A Critical Review of Grading Systems: Implications for Public Health Policy

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Abstract
Grading instruments are an important part of evidence-based medicine and are used to inform health policy and the development of clinical practice guidelines. They are extensively used in the development of clinical guidelines and the assessment of research publications, having particular impact on health care and policy sectors. The positive effects of using grading instruments are, however, potentially undermined by their misuse and a number of shortcomings. This review found eight key concerns about grading instruments: (1) lack of information on validity and reliability, (2) poor concurrent validity, (3) may not account for external validity, (4) may not be inherently logical, (5) susceptibility to subjectivity, (6) complex systems with inadequate instructions, (7) may be biased toward randomized controlled trial (RCT) studies, and (8) may not adequately address the variety of non-RCTs. This narrative review concludes that there is a

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need to take into account these criticisms and domain-specific limitations, to enable the use and development of the most appropriate grading instruments. Grading systems need to be matched to both the research question being asked and the type of evidence being used.

**Keywords**
grading evidence, evidence-based medicine, epidemiological studies, policy, clinical guideline

**Background**

Systems that synthesize and grade the quality and robustness of scientific results are used by researchers and policy makers to develop summaries or conclusions about the state of science on particular topics. Hence, grading systems can play a critical role in whether and how research findings are ultimately translated into policy and practice. The application of a grading system to a body of evidence can provide an authority to the findings of a review or report that is used to justify changes in policy and practice at national levels. It is therefore essential that scientists are aware of the potential limitations of these systems and how they may ultimately impact whether scientists’ research findings are used to inform policy and practice. Here, we present a narrative synthesis and commentary of issues raised in recent critical and systematic reviews of evidence grading systems. We focus on the use of grading systems primarily in epidemiological research and critically evaluate their limitations and potential for misuse.

**Search Methodology**

Our search included MEDLINE and Cochrane Library (Methodology Register) up to 2013, and selected articles up to 2015, that reported and reviewed grading systems, quality rating scales, observational epidemiological evidence, randomized controlled trials (RCTs), critical appraisals of evidence grading, and systematic reviews. Initial attempts to perform a systematic search using a range of search terms (including key words “grading system,” “grading evidence,” “rating quality of evidence,” and “evidence-based medicine review”) did not yield useful results. In keeping with the data collection challenges outlined in West et al. (2002, p. 75), results from these early search attempts tended to return primary studies
where a grading instrument had been used. As such, a snowball approach was used to retrieve papers that reviewed, discussed, and/or critiqued grading instruments. This was achieved primarily through the use of citations and bibliographies. Data collection ceased once saturation was achieved and no new results emerged. This technique ultimately yielded 52 papers, which were included in this review. It was agreed that a narrative approach was more appropriate, given the subject matter and that a systematic literature review was not feasible.

**Definition and Characteristics of Grading Systems**

There are many instruments available for grading evidence (Bai, Shukla, Bak, & Wells, 2012; West et al., 2002). Broadly speaking, grading instruments are used to quantify the quality of research evidence. Gugiu (2015) says a grading system “is a qualitative rubric employed by researchers and clinicians for the purpose of developing an empirical base of evidence in support (or opposition) of a particular course of treatment. Hence, grading systems have come to define what researchers view to be credible evidence. Moreover, the study designs promoted by these guidelines are deemed by many researchers, including government agencies, as more worthy of receiving research funding than other study designs” (p. 150). The usual output of a grading instrument is a rating score or “grade” that allows the user to compare the quality of different pieces of research. Some also provide a “strength” rating of recommendations based on the given topic. Grading instruments supply criteria for deriving grades as well as some form of instruction as to how to apply those criteria. Some instruments are designed for the grading of multiple kinds of research (e.g., RCTs and non-RCTs including observational studies), while others are specialized for use in evaluating only specific research designs (e.g., RCTs or observational studies). Additionally, instruments can be divided into those designed for rating a single piece of evidence, such as an individual research paper, and those designed to evaluate a body of evidence, such as a systematic review or a meta-analysis (see Table 1, e.g., of grading instruments designed for different purposes).

Grading systems meet the need of scientists and policy makers to derive some conclusions from the often overwhelming volume of information published on specific topics. However, the use of grading instruments itself implies certain assumptions about their validity which many users may not question or be in a position to evaluate. A potential area of concern lies with the fact that it is possible to misuse grading instruments or misinterpret the
meaning of ratings, if there is insufficient awareness of their measurement properties and validity. This article offers a critical review of grading instruments and provides a thoughtful debate about their use and potential for misuse, especially in relation to public health research.

Critics of the grading approach call into question the role that grading should play in both evidence-based medicine (EBM) and evidence-based policy (Glasziou, Vandenbroucke, & Chalmers, 2004; Gugiu, Westine, Coryn, & Hobson, 2013). Stemming often from the social sciences, philosophy, and methodology, there is a risk that discussion of grading systems is not accessible to data-based researchers. Here we aim to explain the key criticisms for those using grading systems in substantive research. These points are described in the following sections.

**Grading Systems May Lack Information on Their Validity and Reliability**

Many of the existing grading instruments have not been validated, and hence, there is a lack of data to support the claim that these systems are

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<th>Table 1. Examples of Grading Instruments.</th>
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<td><strong>Purpose</strong></td>
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*Note.* RCT = randomized controlled trial.
effective, avoid bias, and generally work in the way they claim to work. In a review of 44 grading scales published through a peer-review process, Crowe and Sheppard (2011) found an alarming number did not undertake checks for concurrent validation or attempt construct validation, and the majority did not report data on their reliability. A systematic review of 121 published critical appraisal tools similarly found that few had provided evidence of item validity or reliability of use (Katrak, Bialocerkowski, Massy-Westropp, Kumar, & Grimmer, 2004).

Some authors have argued that the lack of validation of the most widely used grading instrument, grading of recommendations, assessment, development and evaluation (GRADE), is reason to argue against its utility (Kavanagh, 2009). Gugiu and Gugiu (2010) contend that EBM is based on the ability of reviewers to discern and differentiate between evidence, and if the systems used to do so are not actually proven to be valid, it amounts to an undermining of both EBM principles and scientific progress.

It can plausibly be argued that there is no standard against which a grading system could be validated because they themselves are set up as the gold standard, and implicit in the use of grading systems is that there is a metric of quality and strength of association, if not certainty, associated with types of evidence. It is theoretically possible to use a meta-analysis approach to compare the grading outcomes of different systems for a given set of studies—with the assumption that greater variability in effect sizes at each level of a grading system would represent greater bias. Nevertheless, there are exceptions where authors have performed tests of face validity (Downs & Black, 1998; Kim et al., 2013), content validity (Shea et al., 2009; Slim et al., 2003), construct validity (Kim et al., 2013; Oxman & Guyatt, 1991), and criterion validity (Downs & Black, 1998). Concurrent and external validity are explored further below.

**Grading Systems May Have Poor Concurrent Validity**

Concurrent validity is established when instruments designed to measure the same underlying construct are highly correlated. For example, if three questionnaires measuring depressive symptoms produced highly intercorrelated scores, we could conclude that they have high concurrent validity. If grading systems have concurrent validity, then different systems designed for similar purposes should be highly correlated. In addition, the interrater agreement of each scale should be high, as this would impact on scale reliability and the accuracy of measures of concurrent validity. Unfortunately, interrater reliability and concurrent validity of most scales are low...
A review of 16 quality assessment tools (Colle, Rannou, Revel, Fermanian, & Poiraud, 2002) compared raters’ findings for a systematic review of exercise therapy and lower back pain across different scales. Low levels of interrater agreement were observed across the majority of tools, and there was low correlation between the different tools used, resulting in widely different interpretations of the results. The most severe example showed one scale rating only 5 of the reviewed RCTs as high quality, while 17 were rated as high quality by another scale. Another study comparing the quality rating of a systematic review on the impact of tai chi on bone mineral density in postmenopausal women (Alperson & Berger, 2011) found that when using two different instruments (Oxman’s criteria vs. the Jadad Scale), the research was either dismissed as low quality or conversely rated as safe and effective.

Correlation between scales designed to evaluate the quality of clinical trials was evaluated in a study that conducted the same meta-analyses 25 times using weighted scores from 25 different quality rating scales in meta-regression analyses (Jüni et al., 1999). The median quality of the trials ranged from 32.5% to 82.9% of maximum score across the 25 analyses. Across the 25 scales, there was wide variation in the pooled relative risks from trials rated as high quality, leading to significant discrepancy in the conclusions drawn depending on the scale used. The authors note importantly that methodological elements central to trial quality need to be specified a priori (Jüni et al., 1999). In clinical trials, there will always be some methodological criteria that are common, but there are also domain-specific methodological characteristics that need to be measured.

A systematic review of 121 published critical appraisal tools in the context of allied health determined that 87% were created for a specific study design and that there was wide variation in the items used (Katrak et al., 2004), and this variation is likely to explain why their application leads to variation in results. Two reviews of tools used to assess nonrandomized studies also found a lack of between-instrument consistency in terms of the domains used to assess study quality and the weighting of criteria within those domains (Sanderson, Tatt, & Higgins, 2007; Shamliyan, Kane, & Dickinson, 2010).

Despite the reliance of clinical practice guidelines committees on grading systems such as GRADE, there is evidence that the use of different instruments to evaluate RCTs may lead to different conclusions and clinical recommendations. For example, the impact of using different quality assessment scales
on practice guidelines has been evaluated in the field of cancer (Brouwers et al., 2005). A study of four validated scales found that the number of RCTs evaluated as high quality or low quality depended on which scale had been used (Brouwers et al., 2005) and that the quality rating of studies was not predictive of the pooled odds ratio in the final meta-analyses. The authors of this study decided that they would not recommend the use of quality ratings in the development of clinical guidelines in their specific clinical area.

**Grading Systems May Not Account Adequately (or at All) for External Validity**

External validity is primarily used to refer to the generalizability of research results to real-life populations but can also include the feasibility of applying findings in the real world and the sustainability of an intervention over time (Persaud & Mamdani, 2006). Some argue that external validity is as important to the evidence rating process as internal validity especially in the clinical practice sphere (Persaud & Mamdani, 2006). It has been argued that recommendations formed by grading instruments should take into account such factors as the pros and cons for the populations that would be impacted, costs to individuals as well as health systems, difficulties with implementation of a health intervention, differences in applicability across different regional settings, and more (Barbui et al., 2010; Brouwers, Somerfield, & Browman, 2008). However, grading instruments are often designed to focus solely on scientific robustness and do not include any metrics for evaluating the external validity of findings.

Black (1996) argued that external validity is more easily derived from qualitative and observational evidence, which provide context and applicability data as well as tending to be larger and more inclusive with their study participants than RCTs. The need for incorporating a measure of external validity into evidence grading methods is highlighted by studies that have shown that RCTs graded as high quality often use highly selected samples when compared to population rates of those meeting eligibility criteria, resulting in their samples being homogenous and unrepresentative, thereby reducing its applicability (e.g., Persaud & Mamdani, 2006; Rothwell, 2005; Sanson-Fisher, Bonevski, Green, & D’Este, 2007).

**Grading Systems Are Not Necessarily Inherently Logical**

Grading systems sometimes imbue the synthesis of research findings with “academic authority,” and there is often an implication that there is an
ontological basis for their hierarchy. In fact, there is no essential way in
which different elements of quality combine to create higher quality. It has
been argued that grading systems are not necessarily inherently logical
(Kavanagh, 2009). For example, GRADE requires that a non-RCT has a
significantly higher effect size before its evidence is elevated—which effec-
tively rewards non-RCT studies with biased effect sizes over well-
conducted non-RCTs with effect sizes more in line with those generated
by well-conducted RCTs. Furthermore, arriving at a final grade level
involves trading off different elements against each other to establish the
level of quality. For example, sparse data can be traded off against a dose–
response effect even though these two characteristics are not interchange-
able. Similarly, the American College of Chest Physicians rating system
automatically elevates evidence from RCTs with major methodological
flaws above evidence from non-RCTs regardless of quality. This also limits
its use in areas such as public health where RCTs may be impractical or
invalid (Atkins et al., 2004).

Grading Systems Are Susceptible to Subjectivity and Low Interrater
Reliability

Another key criticism of grading instruments is their susceptibility to sub-
jectivity (Baker, Potter, Young, & Madan, 2011; Barbui et al., 2010;
Brouwers et al., 2008; Calderón et al., 2006). By subjectivity, we do not
mean a reliance on expert judgment as part of the assessment process,
though this is an explicit feature of some instruments, but rather that there
is room for interpretation of the grading guidelines which can mean the
assessor’s own background, skills, or views can have a unique impact on
how guidelines are operationalized. This problem has been raised regarding
a large number of grading instruments. Prior to the release of GRADE, the
GRADE working group undertook a study of six prominent grading sys-
tems, evaluating them against a predetermined set of criteria (Atkins et al.,
2004). Twelve assessors were recruited to evaluate the systems based upon
preagreed criteria about their sensibility. The conclusions of the assessors
were divergent and highlighted not only the inconsistencies among different
grading systems but also how great an impact the experience of those using
the systems can have. Different professional backgrounds, perhaps even
personality types, may gravitate toward different systems. Furthermore,
they may use these systems in different ways suited to their knowledge and
sensibilities. Qualitative research confirms that subjectivity and reliance on
value judgments are a serious problem for grading instrument users
Calderón et al., 2006). Nineteen evidence raters reported to be highly experienced with the evaluation of clinical guidelines and were given training on GRADE before undertaking a rating task. Nonetheless, participants reported that they felt “professional background, prior experience and the degree of leadership” all conditioned their responses to some degree (Calderón et al., 2006). Susceptibility to subjectivity is one potential cause of low levels of interrater reliability. While there is often a lack of testing performed to verify reliability levels, low interrater reliability has indeed been found with several instruments (Atkins et al., 2004, 2005).

Grading Systems Often Have Inadequate Instructions and Are Overly Complex

Vulnerabilities to subjectivity may stem from a failure to provide clear instructions and define terms, especially in light of how complex some of the grading instruments are (Baker et al., 2011; Barbui et al., 2010; Ibargoyen-Roteta et al., 2010; Kavanagh, 2009). In their review of 121 critical appraisal tools, Katrak, Bialocerkowski, Massy-Westropp, Kumar, and Grimmer (2004) found that more than half (57%) of the tools assessed failed to provide guidelines regarding their administration. In a study by Baker, Potter, Young, and Madan (2011) comparing the use of three different grading instruments (GRADE, the Scottish Intercollegiate Guidelines Network [SIGN], and the National Service Framework for Long-Term Conditions [NSF-LTC]), 2 of the 12 assessors refused to use GRADE at all because they felt the system was too complicated. Eighty percent of those who did use GRADE thought that training was required to use the tool properly (conversely, the NSF-LTC was considered to be too simplistic, while SIGN was thought to require too much replication). These kinds of issues should not necessarily be dismissed as the result of underexposure to grading systems or guideline development in general. Even expert guideline assessors who had been given training in GRADE, such as those in Calderón et al.’s (2006) study, felt that the method was not clear or easy to apply.

It has been suggested that the absence of instructions, or the vagueness of those supplied, could result in raters deliberately or unintentionally arriving at conclusions that suit their own preconceptions (Barbui et al., 2010; Brouwers et al., 2008). Even when some form of instruction is provided, key terms are often left undefined forcing raters to come up with their own interpretations. Gugiu and Gugiu (2010) point out that such vagueness implies that judgment about the meaning of terms such as “strength of
evidence” can largely be a reflection of the individual reviewer’s opinion of what constitutes strong evidence, rather than an objective view arrived at by scientific consensus. Apart from furthering susceptibility to bias, the lack of instructions can also result simply in the incorrect use of the instruments. In a comparison of GRADE, SIGN, and NSF-LTC, the research team found that the majority of assessors (68%) used the tools incorrectly in some way (6/10 for GRADE, 6/12 for SIGN, and 11/12 for NSF-LTC; Baker et al., 2011). Given a common lack of clear instructions, it is not surprising that many grading instruments are not used appropriately and that both interrater reliability and construct validity are low.

Some Grading Systems Are Biased Toward RCTs

RCTs are the best available method for eliminating bias (Guyatt et al., 2011) and where possible are often an optimal research design. However, it does not always follow that any RCT is necessarily better than other types of evidence. A number of researchers have suggested some of the mainstream grading instruments are inherently biased in favor of RCTs (Baker et al., 2011; Baker, Young, Potter, & Madan, 2010; Barbui et al., 2010; Gugiu, 2015). Typically with such instruments, an RCT would be automatically assigned the highest level grade, while non-RCT studies would be assigned a lower one. Although randomization is the best method of minimizing bias at the outset of a study, few grading systems take into account a non-RCT’s attempts to establish equivalent groups at baseline between treatment and control and differentiate this from non-RCTs that do not do this. Furthermore, authors have identified that grading systems do not consider some methodological limitations of RCTs such as dosage, differential attrition between groups, or flawed randomization (Gugiu, 2015). Depending on the system, there may be a means for upgrading or downgrading the grade based on fairly generic criteria. Even with the ability to upgrade and downgrade ratings, critics have pointed out that the majority of grading instruments do not sufficiently allow for ranking of a poor RCT against a good non-RCT, or they grade flawed RCTs higher than strong non-RCTs (Gugiu & Gugiu, 2010). There is also a lack of recognition of the large methodological differences inherent to different study types that fall under the non-RCT or “observational” umbrella (Hartling, Fernandes, Seida, Vandermeer, & Dryden, 2012).

An additional criticism of the default high ratings for RCTs comes from the limitations in conducting them. Despite the fact that RCTs may be the gold standard in EBM, there are areas of research that cannot ethically be
undertaken as an RCT, or for which RCTs are inappropriate (Bagshaw & Bellomo, 2008). Although this fact does not change the importance of RCT methodology, it also means that there are areas of research which may not be appropriately evaluated if they are measured against a hierarchy that emphasizes RCTs.

In some circumstances, observational studies such as large population-based studies have higher external validity, providing more applicable research findings (Persaud & Mamdani, 2006). Further arguments are made that while RCTs may reduce the risk of selection bias, they are typically smaller in size. Observational studies in contrast tend to have a larger number of participants, can be ecologically more valid, and are able to include more diverse patient populations (i.e., from different ethnic and social backgrounds, greater range of ages, etc.; Lohr, 2004).

**Grading Systems May Not Adequately Address Different Types of Observational Research**

Grading systems have also been criticized for assuming that all non-RCTs are simply observational research, without taking into account the different types of non-RCT studies. Some non-RCTs, for example, matched samples, when carefully conducted, do approximate the equivalent of trial designs where participants are randomized (Gugiu, 2015). Gugiu makes the distinction therefore between equivalent controlled trials and noncontrolled trials (Gugiu, 2015).

**Implications of Limitations in Grading Systems for Public Health**

Evidence to inform effective public health policy requires a high level of external validity and knowledge of the long-term outcomes of public health interventions as well as the population prevalence and incidence of a given health problem (Bammer, Michaux, & Sanson, 2010). This type of knowledge cannot be gained through RCT methodology but is gathered using observational studies (Dreyer et al., 2010; Fleurence, Naci, & Jansen, 2010; Morrato, Elias, & Gericke, 2007; Orton, Lloyd-Williams, Taylor-Robinson, O’Flaherty, & Capewell, 2011).

Given the risks and limitations associated with using grading instruments, particularly for the rating of non-RCT studies, those involved in public health research and policy making have cause to be concerned. As outlined above, grading instruments are subject to weaknesses that allow different raters and different systems to result in disparate grading of the
same piece of research or body of evidence. To further illustrate this point
as regards epidemiological research in public health, we will use two exam-
pies to demonstrate instances where the use of a grading instrument has or
could result in the rating of arguably high-quality observational evidence as
low quality.

Risk Factors for Alzheimer’s Disease

In recent years, the U.S. Department of Health and Human Services
National Institutes of Health published a report on “Preventing Alzheimer’s
Disease and Cognitive Decline” (Williams, Plassman, Burke, Holsinger, &
Benjamin, 2010). In this report, evidence for several established risk factors
was rated as low because they could only be studied using observational
research designs, and the GRADE system was applied to the meta-analyses.
In the use of GRADE, as with many other grading instruments, RCTs
default to an initial “high”-quality rating while observational research
defaults to “low.” While it is possible to upgrade observational research
using GRADE, it appears to rely on the judgment of the researcher to assess
whether or not potential sources of bias in the research were reduced.
Regardless, there are two flaws in using GRADE for this assessment. One
is that a major limitation of the GRADE approach is that it does not consider
biomedical research that identifies mechanisms underlying the associations
identified in observational studies. Secondly, the rating process does not
take into account the necessity of relying on observational evidence in this
context and used a grading system that was arguably not optimally suited to
the task. For example, research on smoking is inherently nonexperimental
because it is not ethical to perform a study where participants would be
required to take part in behaviors known to be deleterious to their health
(such as would be the case with an RCT on smoking). Given this limitation,
observational studies are the strongest possible direct form of evidence
available in humans. However, evidence is also available from animal
studies and from RCTs of smoking cessation where cognitive function is
a secondary outcome (e.g., Almeida et al., 2011).

Helicobacter pylori and Gastric Ulcers

In the early 1980s, Marshall and Warren (1984) observed that patients with
chronic gastritis hosted a little-known bacterium that is now known as
Helicobacter pylori. At the time, the accepted scientific wisdom was that
stomach ulcers were usually caused by stress, spicy food, and too much
gastric acid, and most specialists in the field regarded Marshall and Warren’s findings with great skepticism. In fact, so much so that in 1984 and after a number of attempts at convincing the scientific community of the importance of these findings, Marshall decided to infect himself with a petri dish culture of the bacterium. Over the following 8 days, he became sick and developed acute gastritis. This diagnosis was confirmed by an endoscopy and biopsy which had also been performed before infection. There is no suggestion here that Marshall and Warren’s scientific evidence was downgraded by rating tools but rather that if it had been rated, it would have been penalized for its observational nature, despite the fact it was backed up by substantial laboratory culture evidence and that an RCT would not have been possible because the scientific views on this topic were so entrenched. Indeed, after failing to infect animals with the bacterium (which is now known to be present only in humans), Marshall and Warren were eventually awarded a Nobel Prize for this discovery (Warren, 2006).

**Summary**

Grading instruments are now an integral part of EBM and used extensively in the development of clinical guidelines and other far-reaching publications. Because of their importance to such a wide array of health-related organizations and professions, it is vital that their limitations and shortcomings be understood. Grading instruments are susceptible to misuse because of their complexity, insufficient instructions, and their reliance on the traditional evidence hierarchy that places RCTs at the apex irrespective of context. It must also be remembered that the majority of instruments have not been validated, and of those that have been subjected to tests of reliability, the results have tended to be unfavorable.

It should be borne in mind that grading instruments are not a uniform set of tools but vary widely and the choice of the appropriate grading system for a particular context is essential. The purpose of this article is to highlight the need for those who rely on EBM or policy to be aware of the potential shortcomings of the tools used to decide what research will form a particular evidence base and to question how those conclusions were derived.

The consequences of inaccurate grading are serious. If the strongest possible evidence can be graded as low due to ineffective use of grading instruments, it is possible that health practitioners and policy makers may conclude that there are no grounds to recommend reduction or cessation of behaviors known to be risk factors for causes of morbidity and mortality. It
also provides a possible avenue for the public or parties with vested interests to misinterpret or misuse evidence grades. For example, smokers or tobacco companies could directly interpret a “low evidence” rating of research that links smoking and Alzheimer’s disease as reassurance that their use of tobacco is not linked with increased risk.

Such possibilities highlight the need to apply the most appropriate grading instrument to both the research question being asked and the type of evidence being used. It has also been suggested by numerous researchers and members of peak bodies that certain fields might be better served by a specifically developed grading instrument to meet their needs (e.g., re Allied Health: Burnett, Kumar, & Grimmer, 2005; Katrak et al., 2004).

Regarding those instruments that already exist and those in development, the best case scenario would see that the issues raised by critics were addressed. This has the potential to result in improved instructions that minimize subjective interpretations, validation, and reliability testing that is transparently reported as well as consistently undertaken across grading instruments, and a reconsideration of the default ranking of RCTs as high and observational evidence as low in all circumstances.

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