Mental Health Assessments in ICU and Acute Care Megan Altom MOT/S Taylor Harrell MOT/S Carolynn Keane MOT/S Erica Smith MOT/S Danielle Woodward MOT/S March 2020

Evidence-Based Practice Question

In adults with burn injuries in ICU and acute care settings, which mental health assessments are effective for assessing quality of life during initial evaluation?

Clinical Scenario

Sandy Fletchall, an Occupational therapist at the Firefighters Burn Center-Regional One Health, is looking for assessments in the ICU and acute care settings that are focused on emotional states and quality of life, since current assessments are cognitive-based (i.e. SLUMS, MoCA, and the MMSE). Assessments need to be short, easy to administer assessments that are not expensive or are freely available. Evidence-based assessments are important for accurate identification of emotional and trauma related mental health impairments that can be used to gather baseline data for patients in the ICU and acute care settings.

Databases and Sources Searched -Google Scholar, Cochrane, PubMed, MEDLINE, OVID, CINAHL, Veterans Affairs website (www.va.gov)Search Terms -Mental health assessments, Adult with burn injuries Quality of life, Psychometrics and mental health assessments, Trauma, PTSD	Limits Used -English language, Full-text articles only, Age (18+)
---	---

PICO Question Categories

Search Terms Used

Population: Adults with burn injuries in ICU and acute care settings

Outcomes: Quality of life, baseline mental health assessment

Inclusion Criteria for Articles

- Full text articles
- English language
- Adults or older adults
- Assessment- short, easy-to-administer

Exclusion Criteria for Articles

- Pediatric clients
- Adolescents
- Studies occurring more than 20 years ago

Review Process

Our facility needs valid and quick assessments that they can use upon admission to the burn unit that can help guide in interventions. Abstracts and full-text articles were scanned to determine if they were applicable to our PICO question, useful in the ICU, and relevant to the facilities needs. Assessments did not need to be specifically for burns but more so focused on mental health in the ICU. Data extraction was performed by each member of the team to analyze the articles collected during the initial data search. Mentor and professor review of analyses. Articles were appraised for quality using particular forms from Law & MacDermid, 2014).

Search Results by Level of Evidence				
Level of Evidence	Study Design	# of Articles Included		
1	Systematic Review	2		
3	Longitudinal study	1		
4	Comparative Analysis (1), Correlational Analysis (2), Methodological (2)	5		



Modified PRISMA 2009 Flow Diagram (awm 2018)



Article	Level of Evidence	Quality Score	Name of Assessment	Purpose of Assessment	Reliability	Validity	Limitations
 Wei, L. A., Fearing, M. A., Sternberg, E. J., & Inouye, S. K. (2008). The Confusion Assessment Method: A Systematic Review of Current Usage. <i>Journal of</i> <i>the American Geriatrics Society</i>, 56(5), 823–830. Doi:10.1111/j.1532-5415.2008.01674.x 	1	88.9%	The Confusion Assessment Method	Assesses presence, severity, and fluctuation of 9 delirium features: acute onset, inattention, disorganized thinking, altered level of consciousness, disorientation, memory impairment, perceptual disturbances, psychomotor agitation or retardation, and altered sleep-wake cycle	-7 studies- sensitivity rate of 94% (95% Cl 91-97%) -combined sensitivity rate of 89% (95% Cl 85-94%)	-N/A	-Future work needed to validate summary scores or adaptations to measure delirium severity
Wang, Y. P., & Gorenstein, C. (2013). Psychometric properties of the Beck Depression Inventory-II: a comprehensive review. <i>Brazilian Journal of</i> <i>Psychiatry</i> , <i>35</i> (4), 416-431.	1	92%	Beck Depression Inventory-II	BDI-II is a self-report measure of depression in a variety of settings and populations.	- internal consistency around 0.9. -retest reliability ranged from 0.73 to 0.96	-criterion-based validity showed good sensitivity and specificity for detecting depression in comparison to the adopted gold standard. -factor analysis showed a robust dimension of general depression composed by two constructs: cognitive-affective and somatic-vegetative.	-Spectrum bias -Self-report
Augustin, M., Conde Montero, E., Zander, N., Baade, K., Herberger, K., Debus, E. S.,Blome, C. (2017, September). Validity and feasibility of the wound-QoL questionnaire on health-related quality of life in chronic wounds.	Ш	62.5%	Wound-QoL	Wound-QoL (Quality of Life) is a patient-reported assessment used to determine the quality of life of a patient who is experiencing a chronic wound.	N/A	-high convergent validity, -high construct validity, -high longitudinal validity,	-patient-reported
 Amoyal, N. R., Mason, S. T., Gould, N. F., Corry, N., Mahfouz, S., Barkey, A., & Fauerbach, J. A. (2011). Measuring coping behavior in patients with major burn injuries: a psychometric evaluation of the BCOPE. <i>Journal of Burn Care & Research</i>, <i>32</i>(3), 392-398. 	IV	87.5%	BCOPE	Evaluate the BCOPE in measuring coping behaviors, to establish factors underlying coping behaviors in the trauma population and evaluate construct validity.	-Item scale correlation (r = .23 to .76) -Test-retest reliability (baseline to 6 months) ranged from r = .352 -Self-distraction to r = .855 for humor.	-Exploratory factor analysis yielded two factors: active coping and avoidance coping.	-potential differences in coping strategies used in a controlled environment vs. natural setting.
Mouthaan, J., Sijbrandij, M., Reitsma, J. B., Gersons, B. P. R., & Olff, M. (2014). Comparing Screening Instruments to Predict Posttraumatic Stress Disorder. <i>PLoS ONE</i> , <i>9</i> (5). Doi: 10.1371/journal.pone.0097183	IV	91%	SPAN, TSQ, IES-R	SPAN - Assesses frequency and severity of PTSD symptoms over the past week TSQ - Assesses the presence of 5 intrusion items and 5 hyperarousal items over the past week ISE-R - Measures intrusion, avoidance, and hyperarousal	-SPAN - Specificity of 0.64 -TSQ - Specificity of 0.59 -IES-R - Specificity of 0.72 -SPAN (ICC = 0.98) -TSQ (ICC = 0.82) -IES-R (ICC = 0.83)	N/A	-Screening instruments were administered at around 3 weeks following injury, which limited generalizability towards the immediate aftermath of traumatic events -Researchers were unable to collect a clinical PTSD diagnosis of 211 patients at 6 months, whose results could not be included in the index tests.

Aaron, L. A., Patterson, D. R., Finch, C. P., Carrougher, G. J., & Heimbach, D. M. (2001). The utility of a burn specific measure of pain anxiety to prospectively predict pain and function: a comparative analysis. <i>Burns</i> , <i>27</i> (4), 329-334.	IV	68%	BSPAS	Designed to assess burn-related anxiety	-Internal (a -coefficient = 0.90)	-High predictive validity in predicting decreased physical role functioning at time of discharge	-need a larger sample of burn patients and the long-term consequences of reduced physical functioning at discharge should be evaluated both in terms of future function, and psychological adjustment following discharge from hospital
Guenther, U., Popp, J., Koecher, L., Muders, T., Wrigge, H., Ely, E. W., & Putensen, C. (2010). Validity and reliability of the CAM-ICU Flowsheet to diagnose delirium in surgical ICU patients. <i>Journal of critical care</i> , <i>25</i> (1), 144-151.	IV	73%	CAM-ICU Flowsheet	The CAM-ICU Flowsheet is a practical, time-sparing algorithm to assess delirium criteria in intubated patients.	-Very high interrater reliability (κ , 0.96; 0.87-1.00).	-investigators had sensitivities of 88% (95% confidence interval, 69%-98%) and 92% (74%-99%); and specificities of 100% (85%-100%).	-has not been validated as severity scales to distinguish between high and low severity states -included a modest number of intubated patients.
 Wu, CY., Lee, MB., Lin, YY., & Liao, SC. (2019). Development and validation of the 9-item Concise Mental Health Checklist (CMHC-9) for suicide risk assessment. <i>Journal of the Formosan Medical</i> <i>Association</i>, <i>118</i>(9), 1308–1316. doi: 10.1016/j.jfma.2019.05.025 	IV;	68%	CMHC-9	Suicide risk assessment with a three factor structure of demographics, psychopathology and suicidality	-CMHC-9 with AUCs (recent SI 92.9%: 95% CI= 91.5-94.3; lifetime SI 75.9%: 95% CI= 73.9-77.9)	-Crohnbach's <i>a</i> values for total sample: (alpha= 0.79); three sub-samples: (alpha= 0.76 psychiatric; 0.67 community subjects; and 0.69 for medical outpatient subjects)	-use of suicidal ideation rather than other risk factors due to study duration and low prevalence of completed suicide

Bottom Line and Recommendations: There is limited research available specifically for mental health assessments in burn ICU. However, many of the assessments we found, look at aspects that are relevant to the burn population. These assessments can be modified to fit the needs of each client seen in the burn ICU.

REFERENCES

- Aaron, L. A., Patterson, D. R., Finch, C. P., Carrougher, G. J., & Heimbach, D. M. (2001). The utility of a burn specific measure of pain anxiety to prospectively predict pain and function: a comparative analysis. *Burns*, 27(4), 329-334.
- Amoyal, N. R., Mason, S. T., Gould, N. F., Corry, N., Mahfouz, S., Barkey, A., & Fauerbach, J. A. (2011). Measuring coping behavior in patients with major burn injuries: a psychometric evaluation of the BCOPE. *Journal of Burn Care & Research*, 32(3), 392-398.
- Augustin, M., Conde Montero, E., Zander, N., Baade, K., Herberger, K., Debus, E. S., ... Blome, C. (2017, September). Validity and feasibility of the wound-QoL questionnaire on health-related quality of life in chronic wounds. *Wound Repair Regeneration* 25(5):852-857
- Guenther, U., Popp, J., Koecher, L., Muders, T., Wrigge, H., Ely, E. W., & Putensen, C. (2010). Validity and reliability of the CAM-ICU Flowsheet to diagnose delirium in surgical ICU patients. *Journal of Critical Care*, 25(1), 144-151.
- Hissong A.N., Lape, J.E., & Bailey, D.M. (2015). *Bailey's research for the health professional* (3rd ed.), Philadelphia: FA Davis.
- Law, M. & McDermid, J., (Eds.). (2014). Evidence-based rehabilitation: A guide to practice (3rd ed.). Thorofare, NJ: SLACK. Inc.
- Mouthaan, J., Sijbrandij, M., Reitsma, J. B., Gersons, B. P., & Olff, M. (2014). Comparing screening instruments to predict posttraumatic stress disorder. *PloS* one, 9(5), e97183. https://doi.org/10.1371/journal.pone.0097183
- Wang, Y. P., & Gorenstein, C. (2013). Psychometric properties of the Beck Depression Inventory-II: a comprehensive review. *Brazilian Journal of Psychiatry*, 35(4), 416-431.
- Wei, L. A., Fearing, M. A., Sternberg, E. J., & Inouye, S. K. (2008). The Confusion Assessment Method: A Systematic Review of Current Usage. *Journal of the American Geriatrics Society*, 56(5), 823–830. doi: 10.1111/j.1532-5415.2008.01674.x
- Wu, C.-Y., Lee, M.-B., Lin, Y.-Y., & Liao, S.-C. (2019). Development and validation of the 9-item Concise Mental Health Checklist (CMHC-9) for suicide risk assessment. *Journal* of the Formosan Medical Association, 118(9), 1308–1316. doi: 10.1016/j.jfma.2019.05.025

Date Completed:
3/4/2020

Critically Appraised Papers Team #8 March 2020

Critically Appraised Paper #1 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Aaron, L. A., Patterson, D. R., Finch, C. P., Carrougher, G. J., & Heimbach, D. M. Year: 2001 Rater: Danielle Woodward, MOT/S

	DATA EXTRACTED
	Population studied
Population	27 consecutively admitted patients, 24 men and 3 women. The
	mean age of the group was 35 years of age. Participants were
	mostly White but included one African-American and one Hispanic.
	All were admitted to a major medical burn trauma center.
Intervention	N/A, since the study looked only at different assessments.
	Reliability
Reliability (relative)	α -coefficient for the abbreviated Burn Specific Pain Anxiety Scale =
	0.90
Reliability (absolute)	N/A
Minimum detectable	N/A
change	
	Content/Structural validity
Internal consistency	N/A
Content validity	Results in the study concluded that the Burn Specific Pain Anxiety
	Scale (BSPAS) and its contents confidently predict how physical
	health will affect daily activities.
Floor-ceiling effects	Floor-ceiling effects were not discussed in this study.
Factorial validity	The purpose of the study was to see if the BSPAS had predictive
	validity, and after conducting the assessments on the patients, the
	researchers confirmed that the assessment was a very good
	predicator of physical health outcomes.
Item response/Rasch	This study did not have items that were ranged in difficulties
analyses	
	Construct/Criterion validity
Known groups	The researchers already knew about the high reliability of the
	BSPAS and wanted to test for its predictive validity.
Convergent	Researchers discussed how the BSPAS compares to the Profile of
	Mood States (POMS) and State-Trait Anxiety Inventory (STAI) in
	assessing/measuring anxiety and tension.
Divergent	The BSPAS, POMS, and STAI measured anxiety, while the SF-36
	was used to measure physical and emotional functioning and the
	visual analogue scale (VAS) was used to measure pain.
Longitudinal validity	The assessments were used to establish a baseline for patients'
	anxiety and pain levels when they are admitted to the burn center.

Concurrent criterion	The researchers discussed how the BSPAS shows incredibly high
	levels of both criterion and concurrent validity, especially when it
	came to being correlated with a patient's procedural pain and burn
	severity.
Predictive criterion	Results from the study showed the BSPAS being a reliable tool to
	predict how physical dysfunction will affect performance in daily
	activities.
	Responsiveness/Clinical change
Responsiveness	The assessments studied were not used to measure change over time
Minimally clinical	CID/MID was not discussed in the study
important difference	
	Usefulness/practicality
Readability	The patients who were given the assessments were asked to give
	consent and were able to understands the assessments'
	words/phrases
Interpretability	Researchers discussed that the scoring of the results shows a high
	correlation between procedural pain and anxiety, while also showing
	a connection between physical dysfunction and daily activity
	performance.
Time to administer	Less than 10 minutes
Administration burden	Researchers found the results to be significant, but recommend the
	study be replicated with a larger sample size to gain more
	differences in assessments.
Cultural applicability	Suffering from burns does not discriminate, and thus, the
	applicability of this study to many different cultures/backgrounds is
	very relevant, as burns can happen to anyone, and returning to
	normalcy can be very difficult for anyone.

Rationale: The reason I feel this article works for our PICO is because the assessment specifically targets burn patients and how their anxiety could potentially affect their function to perform daily activities in the future.

Critically Appraised Paper #1 Quality Appraisal for Clinical Measurement Research Reports Evaluation Form

Authors: Aaron, L. A., Patterson, D. R., Finch, C. P., Carrougher, G. J., & Heimbach, D. M. Year: 2001

Rater: Danielle Woodward, MOT/S

Evaluation Criteria			
Study Question	2	1	0
1. Was the relevant background work cited to define what is currently		1	
known about the measurement properties of measures under study,			
and the potential contributions of the current research question to			
informing the knowledge base?			
Study Design			
2. Were appropriate inclusion/exclusion criteria defined?		1	
3. Were specific clinical measurement questions/hypotheses	2		
identified?			
4. Was an appropriate scope of measurement properties considered?		1	
5. Was an appropriate sample size used?	2		
6. Was appropriate retention/follow-up obtained? (For studies	N/A	N/A	N/A
involving retesting; otherwise n/a)			
Measurements			
7. Were specific descriptions provided of the measure under study	2		
and the method (s) used to administer it?			
8. Were standardized procedures used to administer all study		1	
measures?			
Analyses			
9. Were analyses conducted for each specific hypothesis or purpose?	2		
10. Were appropriate statistical tests used to obtain point estimates of		1	
the measurement of properties?			
11. Were appropriate ancillary analyses done to quantify in the			0
estimates of the clinical measurement property or the confidence in			
the point estimate (confidence intervals, benchmark			
comparisons/ROC curves, alternate forms of analysis like			
SEM/MID, etc.)?			
Recommendations			
12. Were clear, specific, and accurate conclusions made about the	2		
clinical measurement properties and supported by the study			
objectives, analysis, and results?			
Subtotals (of columns 1 and 2)	10	5	
Total score (sum of subtotals divided by 24 x 100).	68%		
If for a specific paper or topic an item is deemed inappropriate then you			
can sum of items / 2 x number of items x 100			

Critically Appraised Paper #2 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Amoyal, N.R., Mason, S.T., Gould, N. F., Corry, N., Mahfouz, S., Barkey, A., & Fauerbach, J.A. Year: 2011 Rater: Taylor Harrell, MOT/S

	DATA EXTRACTED
	Population studied
Population	362 participants admitted to Johns Hopkins Bayview Medical
	Center Burn Unit. Primarily men (75%), white (65%), employed at
	time of injury (72%). The most common types of burn injuries were
	flame (53%) and scald (17%). Participants had an average TBSA
	(total burn surface area) burned of 15% and an average TBSA
	grafted of 7%. Participants met American Burn Association criteria
	for a major burn.
Intervention	Participants completed the BCOPE survey at discharge from an
	acute stay hospital and then 24 months after discharge. Also, the
	Davidson trauma scale, satisfaction with appearance scale, and
	short-form health survey (SF-12). The BCOPE is a 28-item measure.
	The participants are asked to rate their agreeance one a 4-point
	scale.
	Reliability
Reliability (relative)	Results of the aforementioned studies yielded five or more factors
	across solutions, under scoring the need for a better understanding of
	coping behaviors in specific populations
Reliability (absolute)	DTS: test refeat reliability with a coefficient of $0.86 (P < .01)$
	Acceptance was positively associated with less affective distress at
	discharge, pre-burn, and 12 months after discharge.
	Lest-retest reliability (from baseline to 6 months) ranged from r.352
Minimum data atalula	Tor self-distraction to r.855 for numor (Table 1).
	N/A
change	Contont/Structural validity
Internal consistency	a coefficients for the coeles ranging from 0.50 to 0.0
Internal consistency	DTS: high internal consistency $(a = 0.90)$ for the frequency and
	D 15. fight internal consistency ($a = 0.99$) for the frequency and severity items
	SWAP: Good internal consistency has been reported for the SWAP
	internal consistency (Cronbach's $a = 0.87$)
Content validity	N/A
Floor-ceiling effects	N/A
Factorial validity	Exploratory factorial analysis was conducted using maximum
	likelihood extraction method.

	Solutions were evaluated using Kaiser's criterion.24 scree plot
	analysis, and cohesion of the factors on conceptual grounds.
	Exploratory factor analysis yielded a seven-factor solution that
	accounted for 51% of the total variance
	Correlational analyses revealed that interitem correlations ranged
	from 0.01 to 0.76
Itom mananaa/Daaah	
nem response/ Rasch	N/A
analyses	Construct/Criterian validity
Known groups	
Convergent	Spearman correlations were used to determine the relationship
	between the BCOPE and the SF-12, DTS, and SWAP.
	Results indicated that the active and avoidance factors were each
	positively associated with total DTS scores at 6 and 12 months after
	discharge.
	Heightened post-traumatic stress disorder and distress and highly
	linked to ambivalent coping.
Divergent	Spearman correlations were used to determine the relationship
_	between the BCOPE and the SF-12, DTS, and SWAP.
	Acceptance was negatively associated with SWAP at discharge.
	Using either approach or avoidant coping—but not both—has been
	shown to be related to less distressing outcome when compared to
	ambivalent coping
Longitudinal validity	N/A
Concurrent criterion	N/A
Predictive criterion	Many of the assessments were done up to 12 months after discharge
	Results indicated that the active and avoidance factors were each
	positively associated with total DTS scores at 6 and 12 months after
	discharge Regarding subscale scores, active coning was positively
	associated with DTS intrucion at 1 and 6 months: DTS
	associated with DTS influsion at 1 and 0 months, DTS
	avoidanceati,o, and 12 months, and hyperarousal at 1, 0, and 12
	months. Avoidance coping was significantly correlated with DTS
	intrusion at 1 and 12 months and DTS avoidance at all time points
	Responsiveness/Clinical change
Responsiveness	Measured from time of discharge in acute care (baseline) to 6
	months.
Minimally clinical	N/A
important difference	
	Usefulness/practicality
Readability	The study was designed to assess the BCOPE, Davidson trauma
	scale, satisfaction with appearance scale, and short-form health
	survey (SF-12). The study looked at the different items on each test
	and compared if they would be relevant for the acute care settings to
	use with burn victims.

Interpretability	The results were interpreted, and the study found certain parts of
	each test proved to distinguish different factors.
	Item scale correlations ranged from $r = .23$ to .76
Time to administer	The study talks about the need for a quick, valid and reliable
	assessment to use on burn patients in the acute care settings to aid in
	intervention and coping strategies.
Administration burden	N/A
Cultural applicability	N/A

Critically Appraised Paper #2 Quality Appraisal for Clinical Measurement Research Reports Evaluation Form

Authors: Amoyal, N.R., Mason, S.T., Gould, N. F., Corry, N., Mahfouz, S., Barkey, A., & Fauerbach, J.A. Year: 2011 Rater: Taylor Harrell, MOT/S

Evaluation Criteria		Score		
Study Question			1	0
1. Was the relevant background work cited to define what is currently	7	2		
known about the measurement properties of measures under study,				
and the potential contributions of the current research question to				
informing the knowledge base?				
Study Design				
2. Were appropriate inclusion/exclusion criteria defined?		2		
3. Were specific clinical measurement questions/hypotheses identified	d?	2		
4. Was an appropriate scope of measurement properties considered?		2		
5. Was an appropriate sample size used?		2		
6. Was appropriate retention/follow-up obtained? (For studies involvi	ng	2		
retesting; otherwise n/a)	_			
Measurements				
7. Were specific descriptions provided of the measure under study and	d		1	
the method (s) used to administer it?				
8. Were standardized procedures used to administer all study measure	es?		1	
Analyses				
9. Were analyses conducted for each specific hypothesis or purpose?		2		
10. Were appropriate statistical tests used to obtain point estimates of t	he	2		
measurement of properties?				
11. Were appropriate ancillary analyses done to quantify in the estimat	es		1	
of the clinical measurement property or the confidence in the point				
estimate (confidence intervals, benchmark comparisons/ROC curve	es,			
alternate forms of analysis like SEM/MID, etc.)?				
Recommendations				
12. Were clear, specific, and accurate conclusions made about the		2		
clinical measurement properties and supported by the study				
objectives, analysis, and results?				
Subtotals (of columns 1 and 2)		18	3	
Total score (sum of subtotals divided by 24 x 100).	T	87.5%		
If for a specific paper or topic an item is deemed inappropriate, then you c	an			
sum of items / 2 x number of items x 100				

Critically Appraised Paper #3 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Augustin, M., Conde Montero, E., Zander, N., Baade, K., Herberger, K., Debus, E. S., Blome, C. Year: 2017 Rater: Megan Altom, MOT/S

	DATA EXTRACTED
	Population studied
Population	227 Chronic Wound patients
Intervention	N/A
	Reliability
Reliability (relative)	This assessment was administered as a baseline and then two more
	times throughout an 8-week span.
Reliability (absolute)	N/A
Minimum detectable	N/A
change	
	Content/Structural validity
Internal consistency	Internal consistency was high in 3 time points; Cronbach's
	alpha=0.928, Cronbach's alpha=0.937, Cronbach's alpha=0.947
Content validity	Adult patients (age 18 years) with an existing chronic wound by
	definition were included at any chosen time point within the
	therapeutic program. Patients with a lack of mental, physical, or
	linguistic ability were excluded. Written consent was provided, and
	written data protection policy was available.
Floor-ceiling effects	A low floor effect was observed in T1: 0.5%, T2: 1%, and T3: 4%.
	Ceiling effects were also low (0%, 1%, 0.5%, respectively)
Factorial validity	As a statistically significant change in a PRO (patient-reported
	outcomes) score does not necessarily represent a clinically important
	improvement, and as it can be difficult to know if a PRO score is
	acceptable from the patient's point of view.
Item response/Rasch	The wound-QoL instrument is a patient reported assessment. It gives
analyses	them the chance to answer the questions on a range from none at all,
	a little, moderately, quite a bit, and very much. Item selectivity
	ranges were very similar over time. In 11, item selectivity ranged
	Irom 0.358 to 0.834; 12: 0.441 to 0.821, and 13: 0.538 to 0.828.
V	Construct/Criterion Validity
Known groups	N/A
Convergent	Correlation between FLQA- wk and wound-QoL was high for both
	global score and subscales. This was observed in all 5 time points (bigher then $0.8 < 0.001$)
Divergent	1100000000000000000000000000000000000
Longitudinal validity	Wound Ool was found satisfactory in langitudinal validity since
	there were significantly $(n = 0.001)$ and correlations with the EQ
	= under were significantly (p = 0.001) good contentions with the EQ-

	5D (range 5 0.5–0.7) and FLQA-wk global score (r>0.8) at every
	time point throughout the 8 weeks.
Concurrent criterion	The correlation of the change in Wound-QoL subscales with the change in FLQA-wk subscales was also highly significant, ranging from r 5 0.514 (psyche) to r 5 0.661 (body) for T1-T2 and from r 5 0.577 (psyche) to r 5 0.698 (body) for T1-T3
	These correlations show concurrent validity with the Wound-QoL and the FLQA-wk assessment.
Predictive criterion	N/A
	Responsiveness/Clinical change
Responsiveness	High correlation was found for the change in Wound-QoL global score with change in the wound size. The Wound-QoL global score was significantly better among those with a healed wound compared to the others (0.5 vs. 1.6; $p = 0.002$).
Minimally clinical	The newly developed Wound-QoL was found to be valid and
important difference	responsive and is useful as a short instrument for assessing health- related quality of life.
	Usefulness/practicality
Readability	N/A
Interpretability	N/A
Time to administer	Brief questionnaire, 17 questions. Time depends upon client answering and comprehending the questions
Administration burden	N/A; Article didn't specifically address administrator burden; however, it is a short, patient-reported assessment.
Cultural applicability	As a limitation, the current data were derived from specialized centers with a high expertise in wound care. There may thus be a selection of hard-to-heal wounds. Nevertheless, the study included patients from hospitals as well as from office-based physicians and a large variety of indications, thus providing a certain level of variability.

Critically Appraised Paper #3 Quality Appraisal for Clinical Measurement Research Reports Evaluation Form

Authors: Augustin, M., Conde Montero, E., Zander, N., Baade, K., Herberger, K., Debus, E. S., Blome, C. Year: 2017

Rater: Megan Altom, MOT/S

Evaluation Criteria		ore	
Study Question		1	0
13. Was the relevant background work cited to define what is currently	2		
known about the measurement properties of measures under study,			1
and the potential contributions of the current research question to			1
informing the knowledge base?			1
Study Design			
14. Were appropriate inclusion/exclusion criteria defined?		1	
15. Were specific clinical measurement questions/hypotheses identified?	2		
16. Was an appropriate scope of measurement properties considered?			0
17. Was an appropriate sample size used?		1	
18. Was appropriate retention/follow-up obtained? (For studies involving			0
retesting; otherwise n/a)			1
Measurements			
19. Were specific descriptions provided of the measure under study and	2		
the method (s) used to administer it?			
20. Were standardized procedures used to administer all study measures?		1	
Analyses			
21. Were analyses conducted for each specific hypothesis or purpose?	2		
22. Were appropriate statistical tests used to obtain point estimates of the			
measurement of properties?			
23. Were appropriate ancillary analyses done to quantify in the estimates			0
of the clinical measurement property or the confidence in the point			1
estimate (confidence intervals, benchmark comparisons/ROC curves,			1
alternate forms of analysis like SEM/MID, etc.)?			
Recommendations			
24. Were clear, specific, and accurate conclusions made about the	2		
clinical measurement properties and supported by the study			
objectives, analysis, and results?			
Subtotals (of columns 1 and 2)		3	
Total score (sum of subtotals divided by 24 x 100).			
If for a specific paper or topic an item is deemed inappropriate then you can			
sum of items / 2 x number of items x 100			1

Critically Appraised Paper #4 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Guenther, U., Popp, J., Koecher, L., Muder, T., Wrigger, H., Ely, E. W., & Putensen, C. Year: 2010 Rater: Erica Smith, MOT/S

	DATA EXTRACTED	
Population studied		
Population	Surgical ICU patients.	
	Minimum sample size of 41 patients.	
	Patients' enrollment. One hundred two patients were screened, 48	
	non-German speaking) with another 23 immediate postoperative	
	non-Oerman speaking) with abouter 25 minieutate postoperative natients still under residual effects of anesthesia. One was unwilling	
	to complete the assessment rendering 54 to enroll who were then	
	evaluated by the reference rater using DSM-IV criteria and the	
	CAM-ICU Flowsheet raters.	
Intervention	N/A	
	Reliability	
Reliability (relative)	Delirium prevalence of 40% derived from preliminary studies in the	
• • • •	ICU; and a 95% confidence interval.	
Reliability (absolute)	The agreement (interrater reliability) between the 2 CAM-ICU	
	Flowsheet investigators was very high (Cohen κ , 0.96; 95%	
	confidence interval, 0.87-1.00).	
Minimum detectable	N/A	
change		
	Content/Structural validity	
Internal consistency	N/A	
Content validity	Delirium can be subdivided by assessing motoric symptoms: (1)	
	hyperactive or "agitated" delirium with positive symptoms; (2)	
	hypoactive or "quiet" delirium with negative symptoms; and (3)	
	mixed type, if both subtypes appear alternately over time.	
Floor-ceiling effects	N/A N/A	
Factorial validity	N/A	
Item response/Rasch	N/A	
analyses		
17	Construct/Criterion validity	
Known groups	Analyzed with the Mann-Whitney U test, except for the rate of	
	The characteristic ventiliation and sex analysis, which were done with the Eisher event test. Tests were 2 sided and a $D \leq 0.5$ was considered.	
	risher exact test. Tests were 2-sided, and a $P \le .05$ was considered	
	statistically significant.	

Convergent	N/A
Divergent	N/A
Longitudinal validity	N/A
Concurrent criterion	N/A
Predictive criterion	The CAM-ICU Flowsheet has high sensitivity and high specificity.
	The CAM-ICU Flowsheet investigators had sensitivities of 88% (95% confidence interval, 69%-98%) and 92% (74%-99%), specificities of 100% (85%-100%).
	Screened for delirium (1) by a psychiatrist as the reference rater using the 4 delirium criteria of the Diagnostic and Statistical Manual of Mental Diseases, Fourth Edition (DSM-IV).
	The "CAM-ICU Flowsheet," derived from the Confusion Assessment Method for Intensive Care Units (CAM-ICU), provides an algorithm by which to assess the 4 delirium criteria of the DSM- IV in a standardized fashion in intubated patients.
	Sensitivity (true positives [TP]/true positives [TP] + false negatives [FN]), specificity (true negatives [TN]/true negatives [TN] + false positives [FP]), positive predictive values (TP/TP + FP), negative predictive values (TN/TN + FN), and overall accuracies (TP + TN/TP + FP + TN) were calculated from 2×2 tables (Prism4 Software for Macintosh, GraphPad Software Inc. San Diego, Calif).
	Responsiveness/Clinical change
Responsiveness	N/A
Minimally clinical	N/A
important difference	
	Usefulness/practicality
Readability	N/A
Interpretability	Accuracies of 94% to 96%
	It is important to note that the CAM-ICU Flowsheet switches the original numbering of features 3 and 4 for simplicity because most ICU patients with delirium are positive in the order of the flow sheet, thus allowing the CAM-ICU Flowsheet to be completed in just 3 features and only needing to include the fourth feature in a minority of patients.
	Delirium subtypes were classified into a motoric subtype grouping according to the Richmond Agitation Sedation Scale (RASS), which was rated for the feature 3 ("altered level of consciousness") of the CAM- ICU Flowsheet.

	Patients were deemed as having hypoactive delirium if they were DSM positive by the reference rater and had RASS -3 to 0, or deemed as having hyperactive delirium if their RASS was between +1 to +4. Mixed type is defined by alternating between either state, but because we only evaluated each patient once to avoid repeat- observer bias, mixed-type delirium was not diagnosed in this investigation. Patients with a RASS of -4 or -5 were considered
	comatose and were excluded from the study because comatose patients cannot be assessed for delirium.
Time to administer	Quick assessment instrument.
	Median assessment times of less than a minute
	The median time to complete the CAM-ICU Flowsheet assessment was 45 seconds (IQR, 40-75 seconds) in patients without delirium vs 50 seconds (IQR, 40-120 seconds; maximum, 180 seconds) in patients with delirium.
Administration burden	N/A
Cultural applicability	The CAM-ICU Flowsheet is measured similarly across cultures.
	The German CAM-ICU Flowsheet was translated according to the Principles of Good Practice for the Translation and Cultural Adaptation Process for Patient- Reported Outcomes Measures into German language [31,32]. This included the "forward translation" of the original English CAM-ICU Flowsheet, the revision by a geriatric psychiatrist, and the back-translation into English by 2 physicians who were unaware of the original.
	The German version is also available online at www.icudelirium.org.

Critically Appraised Paper #4 **Quality Appraisal for Clinical Measurement Research Reports Evaluation Form**

Authors: Guenther, U., Popp, J., Koecher, L., Muder, T., Wrigger, H., Ely, E. W., & Putensen, C. Year: 2010

Rater: Erica Smith, MOT/S

Evaluation Criteria		Score	
Study Question		1	0
1. Was the relevant background work cited to define what is currently	2		
known about the measurement properties of measures under study, and			
the potential contributions of the current research question to			
informing the knowledge base?			
Study Design			
2. Were appropriate inclusion/exclusion criteria defined?		1	
3. Were specific clinical measurement questions/hypotheses identified?		1	
4. Was an appropriate scope of measurement properties considered?	2		
5. Was an appropriate sample size used?		1	
6. Was appropriate retention/follow-up obtained? (For studies involving			0
retesting; otherwise n/a)			
Measurements			
7. Were specific descriptions provided of the measure under study and	2		
the method (s) used to administer it?			
8. Were standardized procedures used to administer all study measures?	2		
Analyses			
9. Were analyses conducted for each specific hypothesis or purpose?		1	
10. Were appropriate statistical tests used to obtain point estimates of the		1	
measurement of properties?			
11. Were appropriate ancillary analyses done to quantify in the estimates		1	
of the clinical measurement property or the confidence in the point			
estimate (confidence intervals, benchmark comparisons/ROC curves,			
alternate forms of analysis like SEM/MID, etc.)?			
Recommendations			
12. Were clear, specific, and accurate conclusions made about the clinical	2		
measurement properties and supported by the study objectives,			
analysis, and results?			
Subtotals (of columns 1 and 2)	10	6	
Total score (sum of subtotals divided by 24 x 100).	73%		
If for a specific paper or topic an item is deemed inappropriate then you can			
sum of items / 2 x number of items x 100			

Critically Appraised Paper #5 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Mouthaan, J., Sijbrandij, M., Reitsma, J. B., Gersons, B. P. R., & Olff, M. Year: 2014 Rater: Danielle Woodward MOT/S

	DATA EXTRACTED	
Population studied		
Population 311 participants, with injuries sustained in a traumatic event.		
Intervention	No intervention was present in the study.	
	Reliability	
Reliability (relative)	Clinical interview of the Clinician Administered PTSD scale	
	(CAPS) - (ICC = 0.98)	
	Startle, Physiological arousal, Anger, and Numbness (SPAN) - (ICC	
	= 0.83)	
	Trauma Screening Questionnaire $(TSQ) - (ICC = 0.82)$	
	Impact of Event Scale-Revised (IES-R) $-$ (ICC = 0.83)	
Reliability (absolute)	No standard error of measurement was reported in the study	
Minimum detectable	The MDC was not identified in this; the one thing specified was if a	
change	patient would be diagnosed with PTSD, their score would have to	
	have been at least 45 or higher.	
	Content/Structural validity	
Internal consistency	High internal consistencies were shown for the CAPS subscales	
	(Intrusion: $\alpha = 0.91$, Avoidance: $\alpha = 0.83$, and Hyperarousal: $\alpha = 0.91$	
~ !! !!		
Content validity	All the assessments used (CAPS, SPAN, TSQ, and IES-R) measured	
	what they were made to measure, helping predict future mental	
F1 '1' CC (health issues with patients suffering from trauma-related injuries.	
Floor-ceiling effects	Floor-ceiling effects were not measured in this study.	
Factorial validity	Researchers discussed that the assessments used were excellent tools	
	for detecting future PTSD and other mental filnesses, though further	
	testing must be done in order to accurately diagnosis the linesses in	
Itom magnance/Decel	Items on the accordinate wave not reprod in difficulty	
analyzag	tients on the assessments were not ranged in difficulty.	
anaryses Construct/Criterier velidity		
Known groups	Known groups were not discussed in the study	
Convergent	All assessments used were able to adequately detect PTSD in	
Convergent	nations with trauma-related injury, whether it was a short	
	assessment or long one	
Divergent	Researchers only discussed that further testing/assessing must be	
	done in order to accurately diagnosis mental illness in patients	
Longitudinal validity	After 6 months and being reassessed researchers found that 5.8% of	
	the patients studied were diagnosed with 6-month PTSD. 7.1% were	

	diagnosed with Major Depressive Episode (MDE), and 7.8% with anxiety disorder (AD).
Concurrent criterion	SPAN and CAPS (Pearson $r = 0.65$, $p < 0.001$)
	TSQ and CAPS (Pearson $r = 0.72$, $p < 0.001$).
Predictive criterion	Researchers discussed that the assessments used were good
	predictors at diagnosing PTSD and other mental illnesses in patients
	suffering from trauma-related injuries.
	Responsiveness/Clinical change
Responsiveness	The assessments can be used to diagnose patients with mental illness
	and can be used as a baseline to create and plan interventions that
	might help with those later diagnoses.
Minimally clinical	MDC was not addressed in this study.
important difference	
Usefulness/practicality	
Readability	The patients were able to understand the assessments, and results
	were able to be understood and analyzed appropriately.
Interpretability	Researchers were able to interpret and find moderate to strong
	correlations between the assessments on predicting mental illnesses
	such as PTSD.
Time to administer	Varied, as the SPAN and TSQ were short, and the IES-R and CAPS
	were long.
Administration burden	The CAPS was done by those who were trained to administer it, as
	well as all the other assessments used in the study.
Cultural applicability	Though the study took place in the Netherlands, the assessments
	used are widely known and used often when it comes to trauma-
	related injuries in the ICU.

Rationale: This article, despite looking at mostly other patients besides burn patients, the assessments compared were great for predicting mental health complications for those who are admitted to the ICU, for whatever reason.

Critically Appraised Paper #5 Quality Appraisal for Clinical Measurement Research Reports Evaluation Form

Authors: Mouthaan, J., Sijbrandij, M., Reitsma, J. B., Gersons, B. P. R., & Olff, M. Year: 2014

Rater: Danielle Woodward MOT/S

Evaluation Criteria		Score	
Study Question		1	0
1. Was the relevant background work cited to define what is currently	2		
known about the measurement properties of measures under study,			
and the potential contributions of the current research question to			
informing the knowledge base?			
Study Design			
2. Were appropriate inclusion/exclusion criteria defined?		1	
3. Were specific clinical measurement questions/hypotheses identified?	2		
4. Was an appropriate scope of measurement properties considered?	2		
5. Was an appropriate sample size used?	2		
6. Was appropriate retention/follow-up obtained? (For studies	N/A	N/A	N/A
involving retesting; otherwise n/a)			
Measurements			
7. Were specific descriptions provided of the measure under study	2		
and the method (s) used to administer it?			
8. Were standardized procedures used to administer all study		1	
measures?			
Analyses			
9. Were analyses conducted for each specific hypothesis or purpose?	2		
10. Were appropriate statistical tests used to obtain point estimates of	2		
the measurement of properties?			
11. Were appropriate ancillary analyses done to quantify in the	2		
estimates of the clinical measurement property or the confidence in			
the point estimate (confidence intervals, benchmark			
comparisons/ROC curves, alternate forms of analysis like			
SEM/MID, etc.)?			
Recommendations			
12. Were clear, specific, and accurate conclusions made about the	2		
clinical measurement properties and supported by the study			
objectives, analysis, and results?			
Subtotals (of columns 1 and 2)	18	2	
Total score (sum of subtotals divided by 24 x 100).	91%		
If for a specific paper or topic an item is deemed inappropriate then you			
can sum of items / 2 x number of items x 100			

Critically Appraised Paper #6 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Wang, Y.P, & Gorenstein, C. Year: 2013 Rater: Erica Smith, MOT/S

	DATA EXTRACTED
	Population studied
Population	A variety of populations.
	The instrument was applied to over 60,000 respondents.
Intervention	N/A
	Reliability
Reliability (relative)	Twenty-nine of the 118 retrieved articles (25%) did not report reliability coefficients, indicating that the assumption of test score reliability generally has not prevailed in clinical practice regarding application of the BDI.
Reliability (absolute)	Retest reliability (Pearson's r) showed relative stability through re- application of the BDI-II, with good to excellent coefficients (range, 0.73 to 0.96), with a mean re-application interval of 2 weeks (range, 1 week to 6 months) for the majority of studies (82%).
Minimum detectable change	N/A
	Content/Structural validity
Internal consistency	In comparison to the internal consistency of the previous versions of the BDI (average Cronbach's alpha coefficient around 0.85),8 most studies on BDI-II reported an average alpha coefficient around 0.9, ranging from 0.83 to 0.96.
Content validity	 The English version of the BDI-II has been translated into 17 languages, and is used in Europe, the Middle East, Asia, and Latin America (Table 1). Although the English version prevailed among the studies (65%), the increasing number of language versions suggests inter- national acceptance of the instrument. The content validity of the BDI-II appears to be adequate, but narrower than that of the former version. The BDI-I reflected six of the nine criteria for DSM-based depression, while the BDI-II presented an improved performance on specificity to indicate DSM-based depression.
Floor-ceiling effects	N/A
Factorial validity	Beck reported a median item-total scale correlation of 0.59 for the BDI-II in a sample of college students (n=120). Acceptable item- total scale correlations (rit is less than or equal to 0.5) were described for 17 out of 21 items. Nonetheless, this correlation can vary across studies. For the Arabic version, substantial item-total

	correlation was described for 10 items among Islamic students,
	whereas adequate item-total correlation of the Portuguese version in
	Brazilian samples was reported for 15 items. Factors such as
	language version, type of sample, age range, educational level, and
	severity of depression might affect the difficulty of item
	endorsement. Insight into which items should be assigned to a scale
	can improve its performance through item-level analysis. Factor
	analysis showed a robust dimension of general depression composed
	by two constructs: cognitive-affective and somatic-vegetative.
Item response/Rasch	Most validation studies of the BDI-II were analyzed on the grounds
analyses	of classic test theory (CTT), assuming a true score for each
	respondent and disregarding the measurement error. In other words,
	two individuals with the same total score may differ in terms of the
	relative severity and frequency of symptoms.
	Construct/Criterion validity
Known groups	The confirmatory strategy has been employed to compare the
groups	structure and model fit of previous studies in relation to the
	construct validity of the BDI-II.
Convergent	The convergent validity between the BDI-I and the BDI-II was high.
	with Pearson's product-moment correlation coefficients (r) ranging
	from 0.82 to 0.94
	The convergent validity between the BDI-II and scales that assess
	anxiety – such as the Beck Anxiety Inventory (BAI), the Hamilton
	Anxiety Rating Scale (HAM-A), and the State-Trait Anxiety
	Inventory (STAI) – was also significant, with a wide range of
	correlation coefficients (0.37 to 0.83: rough estimate of 0.50).
Divergent	N/A
Longitudinal validity	N/A
Concurrent criterion	This study reported a comparison of the BDI-II with scales
	measuring depression anxiety and miscellaneous constructs as
	criterion determined at essentially the same time to check for
	concurrent validity
Predictive criterion	In general studies reported a sensitivity of 0.70 Sensitivity should
	be viewed as the most important indicator to minimize the chance of
	false-negative diagnosis of depressive dis- orders
	Responsiveness/Clinical change
Responsiveness	The mean score ranged from 5.1 to 38.4
Minimally clinical	N/A
important difference	
	Usefulness/practicality
Readability	N/A
Internretability	N/Δ
Time to administer	N/A

Administration burden	Scores were found to significantly decrease for the weekly administration group only, indicating that lower retest scores could be the result of a measurement effect and the frequency of administration
Cultural applicability	With the BDI-II being such a popular measure adapted for use in several countries, information on cross- cultural comparability is still remarkably scarce. The cross-cultural equivalence between the versions of the BDI-II stands out as a topic of fervent academic interest: the symptomatology of depression in different culture/races or languages can be compared by testing the measurement of variance of the instrument.

Critically Appraised Paper #6 Quality Appraisal for Clinical Measurement Research Reports Evaluation Form

Authors: Wang, Y.P, & Gorenstein, C. Year: 2013 Rater: Erica Smith, MOT/S

Evaluation Criteria		Score	
Study Question		1	0
1. Was the relevant background work cited to define what is currently	2		
known about the measurement properties of measures under study, and			
the potential contributions of the current research question to			
informing the knowledge base?			
Study Design			
2. Were appropriate inclusion/exclusion criteria defined?		1	
3. Were specific clinical measurement questions/hypotheses identified?	2		
4. Was an appropriate scope of measurement properties considered?	2		
5. Was an appropriate sample size used?	2		
6. Was appropriate retention/follow-up obtained? (For studies involving	2		
retesting; otherwise n/a)			
Measurements			
7. Were specific descriptions provided of the measure under study and	2		
the method (s) used to administer it?			
8. Were standardized procedures used to administer all study measures?			
Analyses			
9. Were analyses conducted for each specific hypothesis or purpose?	2		
10. Were appropriate statistical tests used to obtain point estimates of the	2		
measurement of properties?			
11. Were appropriate ancillary analyses done to quantify in the estimates		1	
of the clinical measurement property or the confidence in the point			
estimate (confidence intervals, benchmark comparisons/ROC curves,			
alternate forms of analysis like SEM/MID, etc.)?			
Recommendations			
12. Were clear, specific, and accurate conclusions made about the clinical	2		
measurement properties and supported by the study objectives,			
analysis, and results?			
Subtotals (of columns 1 and 2)		2	
Total score (sum of subtotals divided by 24 x 100).	92%		
If for a specific paper or topic an item is deemed inappropriate then you can			
sum of items / 2 x number of items x 100			

Critically Appraised Paper #7 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Wei, L. A., Fearing, M. A., Sternberg, E. J., & Inouye, S. K. Year: 2008 Rater: Carolynn Keane, MOT/S

	DATA EXTRACTED		
	Population studied		
Population	Patients with dementia, depression, psychosis		
Intervention	The use of the Confusion Assessment Method to address recognition		
	and management of delirium.		
	Reliability		
- Interrater reliabili	ity (.70-1.00) was moderate to high across all studies		
Gonzalez 2004:			
 Sensitivity 	y: .90		
 Specificity 	y: 1.0		
 Inter-rater 	reliability: .89		
- Laurila 2002:			
 Sensitivity 	y: .8186		
 Specificity 	y: .6384		
 Inter-rater 	reliability:		
- Fabbri 2001:			
 Sensitivity 	y: .94		
 Specificity 	y: .96		
 Inter-rater 	reliability: .70		
- Monette 2001:			
 Sensitivity 	 Sensitivity: .86 		
 Specificity: 1.0 			
Inter-rater	reliability: .91		
- Ely 2001a:			
Sensitivity	y: .95-1.0		
	y: .8993		
■ Inter-rater	reliability: ./995		
- Ely 20016:			
- Sensitivity	y: .93-1.0		
- Specificity	y: .98-1.0		
Polfcon 1000	Tenaolinty90		
- Kollsoll 1999.	ar. 70		
- Sensitivity	y / 0 w. 1 00		
■ Inter_rater	y. 1.00 : reliability:		
- 701 1008.	Tonuomity		
Sensitivity	v· 89		
 Specificity 	y: 1.00		

 Inter-rates 	r reliability: 0.86				
- Pompei 1995:					
 Sensitivity: .46 					
 Specificity: .92 					
 Inter-rater reliability: 					
- Rockwood 1994:	- Rockwood 1994:				
 Sensitivit 	y: .68				
 Specificit 	y: .97				
 Inter-rate 	r reliability:				
- Inouye 1990:					
 Sensitivit 	y: .94-1.0				
 Specificit 	y: .9095				
 Inter-rates 	r reliability: .81-1.0				
Reliability (relative)	ICC not stated				
Reliability (absolute)	SEM not stated				
Minimum detectable	Results were combined across 7 high quality studies (n=1071),				
change	demonstrating an overall sensitivity of 94% (95% confidence				
	interval, CI, 91-97%), and specificity of 89% (95% CI, 85-94%).				
	Content/Structural validity				
Internal consistency Cronbach's alpha values not stated					
Content validity N/A					
Floor-ceiling effects	N/A				
Factorial validity	N/A				
Item response/Rasch	N/A				
analyses					
	Construct/Criterion validity				
Known groups	N/A				
Convergent	- CAM-ICU				
	- CAM for ED				
	- MDS Version 3.0				
	- Nursing Home CAM				
	- Delirium Index				
	- One day Fluctuation Scale				
Divergent	In two validation studies (<u>11–12</u>), the CAM-ICU was compared				
	with ratings by delirium experts using the DSM-IV, yielding a				
	sensitivity of 95–100%, specificity of 93–98%, and interrater				
	reliability of .79–.95. When the non-verbal ratings on the CAM-ICU				
	were directly compared to verbal ratings (24), CAM-ICU ratings				
	demonstrated lower sensitivity of 73% and lower interrater				
	reliability of .64, but maintained a high specificity of 100%.				
Longitudinal validity	N/A				
Concurrent criterion	N/A				
Predictive criterion	- Positive predictive accuracy of 91-94%				
	- Negative predictive accuracy of 90-100%				
	Responsiveness/Clinical change				

Responsiveness	
Minimally clinical	N/A
important difference	
	Usefulness/practicality
Readability	- Comprehensive literature search for articles published
	between January 1, 1991 and December 31, 2006 using
	PubMED, EMBASE, PsychINFO, CINAHL, Ageline, and
	Google Scholar. Full text searches for the search term
	'Confusion Assessment Method'. Applying additional search
	terms, such as 'delirium', 'acute confusional state'. Excluded
	articles based off of inclusion criteria.
	- In over 222 application studies, we have documented that the
	CAM is widely used for both clinical and research
	applications.
Interpretability	Results compared on sensitivity, specificity and inter-rater
	reliability.
Time to administer	Not stated
Administration burden	Some training is recommended for optimal use. (See Confusion
	Assessment Method Training Manual)
Cultural applicability	CAM has been translated into 10 languages.

Critically Appraised Paper #7

Heal	th	Evic	len	ce™
Helping public h	ealth use h	nest evidence in	practice sin	ce 2005

Quality Assessment Tool – Review Articles

Instructions for completion: Please refer to the attached dictionary for definition of terms and instructions for completing each section. For each criteria, score by placing a check mark in the appropriate box.

First Author:	Leslie A. Wei
Year:	2008
Journal:	N/A
Reviewer:	Carolynn Keane, MOT/S

	CRITERIA	YES	ľ
Q1. Did the authors have a clearly focused question [population, intervention (strategy), and outcome(s)]?Q2. Were appropriate inclusion criteria used to select primary studies?		x	
		x	
Q3. Did the authors describe a s	earch strategy that was comprehensive?		T
Circle all strategies used:	 health databases psychological databases social science databases educational databases other handsearching key informants reference lists unpublished 	x	
Q4. Did search strategy cover a	adequate number of years?	x	
Q5. Did the authors describe the	level of evidence in the primary studies included in the review?		-
 Level I → RCTs only Level II → non-randomized, cohort, case-control Level III → uncontrolled studies 		N/A	
Q6. Did the review assess the m	ethodological quality of the primary studies, including:		-
(Minimum requirement: 4/7 o	f the following)		
 Research design Study sample Participation rates Sources of bias (confored) Data collection (measured) Follow-up/attrition rates Data analysis 	inders, respondent bias) rement of independent/dependent variables)		
Q7. Are the results of the review	transparent?	x	
Q8. Was it appropriate to combin	ne the findings of results across studies?	x	
Q9. Were appropriate methods	used for combining or comparing results across studies?	x	
10 Do the data support the aut	nor's interpretation?	x	

Critically Appraised Paper #8 <u>Data Extraction Form for Studies Evaluating the</u> <u>Clinical Measurement Properties of Outcome Measures</u>

Authors: Chia-Yi Wu, Ming-Been Lee, Yi-Yin Lin, Shih-Cheng Liao Year: 2019 Rater: Carolynn Keane, MOT/S

DATA EXTRACTED				
Population studied				
Population	- 3 groups of subjects			
	 931 psychiatric outpatients 			
	 931 non-psychiatric medical outpatients 			
	 2120 community residents 			
	- Recruited from convenience samples of the outpatient clinics			
	of the psychiatric department and the non-psychiatric			
	departments of family medicine, oncology, and emergency			
	medicine in a medical center in Northern Taiwan.			
Intervention	Participants were invited to answer a battery of structured questions			
	in written form (patient subjects) or by telephone form (community			
	subjects) by personnel with specific training for the study. The			
	structured questionnaire including CMHC items and additional			
	questions for demographic characteristics and suicidality was			
	administered via a standardized procedure in different settings.			
	Reliability			
Reliability (relative)	No ICC given			
Reliability (absolute)	No standard error of measurement given			
Minimum detectable	CMHC-9 with AUCs (recent SI 92.9%: 95% CI = 91.5-94.3;			
change lifetime SI 75.9%: 95% CI= 73.9-77.9)				
Content/Structural validity				
Internal consistency	The internal consistency of the CMHC-9 was satisfactory based on			
	Cronbach's alpha values for the total sample (alpha= 0.79) and the			
	three sub-samples (i.e., alpha= 0.76 for psychiatric subjects; 0.67 for			
	community subjects; and 0.69 for medical outpatient subjects).			
Content validity	To reduce the risk of "lack of assessment" that limits opportunities			
	for mental health care, universal screening is recommended			
Floor-ceiling effects	- It can be part of clinical assessment in detecting patients'			
	recent psychopathology and suicidality, potentially			
	improving patient-centered care and subjective and objective			
	intervention outcomes.			
	- The lowest percentage of scoring level of 0 was found in the			
	psychiatric outpatient group, while over 30-45% of the other			
	two samples also scored 0. This demonstrates the floor effect			
	in the psychiatric group as well as distinct discrimination			
	between lower scoring groups from the higher scoring group.			

Factorial validity	- The factorial structure of CMHC-9 was invariant across
	groups and the summed scores could differentiate a high-risk
	group in the psychiatric sub-sample from other medical or
	community sub-samples.
	- The results indicated that three common first-order factors
	were extracted with an eigenvalue greater than one and
	parallel analysis. In total, 54% of the total variance was
	explained by these three factors:
	the highest % of variance was explained by Factor 1
	(31%), including five items of psychopathology (with
	factor loadings of 0.61-0.91);
	 Factor 2 (31%), containing four items of suicidality
	from the CSPS such as lifetime suicide attempt,
	future suicidal intent, substance abuse and lack of
	support (loadings = $0.95, 0.54, 0.42, \text{ and } 0.34,$
	respectively)
	• Factor 3 (8%), consisting of the two items of age and
	marital status (loadings= 0.68 and 0.65 , respectively).
Item response/Rasch	Post hoc analysis using the Turkey HSD test revealed significant
analyses	differences in paired comparisons among community subjects,
	medical patients, and psychiatric patients (Means $=$ 1.2, 1.9, and 4,
	SDs = 1.6, 1.8, and 2.3, respectively, p < 0.001).
	Construct/Criterion validity
Known groups	- The demographics differed significantly among the three
	sub-samples, with a higher rate of females and those
	divorced/separated/widowed in psychiatric subjects and
	younger age in community subjects.
	- The descriptive statistics of the suicide risk factor
	items/scales differed significantly among the three sub-
	samples, with the highest percentages of all risk factor items
	and overall scale scores of the psychopathology and CMHC-
	9 in psychiatric subjects, and the lowest percentages in
	community subjects.
	- The psychiatric subjects had more severe suicide risk and
	psychological distress than the other two groups.
Convergent	- The CMHC-9 was originated from the CMHC, which
	contained 16 items for suicide risk assessment with a three-
	factor structure of demographics, psychopathology and
	suicidality.
	- CMHC consisted of 5 items of psychopathology from the
	BSRS-5R, nine items from the CSPS and 2 additional
	measures for recent and lifetime suicide ideation
Divergent	There is a lack of correlation among the tests.
Longitudinal validity	N/A

Concurrent criterion	Time-specific assessment of suicide risks including recent
	psychopathological symptoms (within seven days), past suicide
	attempt (over the lifetime) and future suicide attempt).
	BSRS-5R- Satisfactory psychometric properties as a measure
	to detect psychiatric morbidity and SI in medical settings or
	in the community.
	• CPS
Predictive criterion	The ROC curves of the CMHC-9 scores revealed that the optimal
	cut-off to predict recent SI was $\frac{3}{4}$ for all subjects with 92.1%
	sensitivity, 82.0% specificity, 99.2% negative predictive value and
	30.6% positive predictive value.
	Responsiveness/Clinical change
Responsiveness	Yes, it can assess suicide risks including recent psychopathological
1	symptoms (within seven days), past suicide attempt (over the
	lifetime) and future suicide intent.
Minimally clinical	The optimal cut-offs to predict recent SI in all subjects and in
important difference	different demographic subgroups were examined. The significance
	of all tests was set at $p < 0.05$.
	Usefulness/practicality
Readability	The structured questionnaire including CMHC items and additional
	questions for demographic characteristics and suicidality (e.g.,
	serious consideration of suicide over the past week or over the
	lifetime) was administered via a standardized procedure in different
	settings.
Interpretability	The CMHC-9 scores among the three samples were compared with
	ANOVA and found significant between-group differences (F (2,
	3979) = 821.5, p < 0.001).
Time to administer	Not given, only 9-items so fairly short
Administration burden	Can be self-administered or interview format
Cultural applicability	Checklist does not change amongst cultures

Critically Appraised Paper #8 Quality Appraisal for Clinical Measurement Research Reports Evaluation Form

Authors: Chia-Yi Wu, Ming-Been Lee, Yi-Yin Lin, Shih-Cheng Liao Year: 2019 Rater: Carolynn Keane, MOT/S

Evaluation Criteria		Score	
Study Question			0
1. Was the relevant background work cited to define what is currently	2		
known about the measurement properties of measures under study,			
and the potential contributions of the current research question to			
informing the knowledge base?			
Study Design			
2. Were appropriate inclusion/exclusion criteria defined?	2		
3. Were specific clinical measurement questions/hypotheses identified?			0
4. Was an appropriate scope of measurement properties considered?		1	
5. Was an appropriate sample size used?			0
6. Was appropriate retention/follow-up obtained? (For studies involving	; n/a	n/a	n/a
retesting; otherwise n/a)			
Measurements			
7. Were specific descriptions provided of the measure under study and	2		
the method (s) used to administer it?			
8. Were standardized procedures used to administer all study measures?	2		
Analyses			
9. Were analyses conducted for each specific hypothesis or purpose?			
10. Were appropriate statistical tests used to obtain point estimates of the	2		
measurement of properties?			
11. Were appropriate ancillary analyses done to quantify in the estimates			0
of the clinical measurement property or the confidence in the point			
estimate (confidence intervals, benchmark comparisons/ROC curves,			
alternate forms of analysis like SEM/MID, etc.)?			
Recommendations			
12. Were clear, specific, and accurate conclusions made about the clinica	1	1	
measurement properties and supported by the study objectives,			
analysis, and results?			
Subtotals (of columns 1 and 2)		2	
Total score (sum of subtotals divided by 24 x 100).	68%		
If for a specific paper or topic an item is deemed inappropriate then you can			
sum of items / 2 x number of items x 100			