Unusual entrapment neuropathy in a golf player

To the Editor: The article by Hsu et al. provided a very elegant demonstration of the value of the technique of inching in the localization of focal lesions of motor and sensory axons and the ability of this technique to characterize the nature of the lesion as demyelinating or axonal. However, the demyelinating is producing focal axonal slowing of conduction, fiber blocking, or both may also be demonstrated and the percentage of total fibers blocked can be estimated. The presence and extent of axonotmesis may also be determined. This information has important prognostic (as the authors point out) and treatment implications and can be obtained in the EMG evaluation of any focal nerve lesion when stimulation proximal and distal to the lesion is possible (e.g., carpal tunnel syndrome [CTS]). The authors feel that the lesion location in their patient (between 2 and 3 cm distal to the distal wrist crease) is “unusual” and speculate on the mechanism of its production.

In our study of 122 patients with a clinical diagnosis of CTS (which was bilateral in 79) the lesion was between 2 and 3 cm distal to the distal wrist crease in 44.5% of the sensory studies and 21.3% of the motor studies. It is a common experience among hand surgeons and electromyographers that patients with long-standing minimal CTS often experience a marked increase in symptoms after unaccustomed heavy hand usage of different types and for varying periods. Sometimes the extent of this usage is less than the reported patient experienced. Patients with no previous symptoms of CTS may also experience such symptoms with comparable increased hand usage. The symptoms of the patient Hsu et al. describe are compatible with CTS. We believe that it is likely that this patient’s median nerve lesion resulted from the usual factors producing CTS.

We agree that compression of the median nerve between 2 and 3 cm distal to the wrist crease is not unusual in patients with CTS, as shown by their study. However, median nerve neuropathy in the palm is uncommon in golf players, more so compression by the first metacarpal bone. The association of median neuropathy and golf practice is clear in two aspects: the temporal relationship and conduction block on the nerve conduction study. This patient developed symptoms after playing golf, which he had not experienced before, and his symptoms remitted after he stopped playing. Further, the clinical consequences of conduction block are weakness and loss of sensation. This patient did not have these symptoms. Therefore, the conduction block shown in our case developed after golfing.

It is difficult to speculate whether this patient had minimal median neuropathy in the carpal tunnel that was aggravated by playing golf or not. This young man did not have any of the well-described risk factors for developing CTS (i.e., obesity, other medical problems, occupation involving repetitive wrist movement, and sex). It remains uncertain how much the transverse carpal ligament contributed to the compression of the median nerve. This is the reason why we preferred median neuropathy in the palm rather than CTS in this case.

What we really wanted to emphasize was the importance of palmar stimulation and that this can distinguish focal demyelination with conduction block, which has a very good prognosis from axonal degeneration. Further, better technique of holding the golf club may prevent this unusual focal neuropathy.

Wei-Chih Hsu, MD, Taipei, Taiwan, Agyepong Oware, MRCP, Bristol, UK

Reply from the Authors: We appreciate the interest in and comments on our report by Drs. White and Johnson. They have two comments: the entrapment site is not unusual, and they speculate that there could have been a pre-existing minimal carpal tunnel syndrome (CTS) that contributed to our patient having median nerve compression.

We agree that compression of the median nerve between 2 and 3 cm distal to the wrist crease is not unusual in patients with CTS, as shown by their study. However, median nerve neuropathy in the palm is uncommon in golf players, more so compression by the first metacarpal bone. The association of median neuropathy and golf practice is clear in two aspects: the temporal relationship and conduction block on the nerve conduction study. This patient developed symptoms after playing golf, which he had not experienced before, and his symptoms remitted after he stopped playing. Further, the clinical consequences of conduction block are weakness and loss of sensation. This patient did not have these symptoms. Therefore, the conduction block shown in our case developed after golfing.


References

Reply from the Authors: We appreciate the critical comments on our paper “Effects of subthalamic nucleus (STN) stimulation on motor cortex excitability.” The authors of the letter have several concerns that we would like to address.

Regarding methodology, we agree that age has a substantial influence on motor cortex excitability. We will publish a paper focusing on this issue in the near future. We have examined two different age groups in this study using paired-pulse transcranial magnetic stimulation (3 vs 13 ms interstimulus interval) and found that intracortical inhibition was significantly greater in older subjects. This result, however, is different from previous reports suggesting a decrease of intracortical inhibition with on-
point out the fact that our experimental protocol and the sample
886
NEUROLOGY 60
886
the receiver
operator characteristic curve, which relates the spec-
–
–
limited value and potentially misleading.
–
–
risk-to-benefit ratio. Unfortunately, the analysis performed is of
considerable risks. Effective predictors could favorably shift the
decision to operate on the oldest
–
–
correlation in middle-aged humans: a study using paired-
–
–
Regarding our results, this issue has been addressed in the
discussion section of our paper in detail.1 We just would like to
point out the fact that our experimental protocol and the sample
for PD patients were different from previous studies,2 possibly ac-
counting for different findings.
–
–
Regarding the discussion, we are thankful for their addi-
tional suggestions. However, at this stage all explanations are
rather hypothetical; an open discussion on this matter would be
appreciated.
–
–
J. Däuper, C. Schrader, J.D. Rolnik, Hannover, Germany
Copyright © 2003 by AAN Enterprises, Inc.

References
532.
3. Cunic D, Reshan L, Khan FI, Lozano AM, Lang AE, Chen R. Effects of
subthalamic nucleus stimulation on motor cortex excitability in Parkin-
internal globus pallidus restores intracortical inhibition in Parkinson’s disease paralleling apomorphine effects: a paired magnetic stimulation study.
6. Kossev AR, Schrader C, Däuper J, Dengler R, Rolnik JD. Increased
intracortical inhibition in middle-aged humans: a study using paired-
7. Peinemann A, Lehner C, Conrad B, Siebner HR. Age-related decrease in
paired-pulse intracortical inhibition in the human primary motor cortex.
8. Siggelkow S, Kossev A, Meil C, Dengler R, Rolnik JD. Impaired sensori-
motor integration in cervical dystonia: a study using TMS and muscle
9. Lewis GN, Byblow WD. Altered sensorimotor integration in Parkinson’s

Predictors of effective bilateral subthalamic
nucleus stimulation for PD

To the Editor: Charles et al. make an important contribution with their article describing possible predictors of deep brain stim-
ulation (DBS) of the subthalamic nucleus efficacy for PD.1 While
DBS is highly effective and FDA approved, the procedure has considerable risks. Effective predictors could favorably shift the risk-to-benefit ratio. Unfortunately, the analysis performed is of limited value and potentially misleading.

A more appropriate analysis would be to report the area under the receiver–operator characteristic curve, which relates the specific-
ity and sensitivity of the tests to age and levodopa responsive-
ness. The goal of any predictive task not only is to avoid surgery for those patients not likely to benefit but also to avoid withhold-
ing surgery from those that would. Visual inspection of the data
represented in the graphs provides little confidence that either age or levodopa responsiveness will have sufficient specificity and sensitivity to be an effective predictor that can be used for patient selection.

In addition, the study of predictors was limited to a retrospec-
tive correlational analysis. Correlation is a mathematically optim-
izing procedure that will find a correlation, even if spurious.2 Thus, it remains unclear how generalized are the regression analyses
performed. That is why it is so important to apply the predic-
tive regression equations in a prospective manner. Often, dividing the sample population into two groups, the first to develop the regression equations and the second to prospectively test those equations, can do this. The large majority of times, the specific-
ity and sensitivity of predictors protectors fall when tested pro-
spectively.

Erwin B. Montgomery, Jr., MD, Cleveland, OH

Reply from the Authors: We thank Dr. Montgomery for his
comments and share his concern about the importance of ade-
quately selecting parkinsonian patients for surgery. Dr. Montgomery’s remark deals with the use of regression and correlational analyses to study the predictive factors of outcome from bilateral subthalamic nucleus stimulation. While it is true that the regression analysis of the data is retrospective, the original patient

enrollment, treatment, and data collection were performed pro-
spectively. The calculation of the sensitivity, specificity, and confi-
dence intervals would need considerably more patients, hardly compatible with this type of therapeutic procedure. The receiver–operating characteristic curves proposed by Dr. Montgomery are frequently used to assess the usefulness of diagnostic markers, but the method also has some disadvantages.3 We think that univariate analysis is one of the most appropriate statistical methods for our study. We agree with the necessity of validating our model in another prospective study. We do not know if “the large majority of times, the specificity and sensitivity of protectors fall when tested prospectively,” but it has been shown that it is not always true.4 Most studies of the surgical treatment of PD found that outcomes from surgery are better in patients with levodopa-
response motor symptoms. Welter et al.5 also used regression
analysis in their series of parkinsonian patients treated with subthalamic nucleus stimulation. In keeping with our results they found that the outcome of STN stimulation was excellent in levodopa-responsive forms of PD. Our results are consistent with the classic inclusion criteria for subthalamic nucleus stim-
ulation and imply that the decision to operate on the oldest patients and/or patients with levodopa-resistant motor symp-
toms should be carefully weighed. The other lesson from our experience is that parkinsonian patients with severe levodopa-
induced motor complications may still be surgical candidates if a fair levodopa response is maintained, i.e., if their best on-
motor score is low. This result is clinically sensible. The rela-
tive young age at the time of surgery could have been expected as a good predictor because young-onset PD is characterized by a good response to levodopa with minimal on-period axial or motor symptoms except fluctuations and dyskinesias.6 Moreover, surgery-related complications are more frequent in an elderly population.

Erwin B. Montgomery, Jr., MD, Cleveland, OH

P.D. Charles, MD, Nashville, TN; N. Van Blercom, MD, P. Krack, MD, Grenoble, France; S.L. Lee, MD, PhD, Nashville, TN; J. Xie, MD, PhD, G. Besson, MD, PhD, A.L. Benabid, MD, PhD, I. Pollak, MD, Grenoble, France

Copyright © 2003 by AAN Enterprises, Inc.
References


Correction

GDAP1 mutations in CMT4: Axonal and demyelinating phenotypes? The exception “proves the rule”

In the recently published editorial titled “GDAP1 mutations in CMT4: Axonal and demyelinating phenotypes: The exception “proves the rule,” (Neurology 2002;59:1835–1836) the authors inadvertently misstated a mutation. The text should have stated “myotubularin-related protein-2 (MTMR2).”
GDAP1 mutations in CMT4: Axonal and demyelinating phenotypes? The exception "proves the rule"

Neurology 2003;60;887
DOI 10.1212/WNL.60.5.887

This information is current as of March 11, 2003

Updated Information & Services
including high resolution figures, can be found at:
http://www.neurology.org/content/60/5/887.full.html

Permissions & Licensing
Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at:
http://www.neurology.org/misc/about.xhtml#permissions

Reprints
Information about ordering reprints can be found online:
http://www.neurology.org/misc/addir.xhtml#reprintsus