A Focus on Subtle Signs and Motor Behavior to Unveil Awareness in Unresponsive Brain-Impaired Patients

The Importance of Being Clinical

Karin Diserens, MD,* Ivo A. Meyer, MD,* Jane Jöhr, PhD, Alessandro Pincherle, MD, Vincent Dunet, MD, Polona Pozeg, PhD, Philippe Ryvlin, MD, PhD, Dafin F. Mureșanu, MD, PhD, Robert D. Stevens, MD, and Nicholas D. Schiff, MD

Abstract

Brain-injured patients in a state of cognitive motor dissociation (CMD) exhibit a lack of command following using conventional neurobehavioral examination tools but a high level of awareness and language processing when assessed using advanced imaging and electrophysiology techniques. Because of their behavioral unresponsiveness, patients with CMD may seem clinically indistinguishable from those with a true disorder of consciousness that affects awareness on a substantial level (coma, vegetative state/unresponsive wakefulness state, or minimally conscious state minus). Yet, by expanding the range of motor testing across limb, facial, and ocular motricity, we may detect subtle, purposeful movements even in the subset of patients classified as vegetative state/unresponsive wakefulness state. We propose the term of clinical CMD to describe patients showing these slight but determined motor responses and exhibiting a characteristic akinetic motor behavior as opposed to a pyramidal motor system behavior. These patients may harbor hidden cognitive capabilities and significant potential for a good long-term outcome. Indeed, we envision CMD as ranging from complete (no motor response) to partial (subtle clinical motor response) forms, falling within a spectrum of progressively better motor output in patients with considerable cognitive capabilities. In addition to providing a decisional flowchart, we present this novel approach to classification as a graphical model that illustrates the range of clinical manifestations and recovery trajectories fundamentally differentiating true disorders of consciousness from the spectrum of CMD.

*These authors contributed equally to this work as first authors.

From the Neurology and Acute Neurorehabilitation Unit (K.D., I.A.M., J.J., P.R.), Department of Clinical Neurosciences, and Department of Radiology (V.D., P.P.), Lausanne University Hospital and University of Lausanne; University Hospital of Old Age Psychiatry (I.A.M.), University of Bern, Switzerland; Neurology Unit (A.P.), Department of Medicine, Hôpitaux Robert Schuman, Luxembourg; Department of Neuroscience (D.F.M.), Lulu Hâţeganu University of Medicine and Pharmacy, Cluj-Napoca, Romania; Departments of Anesthesiology and Critical Care Medicine, Neurology, and Neurosurgery (R.D.S.), School of Medicine, Johns Hopkins University, Baltimore, MD; Feil Family Brain and Mind Research Institute (N.D.S.), Weill Cornell Medical College, New York, NY; and Department of Neurology (N.D.S.), New York Presbyterian Hospital.

Go to Neurology.org/N for full disclosures. Funding information and disclosures deemed relevant by the authors, if any, are provided at the end of the article.

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivatives License 4.0 (CC BY-NC-ND), which permits downloading and sharing the work provided it is properly cited. The work cannot be changed in any way or used commercially without permission from the journal.

Copyright © 2023 The Author(s). Published by Wolters Kluwer Health, Inc. on behalf of the American Academy of Neurology.
Diagnosing CMD: A Clinical and Technical Challenge

The existence of covert awareness was first demonstrated in 2006,4 and the term CMD was introduced in 20155 to describe patients with command following discernible by specific AIE-derived neural signatures but without externally observable motor responses. In a recent article in this journal, an algorithmic flowchart to determine when AIEs should be used was proposed.6 We welcome these efforts to establish criteria for whether AIEs are suitable on an individual patient basis. Currently, AIEs are not warranted in routine clinical practice if evidence of a conscious motor response is observed during bedside neurobehavioral assessment.

Ideally, AIEs should be used for patients with a higher probability of harboring covert awareness, although guidelines establishing such probabilities are only slowly emerging now. Performing AIEs and interpreting their results, especially in the acute setting, requires considerable technical and medical expertise. A possible solution to the logistics posed by these technologies might be a hub-and-spoke model, as proposed by Young et al.,7 whereby peripheral collaborating sites with less resources (i.e., spokes) collect AIE data locally and then send the data to a specialized medical center (i.e., hub) that provides the expertise for processing and analysis. This could help reduce geographic and financial gaps and guarantee the detection of covert awareness in patients who might otherwise be misdiagnosed. Still, there are at least 2 major limitations. First, this approach is currently only viable in health systems with sizeable economical resources. Second, given that patients in the acute setting often have considerable executive, attentional, or language dysfunction from which they would eventually recover, there is an unquantifiable risk of misclassifying such patients as lacking conscious awareness when strict AIE assessment protocols are used. Proving that an unresponsive patient is aware is hard, but proving that the patient is lacking awareness is harder, if not impossible.

Clinical Unmasking of Covert Awareness

We argue that a practical and immediate advance in detecting patients with apparent covert awareness can be achieved by further expanding the current clinical assessment scales. Recent studies indicate that a significant fraction of patients with covert awareness defined by the combination of traditional standardized neurobehavioral assessments and AIEs can be identified clinically, circumventing the need for AIEs in these patients.2,3 Current clinical assessment scales, including the thorough CRS-R, often fail to diagnose awareness in patients who show subtle signs of interaction, even in those retaining some motor localization of the painful stimulus, visual fixation, or visual tracking. In the acute care setting, the rate of patients with covert awareness misdiagnosed as lacking consciousness is at least as high as 15% when using the CRS-R as this is the percentage of unresponsive cases (as classified by the CRS-R) for which AIEs captured evidence of unequivocal brain activation in response to a command.8 In our experience, we estimate the misclassification rate to be around 30% when comparing the CRS-R diagnosis (e.g., vegetative state) at admission to an acute neurorehabilitation unit to the diagnosis at discharge.9

Patients with true disorders of consciousness are not motionless but display an array of reflexive behaviors and, after recovering fragments of awareness, purposeful motor behavior. These patients often progress through a prolonged or permanent confusional state, associated with motor hyperactivity (agitation) stemming from an underlying lack of orientation and incoherent cognition. On the cognitive level, such residual fragments of consciousness may be detected using the CRS-R as limited but overt and reproducible motor responses, with patients failing to regain consistent and accurate communication systems (via speech or gesture). In contrast to the motor patterns observed in patients with true disorders of consciousness, relatively focal lesions that globally affect the motor

Glossary

AIE = advanced imaging and electrophysiology technique; CMD = cognitive motor dissociation; CRS-R = Coma Recovery Scale–Revised; MBT-r = Motor Behavior Tool–Revised.
output channels can cause a characteristic lack of motor or verbal interaction in patients with CMD, hiding their considerable cognitive capability. As mentioned above, the residual signs in such motionless patients may be too subtle to be identified using the CRS-R alone. We have evaluated, and propose using, a complementary clinical tool, the MBT-r (see description in Table), designed to detect subtle motor behaviors that are overlooked by the CRS-R that establishes strict criteria for scoring a specific motor behavior as an expression of consciousness. In a prospective validation study, the MBT-r was shown to identify a subset of patients whose cognitive abilities were underestimated by the CRS-R and demonstrated excellent interrater agreement. Subsequent data from a sample of 141 patients undergoing inpatient rehabilitative care suggested that the MBT-r used in conjunction with the CRS-R leads to greater sensitivity in detecting awareness than the CRS-R alone and identified patients with a high probability of functional recovery at discharge.

**Scrutinize Clues and Pitfalls to Maximize the Clinical Detection Rate of Awareness**

During the assessment of unresponsive patients, it is essential to scrutinize the clues and pitfalls that may support the diagnosis of CMD. Bringing all this together, we created a flowchart that considers clinical, pathophysiologic, radiologic, and electrophysiologic aspects to establish an early diagnosis of either a true disorder of consciousness or CMD (Figure 1). We propose using the suggested flowchart as soon as possible, that is, in the intensive care unit 24 hours after sedation withdrawal, always considering residual anesthesia as a potential confounder. A comprehensive clinical assessment should be conducted at least 3 times a week in conjunction with pathophysiologic considerations and paraclinical investigations if necessary. As indicated in the flowchart, we use conventional

---

**Table** Motor Behavior Tool–Revised (MBT-r)

<table>
<thead>
<tr>
<th>Sign</th>
<th>Observations</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Positive signs</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Spontaneous nonreflexive movement</td>
<td>Any nonstereotypical, noncontextualized, and nonrepetitive intentional motor behavior</td>
<td>Observation of spontaneous behavior without stimulation at baseline or at any moment</td>
</tr>
<tr>
<td>2. Response to a command</td>
<td>Any nonreflective intentional response to a verbal command</td>
<td>Use the CRS-R command-following protocol</td>
</tr>
<tr>
<td>3. Visual fixation or visual pursuit</td>
<td>Any visual fixation or visual pursuit in any direction</td>
<td>Defined as eyes changing from an initial to a new fixation point or eyes tracking a moving target</td>
</tr>
<tr>
<td>4. Response in a motivational context</td>
<td>Any appearance or increase in the frequency of nonreflective motor response in a salient context</td>
<td>For example, on hearing a familiar voice, the patient’s mother tongue or the patient’s own or nickname</td>
</tr>
<tr>
<td>5. Response to a noxious stimulation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>5a. Defensive response: nipple sign</td>
<td>Any attempt of defense when twisting the patient’s nipple while holding the patient’s arm</td>
<td>Before scoring, exclude stereotypical posturing as a confounding factor</td>
</tr>
<tr>
<td>5b. Defensive response: nail bed sign</td>
<td>Any defense gesture to deep pressure applied to a nail bed (test all 4 extremities)</td>
<td>The kinematics of an intentional defense differ from those of a nociceptive withdrawal reflex</td>
</tr>
<tr>
<td>5c. Grimace</td>
<td>Any nonreflective grimace on administering a noxious stimulation</td>
<td>Do not score the reflexive rictus-like grimace of stereotypical posturing or tetanus</td>
</tr>
</tbody>
</table>

**Negative signs (brainstem release signs)**

<table>
<thead>
<tr>
<th>Sign</th>
<th>Observations</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>6. Decorticate posturing, decerebrate posturing, or incessant paroxysmal sympathetic hyperactivity</td>
<td>Spontaneous or stimulus-induced stereotypical posturing (decorticate or decerebrate posturing) or incessant neurovegetative responses (i.e., rapid-onset episodes of tachycardia, hypertension, tachypnea, fever, diaphoresis, and dystonic posturing of up to 30-min duration)</td>
<td>In the absence of brainstem lesions, these signs reflect large and widespread forebrain lesions resulting in functional disconnection of the red nucleus (decorticate posturing), of vestibulo-/tecto-reticulospinal postural reflexes (decerebrate posturing), or of neurovegetative centers</td>
</tr>
<tr>
<td>7. Pathologic conjugate roving eye movements (ping-pong gaze)</td>
<td>Resembles slow eye movements of light sleep; can persist with open or closed eyes, lack total excursion, or move from an extreme gaze to the midline instead of to the opposite extreme; and may present or lack pauses between excursions</td>
<td>Also called short-cycle periodic alternating gaze; caused by large and widespread lesions causing a disconnection between the cortical inhibitory control and brainstem gaze centers</td>
</tr>
</tbody>
</table>

Abbreviation: CRS-R = Coma Recovery Scale–Revised. A positive sign is scored even if subtle if it stands out clearly from a reflexive or stereotypical background movement. Note that repeatability is not necessary for the MBT-r: the observation of a single subtle intentional movement, for example, visual pursuit obviously discernible from the baseline eye movement, is scored as a positive sign. When in doubt, the sign is not recorded. To facilitate interpretation of subtle signs, whenever possible, patients are filmed with the mother tongue or the patient’s own or nickname.
Figure 1 Flowchart for Acute Assessment of Unresponsive Patients With a Suspected Major Cerebral Impairment

Unresponsive patient (absence of overt sign of interaction)

Always consider during clinical evaluation

Neurobehavioral assessment (CRS-R)

Clues and pitfalls that may support (clinical) cognitive motor dissociation diagnosis

Polyneuropathy or myopathy
Brainstem, cranial nerves
Nonconvulsive status epilepticus
Cortical blindness, Terson syndrome
Akinetic mutism
Posterior fossa syndrome
Thalamic involvement
Aphasia

Condition

Paroxysmal Clinical Signs

Areflexia, amyotrophy, flaccidity
Upper motor neuron signs, cranial nerve palsies
Staring, gaze deviation, neglect, myoclonus
No visual interaction
Marked reduction of movements and speech
Marked reduction of movements and speech
Arousal fluctuations, thalamic, arrhythmic syndromes
No command following, mutism

Electroencephalograph Imaging

Structural imaging
EEG
Structural imaging
EEG
Structural imaging
EEG
Structural imaging
EEG

Language-mediated behaviors?

No

Coma, UWS/VVS, or MCS- (based on CRS-R)

≥1 positive sign on MBT-r?

≥1 negative sign on MBT-r?

MRI (repeat if inconclusive)

MRI (repeat if inconclusive)

≥1 positive sign on MBT-r?

Confusion assessment method or neuropsychologic evaluation

Widespread cortico-thalamic damage?

True disorder of consciousness: MCS+
Confusional state or cognitive disability
True disorder or consciousness: MCS-

Circumscribed disruption of modulatory tracts (e.g., in brainstem) causing negative MBT-r signs

MRI (repeat if inconclusive)

Damage to motor outflow system with relative preservation of cortico-sub cortical networks relevant for awareness

ictal, encephalopathic (metabolic, excessive sedation), or flat EEG pattern

Widespread cortico-thalamic damage?

Confusion assessment method or neuropsychologic evaluation

Widespread cortico-thalamic damage?

MRI (repeat if inconclusive)

MRI (repeat if inconclusive)

Widespread cortico-thalamic damage?

No

True disorder of consciousness: MCS–
Confusional state or cognitive disability
True disorder or consciousness: MCS–

Circumscribed disruption of modulatory tracts (e.g., in brainstem) causing negative MBT-r signs

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?

≥1 positive sign on MBT-r?
structural MRI as an adjunctive examination to dichotomize patients into true disorders of consciousness or clinical CMD.12 It is not our intention to challenge the importance of AIEs. In its most complete form, that is, in the complete absence of motor response, CMD is only uncovered if task-based AIEs demonstrate evidence of command following. Resting-state AIE examinations also play an important role, and the pursuit of covert awareness should include an investigation of the functional architecture dynamics using resting-state functional MRI and PET, concentrating primarily on the brain’s ability to shift between the internal (default mode) and external awareness (frontoparietal) networks. The brain’s aptitude to switch between intrinsic and extrinsic network activation has been associated with recovery of consciousness13 and could potentially be used as a biomarker for covert awareness. Neurophysiologic evaluations such as non–task-evoked potentials help to discriminate clinical CMD (especially with aphasia and attention deficit) from patients with a true disorder of consciousness,14 thus contributing to a better understanding of the underlying network mechanisms.

Figure 2 Model of Behavioral and Cognitive Evolution After Severe Brain Impairment

The diagnostic spectrum and clinical evolution after severe brain impairment is represented on a 2-dimensional graph comparing the degree of cognitive content (x-axis) against the degree of behavioral response (y-axis). The red-yellow-green color gradient represents an approximation of the degree of recovery, with red being the worst and green the best. Both cognitive content and behavioral response are determined using the CRS-R in conjunction with the MBT-r, which we developed to expand the range of motor testing to detect subtle, purposeful movements (the so-called positive signs) and brainstem release signs (the so-called negative signs). Concerning the motor/behavioral response of patients with CMD, the functional ambulation category quantifies the degree of functional motor recovery. In cases of a total lack of motor response, AIEs may unveil CMD-defining covert cognition. For patients with a good behavioral response, confusion and neuropsychological assessment protocols distinguish between the different degrees of functional recovery. Patients with a severe brain impairment appear to fall mainly into 2 categories—true disorders of consciousness (spanning from VS/UWS to MCS+, red hues) and CMD/locked-in state (yellow-green hues)—with different underlying lesions (represented by the symbolic brain images), clinical manifestations (e.g., brainstem release signs), and prognosis (represented by the arrows along the recovery pathways). The circles/ellipses surrounding the different diagnoses (VS/UWS, MCS-, etc.) account for the fact that many patients never recover after a specific point in the recovery pathway. Widespread lesions across heteromodal cortical association areas and cortico-thalamic tracts cause the impairment in patients with true disorders of consciousness. Consequently, many of these patients may suffer chronically from a total (VS/UWS) or partial (MCS) lack of awareness. Posturing typically lasts after conscious emergence, causing characteristic clinical motor patterns usually not seen in patients with CMD. Patients with CMD, on the other hand, are characterized by a total (complete CMD, with covert cognition only detectable by AIEs) or near-total (clinical CMD with subtle, purposeful movements) lack of determined movements. Most patients with CMD do not develop brainstem release signs, except for complete locked-in state patients, who typically manifest decorticate/decerebrate posturing. We indicate the potential recovery pathways using symbolic arrows. AIE = advanced imaging and electrophysiology technique; CMD = cognitive motor dissociation; CRS-R = Coma Recovery Scale–Revised; MBT-r = Motor Behavior Tool–Revised; MCS = minimally conscious state (+ = plus and − = minus); VS/UWS = vegetative state/unresponsive wakefulness state. Created using the vector graphics editor Adobe Illustrator (Adobe Inc., San Jose, CA). Brain images were derived from the population-averaged tractography atlas by Yeh et al.24
A Spectrum of Clinical Manifestations and Recovery Trajectories: Neuroanatomic and Physiologic Rationale for a New Model

Given that brain function in patients with CMD is likely to be closer to that of healthy and locked-in state patients than to minimally conscious state plus patients with reliable command following,15,16 we suggest that there is a dividing line separating true disorders of consciousness from CMD and locked-in state patients. However, we propose this partition should be considered a gradient rather than a sharp boundary. In other words, although we should avoid simplistic dichotomizations, this concept forms the basis of a new approach to classification in which unresponsive patients with CMD are in a distinct category from unresponsive patients without CMD. We envision a spectrum of different motor/cognitive states, ranging from complete CMD that cannot be detected even by extended clinical testing to partial CMD (clinical CMD) with subtle clinical signs and typical motor behavior to classical locked-in state (preservation of vertical eye movements/eye blinking) (Figure 2).

The cognitive abilities of patients with CMD may range from the limited language comprehension functions of minimally conscious state plus patients to the almost intact cognitive capabilities of complete locked-in state patients.17 This raises the question of how to distinguish patients with CMD at the lower end of the cognitive spectrum from those with true disorders of consciousness who have regained some command-following abilities (e.g., minimally conscious state patients). Based on pathophysiological considerations, a key feature clinically dichotomizing most CMD and true disorders of consciousness might be the presence of brainstem release signs in the absence of brainstem lesions. Such release signs imply widespread destruction of the cortico-cortical networks relevant for awareness. However, it is essential to initially exclude additional brainstem or cerebellar lesions as these can complicate the clinical assessment for at least 2 reasons. First, these lesions can disrupt the ascending arousal system, affecting wakefulness despite the relative intactness of structures essential for awareness. Second, circumscribed lesions of the corticospinal tract and the tracts connecting the higher modulatory centers and the rubral and/or vestibular nuclei in the rostral part of the brainstem can cause tetraplegia with decorticate or decerebrate posturing, respectively, which may be falsely attributed to larger lesions located more cranially.

Although the exact nature of awareness remains an unsolved complex problem of neuroscience, it is a plausible assumption that it requires the activation of large-scale decentralized cortico-cortical networks.18 This decentralization increases the robustness of awareness against focal damage. It also means that large and widespread lesions will be necessary to significantly disrupt this fundamental function of the brain.12 Classically, neurologic practice has considered bilateral lesions of the cortex or of the upper brainstem and central thalamus or lesions altering both as a requirement for producing coma.19 Because of such widespread lesions, a functional cortical disconnection is detected clinically as decorticate or decerebrate posturing,20 pathologic roving eye movements (ping-pong gaze, i.e., short-cycle periodic alternating gaze),21 and/or incessant paroxysmal sympathetic hyperactivity.22 These clinical manifestations have long been associated with poor outcome and, on a pathophysiological level, are likely caused by a disruption of modulatory (mainly inhibitory) corticorubral-spinal, cortico-vestibular, cortico-mesencephalic, and/or cortico-diencephalic tracts (see negative MBT-r signs in Table).

Patients within the CMD spectrum as opposed to those with true disorders of consciousness are not only characterized by specific clinical, radiologic, and pathophysiological features but, most importantly, by their particular recovery trajectories.9 An early diagnosis of CMD does not automatically imply a good outcome, as the individual prognosis depends heavily on the specific cerebral functions affected; but overall, patients with CMD have far better long-term outcomes, as measured by multiple prognostic scales.9,23

Conclusion

In conclusion, we want to underscore the importance of a thorough clinical assessment targeted to observing the motor behavior, as well as the role of careful clinical and paraclinical screening (Figure 1), selecting conventional structural MRI before proceeding to more sophisticated technical diagnostic tools. We present this approach as a model depicting the spectrum of clinical manifestations and recovery trajectories after significant brain impairment (Figure 2).

Acknowledgment

The authors are grateful to Dr. Melanie Price Hirt for proofreading this manuscript and Mr. Ehsan Faridi for significantly improving the graphical design of Figure 2. The authors also thank Dr. Edlow for his feedback on an earlier version of this work. The authors are particularly indebted to their forebears and mentors who always emphasized the importance of a proper clinical examination.

Study Funding

No targeted funding reported.

Disclosure

The authors report no disclosures relevant to the manuscript. Go to Neurology.org/N for full disclosures.

Publication History

Received by Neurology August 13, 2022. Accepted in final form January 3, 2023. Submitted and externally peer reviewed. The handling editor was Associate Editor Rebecca Burch, MD.
### Appendix Authors

<table>
<thead>
<tr>
<th>Name</th>
<th>Location</th>
<th>Contribution</th>
</tr>
</thead>
<tbody>
<tr>
<td>Karin Diserens, MD</td>
<td>Neurology and Acute Neurorehabilitation Unit, Department of Clinical Neurosciences, Lausanne University Hospital and University of Lausanne, Switzerland</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
<tr>
<td>Ivo A. Meyer, MD</td>
<td>Neurology and Acute Neurorehabilitation Unit, Department of Clinical Neurosciences, Lausanne University Hospital and University of Lausanne, University Hospital of Old Age Psychiatry, University of Bern, Switzerland</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content; additional contributions: creation of the graphical model and flowchart</td>
</tr>
<tr>
<td>Jane Jöhr, PhD</td>
<td>Neurology and Acute Neurorehabilitation Unit, Department of Clinical Neurosciences, Lausanne University Hospital and University of Lausanne, Switzerland</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
<tr>
<td>Alessandro Pincherle, MD</td>
<td>Neurology Unit, Department of Medicine, Hôpitaux Robert Schuman, Luxembourg</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
<tr>
<td>Vincent Dunet, MD</td>
<td>Department of Radiology, Lausanne University Hospital and University of Lausanne, Switzerland</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
<tr>
<td>Polona Pozeg, PhD</td>
<td>Department of Radiology, Lausanne University Hospital and University of Lausanne, Switzerland</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
<tr>
<td>Philippe Ryvlin, MD, PhD</td>
<td>Neurology and Acute Neurorehabilitation Unit, Department of Clinical Neurosciences, Lausanne University Hospital and University of Lausanne, Switzerland</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
<tr>
<td>Dafin F. Mureșanu, MD, PhD</td>
<td>Department of Neuroscience, St. John's Hospital, University of Medicine and Pharmacy, Cluj-Napoca, Romania</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
<tr>
<td>Robert D. Stevens, MD</td>
<td>Departments of Anesthesiology and Critical Care Medicine, Neurology, and Neurosurgery, School of Medicine, Johns Hopkins University, Baltimore, MD</td>
<td>Drafting/revision of the manuscript for content, including medical writing for content</td>
</tr>
</tbody>
</table>

### References

22. Rabstein AA. Parasympathetic Hyperactivity. UpToDate; 2022.
A Focus on Subtle Signs and Motor Behavior to Unveil Awareness in Unresponsive Brain-Impaired Patients: The Importance of Being Clinical

*Neurology* 2023;100;1144-1150 Published Online before print February 28, 2023
DOI 10.1212/WNL.0000000000207067

**This information is current as of February 28, 2023**

<table>
<thead>
<tr>
<th>Updated Information &amp; Services</th>
<th>including high resolution figures, can be found at: <a href="http://n.neurology.org/content/100/24/1144.full">http://n.neurology.org/content/100/24/1144.full</a></th>
</tr>
</thead>
<tbody>
<tr>
<td>References</td>
<td>This article cites 20 articles, 3 of which you can access for free at: <a href="http://n.neurology.org/content/100/24/1144.full#ref-list-1">http://n.neurology.org/content/100/24/1144.full#ref-list-1</a></td>
</tr>
<tr>
<td>Subspecialty Collections</td>
<td>This article, along with others on similar topics, appears in the following collection(s):</td>
</tr>
<tr>
<td></td>
<td><strong>All Imaging</strong> <a href="http://n.neurology.org/cgi/collection/all_imaging">http://n.neurology.org/cgi/collection/all_imaging</a></td>
</tr>
<tr>
<td></td>
<td><strong>All Rehabilitation</strong> <a href="http://n.neurology.org/cgi/collection/all_rehabilitation">http://n.neurology.org/cgi/collection/all_rehabilitation</a></td>
</tr>
<tr>
<td></td>
<td><strong>Clinical neurology examination</strong> <a href="http://n.neurology.org/cgi/collection/clinical_neurology_examination">http://n.neurology.org/cgi/collection/clinical_neurology_examination</a></td>
</tr>
<tr>
<td></td>
<td><strong>Coma</strong> <a href="http://n.neurology.org/cgi/collection/coma">http://n.neurology.org/cgi/collection/coma</a></td>
</tr>
<tr>
<td></td>
<td><strong>Prognosis</strong> <a href="http://n.neurology.org/cgi/collection/prognosis">http://n.neurology.org/cgi/collection/prognosis</a></td>
</tr>
<tr>
<td>Permissions &amp; Licensing</td>
<td>Information about reproducing this article in parts (figures,tables) or in its entirety can be found online at: <a href="http://www.neurology.org/about/about_the_journal#permissions">http://www.neurology.org/about/about_the_journal#permissions</a></td>
</tr>
<tr>
<td>Reprints</td>
<td>Information about ordering reprints can be found online: <a href="http://n.neurology.org/subscribers/advertise">http://n.neurology.org/subscribers/advertise</a></td>
</tr>
</tbody>
</table>