Operationalizing Neuroimaging for Disorders of Consciousness: The Canadian Context

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Severe brain injury affects tens of thousands of Canadians each year, leaving a small but significant proportion who survive the acute phase in a vegetative state (VS) or minimally conscious state (MCS). Patients in both VS and MCS have diurnal patterns of sleep-wake cycles, but have no behavioral response (VS) or only an inconsistent response (MCS) to stimuli. New technological capabilities offer a window into these disorders—not only changing our understanding of the levels of consciousness, but also enabling the potential for better diagnosis. For example, studies using functional magnetic resonance imaging, now extending over a decade and involving many participants, have revealed that a small minority of patients with severe disorders of consciousness (DoC) can modulate their neural states and thereby demonstrate a limited degree of awareness and even reasoning comparable to unaffected persons.1,2

The governments of Canada, its provinces, and territories have recognized and enshrined in legislation and the Constitution moral obligations to persons with disabilities based on the fundamental human rights of autonomy and equality, among others. The right to equality and nondiscrimination is reflected in the Canadian Charter of Rights and Freedoms (section 15), the United Nations Convention on the Rights of Persons with Disabilities (UNCRPD), and provincial human rights legislation requiring reasonable accommodation for persons with disabilities. Canada’s 2010 ratification of the UNCRPD marks a fundamental change in the way Canadian society views persons with disabilities in that they are “no longer considered to be recipients of charity or objects of others’ decisions, but holders of rights.”3,4 This rights-centric view would apply equally to MCS and VS patients as to any other group with disabilities. Given Canada’s moral and legal commitment to persons with disabilities, the evidence of awareness in some patients with disorders of consciousness, and the results of prior phases of research that suggest broad conceptual support from physicians, researchers, and legal experts,5 we believe that Canada, as a society, is obliged to explore whether and how neuroimaging technology could be made available to patients with DoC. To this end, we prioritize the following steps for action, that both newly complement and build upon recommendations advanced by other scholars.

RESEARCH

1. Correlate neuroimaging results with behavioural measures. The Glasgow Coma Scale, the Coma Recovery Scale – Revised, and the Wessex Head Injury Matrix are behavioural measures used to assess patients with disordered consciousness. Even with these tools, historically high rates of misdiagnosis underscore the need for empirically validated measures across all substages of the injured brain.

2. Establish incidence and prevalence of MCS and VS. Data on the course of recovery from coma through VS and MCS to normal consciousness are essential to augment sparse current statistics.

3. Determine the probability that covert consciousness will remain undetected. Signals may go undetected if patients produce neural responses that are too subtle or slow to be detected or if patients do not respond for another reason unrelated to awareness. These clinical-technical challenges underscore the importance of mitigating false-negative results.

4. Harmonize methods, paradigms, and selection of stimuli. Harmonized methods and acquisition paradigms will ultimately yield the greatest power for statistical analysis and comparison to neuropsychological evaluation.

5. Assess the possibility of developing quality-of-life metrics. Assessing quality of life of patients with covert consciousness is a difficult challenge given the limited envelope for communication, but any reproducible strategy will serve to inform the state of these patients and assist health care system decision-making.

TRANSLATION

1. Develop knowledge translation on the meaning and implications of covert consciousness. Specialized knowledge translation tools are essential to capturing and communicating the phenomenon of consciousness after severe brain injury and promoting understanding among physicians, allied health professionals, the media, and the general public.

2. Establish guidelines for timing neuroimaging tests. Timing guidelines should consider many factors, including the nature and severity of the injury, behavioural assessments, and the intended use of the test. They should be aligned with published evidence-based guidelines.

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3. Develop best practices and standards of care for supporting
decision-making. Standards for care should be based on
quantitative and qualitative research about the reaction
of families to choices about testing, their interpretation of
the findings, and their strategies to cope with the results.
Resulting standards should inform how attending physi-
cians, ethicists, and allied health professionals interact with
these families.

4. Develop communication and neuroscientific assessment
protocols effective within technical and patient limitations.
Study designs involving functional magnetic resonance
imaging or other techniques, such as electroencephalography
and magnetoencephalography, involve stimulus-response
paradigms of varying intervals. Questions must be fine-
tuned to allow patients to successfully fulfill the demands of
imaging protocols.

5. Develop protocols appropriate for communicating risks and
decisions. Patients must be able to confirm their under-
standing of the nature and risks of any medical or personal
decisions they are asked to make, and their choice must be
consistent and temporally reliable.

6. Harmonize terminology used to classify covert conscious-
ness. As new DoC nosologies emerge, new terms and scales
for patient management must follow. Harmonization of
classification terminology today will mitigate the diagnostic
confusion and variability seen in the past.

CANADIAN HEALTH CARE AND LEGAL SYSTEMS

1. Integrate respect for the personhood of patients with DoC.
New educational strategies must be developed, tested, and
refined to enable the professional community to embrace
signals of patients’ will and preferences expressed through
neuroimaging techniques.

2. Inventory facilities and staff across Canada. Establish a
trans-Canadian inventory of facilities and staff needed for
assessments of covert consciousness.

3. Clarify the application of capacity laws in the case of DoC.
Unique legal questions raised by covertly conscious patients
should be addressed to ensure that practice complies with
existing legislative and treaty (UNCRPD) obligations.
Article 12 of the UNCRPD commits Canada to take
“appropriate measures” to enable persons with disabilities to
exercise their legal capacity. Although Canada has reserved
the right under the UNCRPD to continue to use substitute
decision-making regimes where appropriate, it has also
indicated that “persons with disabilities are presumed to
have legal capacity.”

4. Establish criteria for evaluating fairness and distributive
justice. A set of common criteria will assist decision-makers to
accommodate regional circumstances and budgets while
retaining publicly defensible accountability-for-reasonableness
frameworks.

Canada has enacted legislation and ratified an international
convention that morally and legally obliges society to respect the
human rights of persons with disabilities. We have identified
priority challenges for research, translation, and policy that must
be resolved to enable patients with DoC to fully exercise their
rights by providing them with neuroimaging technology to
express their will and preferences.

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REFERENCES