## NINDS CDE Notice of Copyright Beck Depression Inventory-II (BDI-II)

Availability:	Please visit this website for more information about the instrument: Beck Inventory and Scales website.
Classification:	Supplemental – Highly Recommended: Epilepsy Supplemental: Amyotrophic Lateral Sclerosis (ALS), Epilepsy, Headache, Multiple Sclerosis (MS), Parkinson's Disease (PD), Sports-Related Concussion (SRC) Subacute (after 72 hours to 3 months) and Persistent/Chronic ( 3 months and greater post concussion), and Traumatic Brain Injury (TBI)
	<b>Exploratory:</b> Unruptured Cerebral Aneurysms and Subarachnoid Hemorrhage (SAH)
Short Description of Instrument:	<b>Construct measured:</b> This scale measures the existence and severity of symptoms of depression.
	Generic vs. disease specific: Generic.
	Means of administration: Self-administered.
	Intended respondent: Self-Report.
	# of items: 21 items.
	<b># of subscales and names of sub-scales:</b> 2 subscales: Affective and Somatic subscales.
	# of items per sub-scale: 8 for affective; 13 for somatic.
Comments/Special instructions:	The Beck Depression Inventory-II (BDI-II) developed in 1996, was derived from the BDI. The 21-item self-administered survey is scored on a scale of 0–3 in a list of four statements arranged in increasing severity about a particular symptom of depression.
Scoring:	<b>Scoring:</b> Each of the 21 items corresponding to a symptom of depression is summed to give a single score for the Beck Depression Inventory-II (BDI-II). There is a four-point scale for each item ranging from 0 to 3. On two items (16 and 18) there are seven options to indicate either an increase or decrease of appetite and sleep. Cut-off score guidelines for the BDI-II are given with the recommendation that thresholds be adjusted based on the characteristics of the sample, and the purpose for use of the BDI-II. Total score of 0–13 is considered minimal range, 14–19 is mild, 20–28 is moderate, and 29–63 is severe.

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Rationale / Justification:	<b>Strengths:</b> Easy to use, widely known, results easy to interpret. Item content improved over BDI-I to increase its correspondence with DSM-IV.
	<b>Weaknesses:</b> Includes several items assessing physical symptoms which may be elevated in ALS patients due to motor neuron degeneration and not depression. However non-ALS clinical studies have provided evidence of the presence of at least two factors, a cognitive-affective factor and a somatic depressive symptom factor, which is more stable than in the BDI. However, this factor structure requires confirmation in ALS.
	Psychometric Properties:
	<i>Feasibility:</i> Easy to complete, relatively short compared to interview- based assessments.
	<i>Reliability:</i> 1 week test-retest stability is high (.93). Internal consistency (coefficient alpha) is .92–.94 depending on the sample.
	Validity: Construct validity was high when compared to the BDI (.93).
	<ul> <li>Sensitivity to Change: Designed to assess mood within the most recent</li> <li>2 week period, so comparison across assessments should reflect</li> <li>change over time.</li> </ul>
	<b>Relationships to other variables:</b> BDI-II scores were not correlated with functional disability (ALSFRS-R scores) (Rabkin et al., 2005) in late-stage ALS patients, but did correlate with suffering, anger, perceived caregiver burden, weariness, and negative effect. In non-ALS studies, BDI-II scores correlate with measures of hopelessness, suicidal ideation and anxiety.
	Purpose of Tool: Screening for severity of depression.
	Used in: Observational studies.
	Administration time: 5 minutes, scoring 1 minute.
	Sport Concussion Specific:
	Advantages: Widely used and accepted instrument. Quantifies depressive symptoms, but is not a diagnostic instrument. Some symptoms overlap with "concussive symptoms". Any study looking at factors contributing to persistent symptoms should use this measure.
	Age Range: age 13 and older

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	ALS References:
	Taylor L, Wicks P, Leigh PN, Goldstein LH. Prevalence of depression in amyotrophic lateral sclerosis and other motor disorders. Eur J Neurol. 2010;17(8):1047–1053.
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	Trail M, Nelson ND, Van JN, Appel SH, Lai EC. A study comparing patients with amyotrophic lateral sclerosis and their caregivers on measures of quality of life, depression, and their attitudes toward treatment options. J Neurol Sci. 2003;209(1-2):79–85.