnar neuropathy, however, normal test results should not completely rule out this diagnosis, given the possibility of false negative results in both of these tests.

# Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors without undue reservation.

#### **Ethics statement**

The studies involving humans were approved by the regional medical ethical commission (CMO Nijmegen – Arnhem) and the hospital ethical commission (LTC) approved this research project. The studies were conducted in accordance with the local legislation and institutional requirements. Written informed consent for participation was not required from the participants or the participants' legal guardians/next of kin in accordance with the national legislation and institutional requirements.

#### **Author contributions**

AK: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Project administration, Visualization, Writing – original draft, Writing – review & editing. JM: Conceptualization, Supervision, Writing – review & editing. RB: Writing – review & editing. WV: Conceptualization, Methodology, Supervision, Writing – review & editing.

#### Funding

The author(s) declare that no financial support was received for the research and/or publication of this article.

#### Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

#### Generative AI statement

The authors declare that no Gen AI was used in the creation of this manuscript.

Any alternative text (alt text) provided alongside figures in this article has been generated by Frontiers with the support of artificial intelligence and reasonable efforts have been made to ensure accuracy, including review by the authors wherever possible. If you identify any issues, please contact us.

#### Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated

organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

#### References

- 1. Norkus SA, Meyers MC. Ulnar neuropathy of the elbow. Sports Med. (1994) 17:189-99.
- 2. Mondelli M, Giannini F, Ballerini M, Ginanneschi F, Martorelli E. Incidence of ulnar neuropathy at the elbow in the province of Siena (Italy). *J Neurol Sci.* (2005) 234:5–10. doi: 10.1016/j.jns.2005.02.010
- 3. Latinovic R, Gulliford MC, Hughes RAC. Incidence of common compressive neuropathies in primary care. *J Neurol Neurosurg Psychiatry*. (2006) 77:263–5. doi: 10.1136/jnnp.2005.066696
- 4. Bartels RHMA, Verhagen WIM, Van Der Wilt GJ, Meulstee J, Van Rossum LGM, Grotenhuis JA. Prospective randomized controlled study comparing simple decompression versus anterior subcutaneous transposition for idiopathic neuropathy of the ulnar nerve at the elbow: part 1. *Neurosurgery*. (2005) 56:522–30. doi: 10.1227/01.NEU.0000154131.01167.03
- 5. Kurver A, Scherf SM, Meulstee J, Verhagen WIM. The necessity of electrodiagnostic studies and ultrasound in ulnar nerve entrapment according to surgeons in the Netherlands. *Clin Neurol Neurosurg*. (2024) 236:108078. doi: 10.1016/j.clineuro.2023.108078
- 6. Kurver A, Smolders J, Verhagen WIM, Meulstee J, Nijhuis FAP. The diagnostic sensitivity for ulnar neuropathy at the elbow is not increased by addition of needle EMG of ADM and FDI when nerve conduction studies are normal. *Front Neurol.* (2019) 10:10. doi: 10.3389/fneur.2019.00196
- 7. AAEM Quality Assurance Committee. Practice parameter for electrodiagnostic studies in ulnar neuropathy at the elbow. Guidelines in electrodiagnostic medicine (2015). Update from: AAEM, practice parameter for electrodiagnostic studies in ulnar neuropathy at the elbow, summary statement. *Muscle Nerve*. (1999) 22:408–11. doi: 10.1002/(sici)1097-4598(199903)22:3<408::aid-mus16>3.0.co;2-7

- 8. Bartels RHMA, Menovsky T, Van Overbeeke JJ, Verhagen WIM. Surgical management of ulnar nerve compression at the elbow: an analysis of the literature. *J Neurosurg.* (1998) 89:722–7. doi: 10.3171/jns.1998.89.5.0722
- 9. Gert van Dijk J, Meulstee J, Zwarts MJ, Spaans F. What is the best way to assess focal slowing of the ulnar nerve? *Clin Neurophysiol.* (2001) 112:286–93. doi: 10.1016/S1388-2457(00)00549-6
- 10. Kothari MJ, Preston DC. Comparison of the flexed and extended elbow positions in localizing ulnar neuropathy at the elbow. *Muscle Nerve*. (1995) 18:336–40. doi: 10.1002/mus.880180312
- 11. Kasius KM, Riphagen JH, Verhagen WIM, Meulstee J. An easily applicable alternative method for warming cold limbs in nerve conduction studies. *Neurophysiol Clin.* (2014) 44:219–26. doi: 10.1016/j.neucli.2014.03.001
- 12. Pelosi L, Arányi Z, Beekman R, Bland J, Coraci D, Hobson-Webb LD, et al. Expert consensus on the combined investigation of ulnar neuropathy at the elbow using electrodiagnostic tests and nerve ultrasound. *Clin Neurophysiol.* (2021) 132:2274–81. doi: 10.1016/j.clinph.2021.04.018
- 13. Beekman R, Visser LH, Verhagen WI. Ultrasonography in ulnar neuropathy at the elbow: a critical review. *Muscle Nerve.* (2011) 43:627–35. doi: 10.1002/mus.22019
- 14. Pelosi L, Mulroy E. Diagnostic sensitivity of electrophysiology and ultrasonography in ulnar neuropathies of different severity. Clin Neurophysiol. (2019) 130:297–302. doi: 10.1016/j.clinph.2018.11.018
- 15. De Kleermaeker FGCM, Meulstee J, Claes F, Kasius KM, Verhagen WIM. Treatment outcome in patients with clinically defined carpal tunnel syndrome but normal electrodiagnostic test results: a randomized controlled trial. *J Neurol.* (2017) 264:2394–400. doi: 10.1007/s00415-017-8637-2