

REVIEW

A review of scales for assessing the risk of developing a pressure ulcer in individuals with SCI

WB Mortenson^{1,2,3}, WC Miller^{2,3,4,5} and the SCIRE Research Team

¹Occupational Therapy, Long-term Care, Vancouver Coastal Health, Vancouver, British Columbia, Canada; ²School of Rehabilitation Sciences, University of British Columbia, Vancouver, British Columbia, Canada; ³GF Strong Rehabilitation Research Lab, Vancouver, British Columbia, Canada; ⁴International Collaboration on Repair Discoveries, Vancouver, British Columbia, Canada and ⁵Department of Occupational Science and Occupational Therapy, University of British Columbia, Vancouver, British Columbia, Canada

Background: Pressure ulcers are a common secondary condition that occur post-spinal cord injury (SCI). These ulcers come at tremendous personal and societal cost. There are a number of scales that can be used to identify those who are at risk.

Objectives: This review critically evaluates risk assessment scales designed for identifying and predicting skin ulcers. Specifically, studies on the psychometric properties and utility for individuals with SCI were assessed.

Methods: The MedLine, CINHAL, Embase, HaPI, Psycinfo, Sportdiscus and Cochrane databases were searched to identify studies. To be included, the scale needed to have at least one study, published in a peer-reviewed journal, which examined its psychometric properties with a sample of individuals with SCI.

Results: Seven scales were included in this review: Abuzzese, Braden, Gosnell, Norton, SCIPUS, SCIPUS-A and Waterlow. None of the tools reported reliability data with this population. Validity evidence ranged from poor to adequate across scales. Most were readily available, quick to administer and had minimal respondent burden; however, the SCIPUS-A and SCIPUS, two scales developed specifically for individuals with SCI, required laboratory blood testing.

Conclusion: Although the SCIPUS-A and SCIPUS show promise, utility issues and limited psychometric testing suggest that these tools cannot be recommended at this time. While the Braden scale has the best combined validity and utility evidence, more specific testing with individuals with SCI is required for it and all other scales included in the review.

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Introduction

Individuals with spinal cord injury (SCI) experience a variety of long-term secondary medical complications. Pressure ulcers, defined as an area of localized damage to the skin and underlying tissue caused by pressure, shear, friction and/or a combination of these (European Pressure Ulcer Advisory Panel, 2007) (Accessed March 7 from <http://www.epuap.org/gltreatment.html>), are one of the most serious and frequent of these problems.^{1,2} These chronic wounds can have harmful personal^{3,4} and societal effects.^{5,6}

Pressure ulcers are extremely common in individuals with SCI. It is estimated 85% of individuals with SCI will

experience a pressure ulcer during their lifetime.⁷ McKinley *et al.*² found that pressure ulcers were the most common secondary medical complication of SCI with prevalence rates ranging from 15.2% 1 year after injury to 29.4% at 20 years after injury. Chen⁸ reported a trend toward increasing pressure-ulcer prevalence in recent years, which is not explained by aging, years since injury or demographic variables.

The medical costs associated with the treatment of pressure ulcers are high. Treatment cost is proportional to the severity of the ulcer, because the healing rate is slower and likelihood of complications is greater.⁵ The total cost for treating pressure ulcers is estimated to be 8.5 billion dollars in the United States⁹ and between 1.4 and 2.1 billion pounds in the United Kingdom, which represents 4% of that country's total health budget.⁵ The cost of pressure-ulcer treatment in Spain represents 5.2% of the total health-care

Correspondence: Dr WC Miller, Department of Occupational Science and Occupational Therapy, University of British Columbia, T325-2211 Wesbrook Mall, Vancouver, British Columbia, Canada V6T 2B5.
E-mail: bcmiller@telus.net

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expenditure.¹⁰ It is estimated that the cost of treating pressure ulcers for individuals with SCI may represent one-quarter of their total cost of care.¹¹

Pressure ulcers have a serious impact on the individual and those around them. Qualitative research has identified that pain is a key feature.^{3,4} As well, pressure-ulcer treatment often necessitates long-lasting activity modifications and restrictions that can have a negative psycho-social impact on the individuals and their families.^{3,4} Other sequellae include social isolation, alteration of body image, lost income, odor and drainage.⁶ In some individuals, pressure ulcers may prove fatal.¹²

A large number of risk and protective factors for the development of pressure ulcers have been described for individuals with SCI. Some of the risk factors that have been identified include: being underweight,¹³ smoking,^{13,14} lower level of activity,^{14,15} incontinence,^{14,15} pulmonary disease,^{14,15} decreased albumin,^{14,15} decreased mobility,^{14,15} extent of paralysis,^{14,15} increasing age,¹⁴ use of medications for sleep,¹⁶ impaired cognitive function,¹⁴ diabetes,¹⁴ living in a hospital or nursing home,¹⁴ spasticity,¹⁴ renal disease,¹⁴ and low hematocrit.¹⁴ Protective factors identified include completing a college degree,¹³ being married,¹³ being employed,¹³ exercise¹⁶ and healthy diet.¹⁶

Clinical practice guidelines for the prevention and treatment of pressure ulcers following SCI advocate the use of risk assessments, which include many of the preceding risk factors, to improve practice in this area.¹⁷ Nixon and McGough¹⁸ identified over 38 such risk assessment instruments. Despite the abundance of available tools, very few have been tested psychometrically¹⁸ especially with individuals with SCI.

It is essential to validate tools with their intended population(s), because validity is not an innate property of a test, but instead represents the evidence that supports the interpretation of the test scores.¹⁹ Tools should be assessed within specific populations,²⁰ because a measure that is reliable with one patient group may be unreliable with another. Moreover, measures intended for one population may lack content validity when used with a different population.²¹ For example, pressure-ulcer risk factors for elderly individuals are often different from those with SCI.²² Finally, in terms of utility, an instrument that is relatively easy to administer with one population may be difficult to administer with another. The purpose of this review was to critically evaluate pressure-ulcer risk assessment tools in terms of their reliability, validity and utility for individuals with SCI.

Methods

A review of pressure-ulcer risk assessment scales used with individuals with SCI was systematically conducted. To be included, measures had to have been assessed in at least one study published in a peer-reviewed journal that examined the psychometric properties of the instrument using a sample of individuals with SCI. Study-specific scales designed for intervention trials were not included in this

review, because they are difficult for researchers and clinicians to obtain and frequently use an alternate method of calculating reliability co-efficients than would be used with tools intended for a variety of users.²³ Additionally, only papers written in English were considered.

Search strategy

MedLine, CINAHL, Embase, HaPI, Psycinfo, Sportdiscus and Cochrane electronic databases were searched (1986 to January 2007) to locate papers describing these measures. Additional searching was conducted by reviewing the references of papers obtained from the electronic search. The key word 'spinal cord injury' was used across each of the databases, while the following terms varied depending on the database used: validation studies, instrument validation, external validity, internal validity, criterion-related validity, concurrent validity, discriminant validity, content validity, face validity, predictive validity, reliability, interrater reliability, intrarater reliability, test-retest reliability, reproducibility, responsiveness, sensitivity to change, evidence-based medicine, outcome measures, clinical assessment tools, scales, measures, pressure ulcer, decubitus, pressure sore and pressure wound. A database file was established using RefWorks to organize potential articles of interest. After eliminating duplicate papers two trained data extractors reviewed each title to determine which abstracts should be reviewed then the abstracts were reviewed to determine which papers should be extracted. When disagreement occurred, a third extractor was consulted to break any ties.

Assessing the tools

Data were extracted from papers by a team of reviewers using a data extraction form designed to record the reliability, validity, interpretability, feasibility and acceptability of the tool items. The methods and standards of data extraction were based on the work of Fitzpatrick *et al*.²⁴ Using the criteria noted in Table 1, the psychometric and utility properties reported for each tool were summarized as excellent, adequate or poor. 'Excellent' indicated that the tool was outstanding in regard to that criteria, 'poor' indicated that serious deficiencies were noted and 'adequate' indicated the 'tool rates somewhere in between'²⁵ (p. S15). For example, a validity score of excellent corresponded to correlations above 0.60 with similar measures or convergent constructs. Respondent burden was considered excellent if administration time was brief (<15 min) and the measure was well accepted by persons with disabilities. Interpretability was considered excellent if the tool was easy to administer, score and interpret. When validity correlations varied across studies (for example, poor in one study and adequate in another), results were summarized as a range (for example, poor—adequate).

Results

Our review identified four studies that provided validity evidence for seven tools that have been tested with

Table 1 Criteria for rating outcome measures

Criterion	Definition	Standard
Reliability	Reliability is the degree to which the score is free from error	ICC and Kappa for inter/intra and test-retest ratings are: excellent (≥ 0.75), adequate (0.4–0.70) or poor (≤ 0.40) ^{25,26}
Validity	The extent to which an instrument measures what it purports to measure	Construct/convergent and concurrent correlations: excellent (≥ 0.60), adequate (0.31–0.59), poor (≤ 0.30) ²⁵ ROC analysis—AUC: excellent (≥ 0.90), adequate (0.70–0.89), poor (< 0.70) ²⁶
Respondent burden	The ease with which a patient can complete the measure	Excellent (brief ≤ 15 min and acceptability high), adequate (either longer (but appropriately so) or some reported problems with acceptability), poor (both length and acceptability are problematic) ²⁵
Administrative burden	The ease with which scores can be calculated and understood.	Excellent (scoring by hand and resulting metric relevant and interpretable for researcher, clinicians and clients) adequate (computer scoring, lack of detail for scoring criteria, more obscure interpretation), poor (costly and/or complex scoring and/or interpretation) ²⁵

individuals with SCI. All of these studies used retrospective data extracted from medical records to determine each subject's risk assessment scale score and presence of pressure ulcers. The measures in these studies included the Abruzzese,²⁷ Braden,²⁸ Gosnell,²⁹ Norton,³⁰ SCIPUS,¹⁴ SCIPUS-A¹⁵ and Waterlow.³¹ A brief overview of each instrument is provided in Table 2. The Norton, created in 1962, is the original pressure-ulcer risk assessment tool and served as a template for many other scales.¹⁸ Overall, the measures cover between 5 and 8 domains. The domains of mobility, activity and continence are common across all measures and nutrition (or albumin level) is a domain in six of the seven scales. The SCIPUS tool has the most items,¹⁵ including many unique items such as autonomic dysreflexia and living setting. The SCIPUS and SCIPUS-A also include items that require laboratory tests of albumin levels, serum creatinine and blood glucose. The SCIPUS and SCIPUS-A were developed exclusively for individuals with SCI using regression analysis to identify variables, collected via retrospective chart review that predicted pressure-ulcer development.^{14,15} Although the risk factors considered for inclusion were not described for the SCIPUS,¹⁴ over 50 potential pressure-ulcer risk factors were evaluated for the SCIPUS-A of which 21 were explicitly described.¹⁵ Those factors included age, tobacco use, alcohol history, nutritional support, level of activity, mobility, mental status, urinary continence, nutritional status, moisture, 'friction and shear', and 10 laboratory blood work variables.¹⁵ For the SCIPUS and SCIPUS-A, the same data were used to generate the scale and to test their validity.^{14,15}

For all scales, items are scored using item-specific descriptive criteria, but the amount of detail provided varies considerably across measures. The Norton and Waterlow provide the least detailed scoring instructions. Both offer only one or two word descriptions for each score. For instance, in the Norton, the domain of physical condition is illustrated by four descriptors 'Good 4, Fair 3, Poor 2, Very Bad 1'³⁰ (p. 225). In contrast, with the Braden,²⁸ the tool that provides the most complete operational definitions, a score of 1 for the domain of sensory perception is described as 'Completely limited: unresponsive to painful stimuli, either

because of either unconsciousness or severe sensory impairment, which limits ability to feel pain over most of body surface' (p. 206). The Abruzzese, SCIPUS and SCIPUS-A measures fall in between, in that although most items are well operationalized, items such as moisture and extent of paralysis for the SCIPUS-A and impaired cognitive function for the SCIPUS are not.

The validity evidence for these instruments is described in Table 3. This includes sensitivity and specificity, area under the curve (AUC), which measures how well the test predicts who will and will not develop a skin ulcer, construct and concurrent validity results. In light of differences in study populations, the psychometric data are presented on per study basis. Validity results tended to vary across studies and between measures. The Braden had the best AUC,³³ but poor construct validity in terms of stage of first pressure ulcer ($r = 0.03$) and conflicting concurrent validity with the Norton ($r = 0.48$) and Waterlow ($r = -0.06$) scales.³² The AUCs ranged from 81% for the Braden to 72% for the Norton.³³

In terms of utility, a number of characteristics are common across measures, as noted in Table 4. All of the scales are available for free either electronically on the World Wide Web or from the original articles in which they were published. Most authors do not indicate that training is required to use the measure, although training for the Braden is suggested.²⁸ A video-taped manual is available for the Braden scale.

All scores are easily calculated by hand based on scoring criteria provided by their authors. For all scales, a total score is computed by adding item scores together. For most scales, higher scores indicate increased risk of developing a pressure ulcer, although for the Braden and Norton, the scoring is reversed and lower scores are indicative of higher risk. For the Norton, Braden, Gosnell and Abruzzese scales, scores are consecutive (1, 2, 3, 4 and so on) and one score per domain is allowed. The scoring is slightly more complicated for the SCIPUS, SCIPUS-A and Waterlow. For the SCIPUS and SCIPUS-A scale, scores are weighted for all domains (for example, extent of paralysis is scored as 0 = none, 1 = paraparesis, 4 = quadriplegia, 8 = paraplegia and 10 = quadriplegia). These weighted values were based on

Table 2 Description of measures

Measure	Abruzzese ²⁷	Braden ²⁸	Gosnell ²⁹	Norton ³⁰	SCIPUS ¹⁴	SCIPUS-A ¹⁵	Waterlow ³¹
Development	Based on Norton and Gosnell scales	Developed on the basis of a conceptual schema of pressure sore development	Items were identified based on research by Norton	Developed on the basis of the researchers' clinical expertise with a geriatric hospital population	Intended for a rehabilitation setting, items for the scale were identified based on statistical analysis of data from 176 individuals with SCI	Intended for acute hospitalization items for the scale were identified based on statistical analysis of data from 226 individuals with SCI	Created to provide better sensitivity and specificity than the Norton by increasing the number of items used ³²
Domains/items (one item per domain unless otherwise noted)	Seven domains (1) General health (2) Mental status (3) Activity (4) Mobility (5) Continence (6) Nutrition (a) Oral nutrition intake (b) Oral fluid intake (7) Predisposing diseases (vascular disease, neuropathies, diabetes, anemias, and so on)	Six domains (1) Sensory perception (2) Moisture (3) Activity (4) Mobility (5) Nutrition (6) Friction and shear	Five domains (1) Mental status (2) Continence (3) Movement control (4) Ability to ambulate (5) Process of food intake Evaluation includes recording of vital signs, skin condition and medications, but these are not scored	Five domains (1) Physical condition (2) Mental condition (3) Activity (4) Mobility (5) Continence	Seven domains (1) Level of activity (2) Levels of mobility (3) Severity of SCI (a) Complete SCI (b) Autonomic dysreflexia or severe spacticity (4) Urine incontinence or constantly moist (5) Pre-existing conditions (a) Age (b) Tobacco use/smoking (c) Pulmonary disease (d) Cardiac disease or glucose > 110 mg dl ⁻¹ (e) Renal disease (f) Impaired cognitive function (6) Residence in a nursing home or hospital (7) Nutrition (a) (albumin < 3.4 or total protein < 6.4) ¹⁵ (b) Anemia (hematocrit < 36.0%)	Six domains (1) Extent of paralysis (2) Incontinence (a) Moisture (b) Continence (3) Mobility (4) Level of activity (5) Nutrition (a) Serum creatinine (b) Albumin (6) Pre-existing conditions (a) Pulmonary disease	Eight domains (1) Age (2) Sex (3) Body build (4) Appetite (5) Continent of urine and feces (6) Mobility (7) Skin appearance in risk areas (8) Special risks (disorders associated with tissue malnutrition (terminal cachexia, cardiac failure, peripheral vascular disease, anemia), neurological deficits, smoking, medication, recent surgery or trauma (including SCI)
Procedure	Each domain item has four response choices. Scoring options are 0, 1, 2, 3 or 0, 1, 4, 6 depending on the item. Scores are summed to produce a total score between 0 (best prognosis) and 30 (worst prognosis)	Each domain item is given a rating of 1–3 or 4 based on a domain-specific ordinal scale. Scores are summed to produce a total score between 6 (worst prognosis) and 23 (best prognosis).	Each domain item is given a rating of 1–4 on a domain-specific ordinal scale. Scores are summed to produce a total score between 5 (worst prognosis) and 20 (best prognosis)	Each domain item is given a rating of 1–4 on a domain-specific ordinal scale. Scores are summed to produce a total score between 5 (worst prognosis) and 20 (best prognosis)	Most items are scored dichotomously as either present or absent, but 4 items have three response options Scores are summed to produce a total score between 0 = best prognosis and 25 = worst prognosis	Response categories are either dichotomous (present/absent; score = 1–2 or 0, respectively) or have 3–5 options Scores are summed together to create a summary score between 0 = best prognosis and 25 = worst prognosis	Items are scored as either dichotomous (yes/no) or on domain-specific scales that range from 0 or 1 to 3–5 Scores are summed to create a total score between 3 (best prognosis) and 45 (worst prognosis)
Cutoff scores	≥ 12	≤ 16 originally proposed. ≤ 10 has been suggested for SCI ¹⁵	—	≤ 14	≥ 6	≥ 18	10+ = risk 15+ = high risk 20+ = very high risk

Table 3 Validity of the pressure ulcer scales

Measure	Abruzzese	Braden	Gosnell	Norton	SCIPUS	SCIPUS-A	Waterlow
Sensitivity	21.8%	74.7%	18.4%	5.8%	36.8%	88.5%	—
Specificity	84.6%	56.6%	90.4%	95.6%	84.2%	59.0%	—
Cutoff	ND	≤10	ND	ND	ND	≥18	—
Accuracy ^{a15}	62.2%	62.3%	62.2%	60.8%	65.9%	71.0%	—
Sensitivity	—	—	—	—	75.6%	—	—
Specificity	—	—	—	—	74.4%	—	—
Cutoff	—	—	—	—	≥6	—	—
+ Predictive	—	—	—	—	92.4%	—	—
– Predictive ^{b14}	—	—	—	—	42.7%	—	—
AUC ^{c33}	—	81% (74–88, 95% CI)	—	72 (64–81, 95% CI)	—	78 (70–85, 95% CI)	76% (68–84, 95% CI)
Concurrent ^{d32}	—	0.48 (Norton) –0.06 (Waterlow)	—	–0.56 (Waterlow) –0.49 (Braden)	—	—	–0.56 (Norton) –0.06 (Braden)
Construct (<i>r</i>)							
Stage of 1st ulcer ^a	0.241	–0.353	0.254	–0.192	0.343	0.488	—
Stage of 1st ulcer ^d	—	0.03	—	–0.28	—	—	0.38
No. of ulcers ^a	0.212	–0.431	0.297	–0.197	0.339	0.519	—

Abbreviations: AUC, area under the curve; CI, confidence interval; ND, not described/reported.

^aFor individuals with SCI during initial hospitalization.

^bFor individuals with SCI during rehabilitation.

^cFor individuals with SCI in a UK spinal cord injuries unit.

^dFor a stratified sample of individuals with SCI with a pressure ulcer.

the relative value coefficients (rounded for simplicity) from a logistic regression model of factors associated with pressure ulcers in individuals with SCI.¹⁵ For the Waterlow most items are weighted, and for many, multiple response categories can be selected. For example, when describing skin type as a risk factor, an individual could have 'dry' (= 1), 'oedematous' (= 1) and/or 'discoloured' (= 2) skin leaving an item-based score ranging from 1 to 4. As with the SCIPUS and SCIPUS-A, scores for each item are not consecutive, but no rationale for the weighting scheme is described.

A summary of the psychometric and utility evidence, according to the Andresen's criteria (Table 1)²⁵ is provided in Table 5. Validity evidence for all the measures ranged from adequate to poor. As all measures took less than 15 min to complete, and were generally non-invasive, most were deemed excellent in terms of their respondent burden. The exceptions were the SCIPUS and SCIPUS-A, which require blood tests. As response descriptions for at least some of the items in most measures were vague, administrative burden was found to be only adequate for all measures except the Braden, which was rated as excellent because of its detailed scoring guidelines.

Discussion

This review identified seven measures that had been tested with individuals with SCI. These included (1) three of the most commonly used and validated scales for pressure-ulcer prediction in the general population (the Braden, Norton and Waterlow);^{10,32} (2) two measures designed specifically for individuals with SCI during acute hospitalization (SCIPUS-A)¹⁵ and rehabilitation (SCIPUS)¹⁴ and (3) two lesser known and validated scales (the Abruzzese and the Gosnell).

The finding that no reliability testing has been conducted in the SCI population for all measures reviewed is a concern,

albeit not unexpected. In their systematic review of risk assessment scales in other populations, Pancorbo-Hidalgo *et al.*¹⁰ identified 12 measures that had been validated using controlled clinical trials or prospective cohort studies. They found no reliability data for 7 of the 12 scales included in their systematic review. Pearson's reliability coefficients for the Braden ranged from 0.83 to 0.99 across 13 studies in different settings (nursing homes, home care and hospital) and populations (orthopedics, geriatric and intensive care). In the three studies that examined the reliability of the Waterlow and Norton scales, reliability scores of $r = 0.99$ and % of observer agreement of 92.5 and 100% were reported, respectively. Because of their inclusion criteria,¹⁰ however, no studies with individuals with SCI were part of this review. Generalizability Theory indicates that these results cannot be applied to individuals with SCI,³⁴ which undermines confidence in these scales to reproduce stable results over time with this population.

Given that over 200 risk factors for developing a pressure ulcer have been reported for individuals with SCI,²² it is important to identify which ones need to be included in a measure to ensure adequate content validity. Although all of the measures included pressure-ulcer risk factors that have been identified for individuals with SCI,²² the SCIPUS and SCIPUS-A were the only instruments developed exclusively for individuals with SCI. The items included in these measures indicate that important risk factors in the general population are not the same for individuals with SCI during different phases of their recovery.¹⁵ The use of linear regression to identify items for the SCIPUS and SCIPUS-A is a strength of these measures, but the limited description of what variables were evaluated makes it difficult to determine if all important variables were considered.

A total of four studies looked at the validity evidence of these measures for individuals with SCI. Overall, no scale demonstrated excellent validity based on the published

Table 4 Utility of the pressure ulcer scales

Measure	Abruzzese	Braden	Gosnell	Norton	SCIPUS	SCIPUS-A	Waterlow
Administrative burden	Response descriptions for some scale items are very brief. No indication formal training is required	Detailed descriptions for items scoring provided by authors. Training suggested	Adequate response descriptions for most items provided. No formal training is required	Descriptors for item scoring are very brief. ³² No indication formal training is required	Response descriptions for some scale items are vague. No indication formal training is required	Response descriptions for some scale items are vague. No indication formal training is required	Response descriptions for some scale items are vague. No indication formal training is required
Respondent burden	5–10 min	5–10 min	Minimal respondent burden for additional items only 5–10 min	5–10 min	Minimal burden if test results for diabetes, albumin and hemocrit, and so on, are available 5–10 min	Minimal burden if test results for albumin and serum creatinine are available 5–10 min	No respondent burden 5–10 min
Advantages and/or limitations		Most well validated tool generally. ¹⁰ Additional information and a video-taped training manual are available at www.braddenscale.com			Items identified based on data from individuals with SCI	Items identified based on data from individuals with SCI	

Table 5 Summary of psychometric properties and utility evidence

Measure	Abruzzese	Braden	Gosnell	Norton	SCIPUS	SCIPUS-A	Waterlow
Reliability	—	—	—	—	—	—	—
Validity	Poor	Poor-adequate	Poor	Poor-adequate	Adequate	Adequate	Adequate
Respondent burden	Excellent	Excellent	Excellent (if skin status assessment is omitted)	Excellent	Excellent (if blood testing results already available, otherwise adequate)	Excellent (if blood testing results already available, otherwise adequate)	Excellent
Administrative burden	Adequate	Excellent	Adequate	Adequate	Adequate	Adequate	Adequate

— Indicates no information available for individuals with SCI.

criteria used in this review.^{25,26} Validity evidence for the Abruzzese and Gosnell was reported in only one study and results were poor.¹⁵ Validity evidence for the SCIPUS and SCIPUS-A was reported in two studies with adequate results. The SCIPUS-A had the best sensitivity and specificity, and construct validity of all measures tested for individuals during acute hospitalization, although the Waterlow was not included in this study.¹⁵ Similarly, the SCIPUS had good sensitivity and specificity for individuals during rehabilitation,¹⁴ but no comparisons were made between other instruments, so no definitive conclusions can be drawn. The sensitivity and specificity of the SCIPUS and SCIPUS-A scores may be inflated, because these scales were developed and tested using the same retrospective data. These scores would likely decrease, if the measure was tested again with a different sample. Validity evidence for the Braden, Waterlow and Norton was reported in three studies with mixed results. Although the AUC was adequate for all of these measures,³³ the Braden and Norton had poor construct validity³² and the concurrent validity of the Braden and Waterlow was poor.³²

Most measures were similar in terms of their utility. With the exception of the SCIPUS, all measures consisted of only eight items or less, so the administration time was very similar. For all instruments, the measure and scoring criteria are available for free and training was only recommended for the Braden scale. Subject burden for most tests is minimal unless results of blood tests are not already available, in which case, the SCIPUS and SCIPUS-A would involve additional respondent burden and might add administrative burden. For instance, in a community setting, administration of the SCIPUS would require a physician to order blood testing and either the individual to visit a laboratory, or have home testing arranged, all of which would add substantially to the direct and indirect cost of this measure. The Braden scale offers the best operationalization of response categories, while response descriptions for at least some of the items from other scales were vague. For example, with the SCIPUS-A raters are to indicate whether the patient experiences moisture rarely, occasionally, very often or constantly; these frequencies are not quantified, however, which may lead to misclassification. Better operationalization of the responses would permit clinicians to extract more useful information.

Comparing the validity results of this review with those obtained by Pancorbo-Hidalgo *et al.*¹⁰ reveals some interesting similarities and differences. The weighted means of sensitivity and specificity of the Braden (57.1 and 67.5%, respectively) noted by Pancorbo-Hidalgo *et al.* are somewhat similar to the sensitivity and specificity noted by Salzberg *et al.*¹⁵ for individuals with acute SCI (sensitivity 74.7% and specificity 56.6%). The sensitivity and specificity of the Norton (46.8 and 61.8%), however, are markedly different (sensitivity 5.8% and specificity 95.6%). These comparisons emphasize the need for population-specific psychometric testing.

The retrospective method used to evaluate validity in all of the studies in this review is a concern. Given problems with the accuracy and completeness of medical records,³⁵ a retrospective analysis of medical data may not represent a

true picture of either the patient's ulcer severity or risk for developing an ulcer.

Recommendations

This review highlights the difficulties inherent in the selection of a pressure-ulcer risk assessment scale for individuals with SCI. Although the SCIPUS and SCIPUS-A have the advantage of being specifically tailored to individuals with SCI and their sensitivity and specificity results are good, they are limited by their lack of reliability data, and the fact that validity testing was conducted with the same data used for their development. Consequently, the use of SCIPUS and SCIPUS-A cannot be recommended without further psychometric testing. Given that the Braden scale performed similarly to the SCIPUS-A in testing, had the largest AUC³³ and, generally, was the most well-validated instrument;¹⁰ the Braden scale seems to be best tool currently available, although it would also benefit from additional testing with individuals with SCI.

A variety of future studies are suggested by the findings of this review. Obviously, all of these measures require reliability testing with individuals with SCI. As well, prospective studies that allow head-to-head comparison of these risk assessment scales would represent a far more robust method to evaluate their concurrent and construct validity. The use of multiple regression analysis to identify scale items seems promising, but all of the factors that are evaluated need to be identified explicitly, as other factors, including psychological factors, socioeconomic status, existence of a previous skin ulcer and etiology of the SCI, may be more critical risk factors for individuals with SCI than for individuals from other populations.²² Because of how these tools are used in clinical settings, responsiveness studies are required and more importantly, studies that evaluate the effectiveness of these tools for pressure-ulcer prevention are needed. Such studies would provide a critical element of validity that is missing among these tools in identifying individuals at risk of developing a pressure ulcer and preventing their occurrence. This is especially important, given that Keast *et al.*³⁶ in their review of best practice recommendations for the prevention and treatment of pressure ulcers, found only level four evidence (expert opinion) supported the use of validated risk assessment tools for the prevention of pressure ulcers.

Limitations

This review included only, peer-reviewed studies written in English. Although a variety of search terms were employed, it is possible that some studies may not have been included in the review, as gray literature, conference abstracts and studies published in other languages were not accessed.

Conclusion

This review identified seven pressure-ulcer risk assessment tools that had been validated with individuals with SCI. Reliability and responsiveness evidence for individuals with SCI was absent in the literature. Validity results indicate that

five of the measures demonstrate adequate validity and two are poor. Although the SCIPUS and SCIPUS-A are promising scales, the Braden seems to be the best tool currently available. Given the importance of healthy skin to individuals and the personal and societal costs of pressure ulcers, further research is vital to evaluate the psychometric properties of these instruments and determine their effectiveness in pressure-ulcer prevention and treatment.

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