

WHEN YOUR PATIENT IS LIVING WITH BRAIN INJURY

A tip card for medical providers treating individuals living with chronic brain injury sequelae

Key points about brain injury (BI):

- BI can affect every aspect of an individual's functioning, leaving some with lifelong challenges.
- BI can be traumatic (TBI) or non-traumatic.
- Injury severity (mild, moderate, severe) does not necessarily predict long-term outcome.
- Many sequelae are difficult to see and therefore may be easy to misinterpret (e.g. lack of initiation, cognitive overload, difficulty recognizing social cues).
- Each injury is unique, like a thumbprint.
- Improvements can occur after initial recovery; re-engagement in therapeutic activities may be beneficial even years post-injury.

Common Sequelae and Subsequent Life Challenges

Areas of Functioning	Specific Sequelae		Subsequent Life Challenges
Motor	Motor planning; coordination; balance; spasticity		Driving/ transportation
Sensory	Changes in vision, hearing, taste, smell or tactile sensation; visual field loss; unilateral neglect; temperature regulation		Following health/wellness recommendations
Cognitive	Attention; concentration; organization; new learning; initiation; memory; problem-solving; judgement; self-awareness; cognitive overload		Communicating needs Relationships, sexuality
Communication	Expressive and receptive communication; dysarthria; tangential speech; following social rules; understanding social cues		Making friends Employment Return to school
Emotional	Regulating emotions; flat affect; easily overstimulated/overwhelmed; increased risk for depression, anxiety and suicidal ideation		Having a sense of meaning in life Behavioral health
Fatigue and Sleep	Physical and emotional fatigue; sleep patterns		Substance use/ misuse



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Interacting with Patients Living with Brain Injury

- Encourage the patient to bring a written list of questions and concerns to the appointment.
- Plan extra time for the appointment to allow for cognitive or communication challenges.
- Encourage the patient to bring a friend/family member to the appointment as a historian/note-taker if needed.
- Encourage compensatory strategies, including
 - o Writing notes in a smartphone or notebook/day-planner organizer;
 - o Using a med-minder; setting alarms on smartphone.
- Find ways to repeat information during the appointment; summarize at the end.
- Have the patient repeat instructions back to you repeat, rehearse, review.
- Provide reminders by email.
- Provide a written summary of the appointment; email a copy of the summary.
- If the patient becomes overwhelmed, model calmness (sit back, take a breath, relax); slow down the information flow; ask how he/she is doing and if they have questions; switch to a lighter topic.
- Encourage an organized approach to wellness (a handout on wellness after BI can be found at https://www.archives-pmr.org/article/S0003-9993(18)30177-1/pdf).
- Encourage socialization and productive activity (support groups, community classes, volunteering).
- Provide resources for support, education and advocacy.

Community Resources - Support, Education, Advocacy

- Brain Injury Association of America <u>www.biausa.org</u>
- United States Brain Injury Alliance <u>www.usbia.org</u> Most states have either a state brain injury association or alliance, offering support groups, resources, education and advocacy. Links to these websites can be found at the two resources above.
- Model Systems Knowledge Translation Center for TBI <u>https://msktc.org/tbi</u>
- Center for Disease Control <u>https://www.cdc.gov/traumaticbraininjury</u>
- Brainline <u>https://www.brainline.org/</u>
- American Stroke Association <u>www.stroke.org</u>
- National Association of State Head Injury Administrators <u>www.nashia.org</u>

This tip card was prepared with support from the Brain Injury Interdisciplinary Special Interest Group (BI-ISIG) of the American Congress of Rehabilitation Medicine (ACRM), by members of the BI-ISIG Chronic Brain Injury Task Force:

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