Critical Appraisal Toolkit (CAT) for assessing multiple types of evidence

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Abstract

Healthcare professionals are often expected to critically appraise research evidence in order to make recommendations for practice and policy development. Here we describe the Critical Appraisal Toolkit (CAT) currently used by the Public Health Agency of Canada. The CAT consists of: algorithms to identify the type of study design, three separate tools (for appraisal of analytic studies, descriptive studies and literature reviews), additional tools to support the appraisal process, and guidance for summarizing evidence and drawing conclusions about a body of evidence. Although the toolkit was created to assist in the development of national guidelines related to infection prevention and control, clinicians, policy makers and students can use it to guide appraisal of any health-related quantitative research. Participants in a pilot test completed a total of 101 critical appraisals and found that the CAT was user-friendly and helpful in the process of critical appraisal. Feedback from participants of the pilot test of the CAT informed further revisions prior to its release. The CAT adds to the arsenal of available tools and can be especially useful when the best available evidence comes from non-clinical trials and/or studies with weak designs, where other tools may not be easily applied.


Introduction

Healthcare professionals, researchers and policy makers are often involved in the development of public health policies or guidelines. The most valuable guidelines provide a basis for evidence-based practice with recommendations informed by current, high quality, peer-reviewed scientific evidence. To develop such guidelines, the available evidence needs to be critically appraised so that recommendations are based on the “best” evidence. The ability to critically appraise research is, therefore, an essential skill for health professionals serving on policy or guideline development working groups.

Our experience with working groups developing infection prevention and control guidelines was that the review of relevant evidence went smoothly while the critical appraisal of the evidence posed multiple challenges. Three main issues were identified. First, although working group members had strong expertise in infection prevention and control or other areas relevant to the guideline topic, they had varying levels of expertise in research methods and critical appraisal. Second, the critical appraisal tools in use at that time focused largely on randomized controlled trials. For feasibility and ethical reasons, clinical trials on randomized controlled trials (RCTs) and non-randomized controlled trials. For feasibility and ethical reasons, clinical trials are rarely available for many common infection prevention and control issues (4,5). For example, there are no intervention studies assessing which practice restrictions, if any, should be placed on healthcare workers who are infected with a blood-borne pathogen. Working group members were concerned that if they used GRADE, all evidence would be rated as very low or as low quality or certainty, and recommendations based on this evidence may be interpreted as unconvincing, even if they were based on the best or only available evidence.

To address these issues, a review of existing critical appraisal tools was conducted. We found that the majority of existing tools were design-specific, with considerable variability in intent, criteria appraised and construction of the tools. A systematic review reported that fewer than half of existing tools had guidelines for use of the tool and interpretation of the items (1). The well-known Grading of Recommendations Assessment, Development and Evaluation (GRADE) rating-of-evidence system and the Cochrane tools for assessing risk of bias were considered for use (2,3). At that time, the guidelines for using these tools were limited, and the tools were focused primarily on randomized controlled trials (RCTs) and non-randomized controlled trials. For feasibility and ethical reasons, clinical trials are rarely available for many common infection prevention and control issues (4,5). For example, there are no intervention studies assessing which practice restrictions, if any, should be placed on healthcare workers who are infected with a blood-borne pathogen. Working group members were concerned that if they used GRADE, all evidence would be rated as very low or as low quality or certainty, and recommendations based on this evidence may be interpreted as unconvincing, even if they were based on the best or only available evidence.

The team decided to develop its own critical appraisal toolkit. So a small working group was convened, led by an epidemiologist with expertise in research, methodology and critical appraisal, with the goal of developing tools to critically appraise studies informing infection prevention and control recommendations.

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This article provides an overview of the Critical Appraisal Toolkit (CAT). The full document, entitled Infection Prevention and Control Guidelines Critical Appraisal Tool Kit is available online (6).

Overview

Following a review of existing critical appraisal tools, studies informing infection prevention and control guidelines that were in development were reviewed to identify the types of studies that would need to be appraised using the CAT. A preliminary draft of the CAT was used by various guideline development working groups and iterative revisions were made over a two year period. A pilot test of the CAT was then conducted which led to the final version (6).

The toolkit is set up to guide reviewers through three major phases in the critical appraisal of a body of evidence: appraisal of individual studies; summarizing the results of the appraisals; and appraisal of the body of evidence.

Tools for critically appraising individual studies

The first step in the critical appraisal of an individual study is to identify the study design; this can be surprisingly problematic, since many published research studies are complex. An algorithm was developed to help identify whether a study was an analytic study, a descriptive study or a literature review (see text box for definitions). It is critical to establish the design of the study first, as the criteria for assessment differs depending on the type of study.

Definitions of the types of studies that can be analyzed with the Critical Appraisal Toolkit*

Analytic study: A study designed to identify or measure effects of specific exposures, interventions or risk factors. This design employs the use of an appropriate comparison group to test epidemiologic hypotheses, thus attempting to identify associations or causal relationships.

Descriptive study: A study that describes characteristics of a condition in relation to particular factors or exposure of interest. This design often provides the first important clues about possible determinants of disease and is useful for the formulation of hypotheses that can be subsequently tested using an analytic design.

Literature review: A study that analyzes critical points of a published body of knowledge. This is done through summary, classification and comparison of prior studies. With the exception of meta-analyses, which statistically re-analyze pooled data from several studies, these studies are secondary sources and do not report any new or experimental work.

Separate algorithms were developed for analytic studies, descriptive studies and literature reviews to help reviewers identify specific designs within those categories. The algorithm below, for example, helps reviewers determine which study design was used within the analytic study category (Figure 1). It is based on key decision points such as number of groups or allocation to group. The legends for the algorithms and supportive tools such as the glossary provide additional detail to further differentiate study designs, such as whether a cohort study was retrospective or prospective.

Separate critical appraisal tools were developed for analytic studies, for descriptive studies and for literature reviews, with relevant criteria in each tool. For example, a summary of the items covered in the analytic study critical appraisal tool is shown in Table 1. This tool is used to appraise trials, observational studies and laboratory-based experiments. A supportive tool for assessing statistical analysis was also provided that describes common statistical tests used in epidemiologic studies.

Table 1: Aspects appraised in analytic study critical appraisal tool

<table>
<thead>
<tr>
<th>Aspect</th>
<th>Type of assessment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample and sampling methods</td>
<td>Representativeness of participants, control of selection bias</td>
</tr>
<tr>
<td>Internal validity</td>
<td>Control of biases: misclassification, information validity and reliability of data collection instruments, adequacy of retention and follow-up</td>
</tr>
<tr>
<td>Control of confounding</td>
<td>Comparability of control and intervention groups, adequacy of control of major confounders</td>
</tr>
<tr>
<td>Ethics</td>
<td>Adequacy of ethical conduct</td>
</tr>
<tr>
<td>Analysis</td>
<td>Adequacy and interpretation of statistical testing, power and sample size</td>
</tr>
<tr>
<td>Screening and applicability questions</td>
<td>Generalizability of results, feasibility of implementation</td>
</tr>
</tbody>
</table>

*Public Health Agency of Canada. Infection Prevention and Control Guidelines Critical Appraisal Tool Kit (6)
ethical conduct. It is used to appraise cross-sectional studies, outbreak investigations, case series and case reports.

The literature review critical appraisal tool assesses the methodology, results and applicability of narrative reviews, systematic reviews and meta-analyses.

After appraisal of individual items in each type of study, each critical appraisal tool also contains instructions for drawing a conclusion about the overall quality of the evidence from a study, based on the per-item appraisal. Quality is rated as high, medium or low. While a RCT is a strong study design and a survey is a weak design, it is possible to have a poor quality RCT or a high quality survey. As a result, the quality of evidence from a study is distinguished from the strength of a study design when assessing the quality of the overall body of evidence. A definition of some terms used to evaluate evidence in the CAT is shown in Table 2.

Table 2: Definition of terms used to evaluate evidence

<table>
<thead>
<tr>
<th>Summative items assessed</th>
<th>Rating</th>
<th>Criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td>Strength of study design</td>
<td>Strong</td>
<td>Meta-analysis &gt; Randomized controlled trial (RCT) &gt; non-randomized controlled trial (NRCT) = lab experiment &gt; controlled before-after (CBA)</td>
</tr>
<tr>
<td>Note: “x &gt; y” means x is a stronger design than y</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Moderate</td>
<td></td>
<td>Cohort &gt; case-control &gt; interrupted time series with adequate data collection points &gt; cohort with non-equivalent comparison group</td>
</tr>
<tr>
<td>Weak</td>
<td></td>
<td>Uncontrolled before-after (UCBA) &gt; interrupted time series with inadequate data collection points &gt; descriptive (cross-sectional &gt; epidemiologic link &gt; ecologic or correlational)</td>
</tr>
<tr>
<td>Quality of the study</td>
<td>High</td>
<td>No major threats to internal validity (bias, chance and confounding have been adequately controlled and ruled out as an alternate explanation for the results)</td>
</tr>
<tr>
<td>Medium</td>
<td></td>
<td>Minor threats to internal validity that do not seriously interfere with ability to draw a conclusion about the estimate of effect</td>
</tr>
<tr>
<td>Low</td>
<td></td>
<td>Major threat(s) to internal validity that interfere(s) with ability to draw a conclusion about the estimate of effect</td>
</tr>
<tr>
<td>Number of studies</td>
<td>Multiple</td>
<td>Four or more studies</td>
</tr>
<tr>
<td></td>
<td>Few</td>
<td>Three or fewer studies</td>
</tr>
<tr>
<td>Consistency of results</td>
<td>Consistent</td>
<td>Studies found similar results</td>
</tr>
</tbody>
</table>

Table 2: Definition of terms used to evaluate evidence (continued)

<table>
<thead>
<tr>
<th>Consistency of results (continued)</th>
<th>Inconsistent</th>
<th>Criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td>Contradictory</td>
<td>Varying results with no clear overall trend related to the effect</td>
<td></td>
</tr>
<tr>
<td>Directness of evidence</td>
<td>Direct evidence</td>
<td>Comes from studies that specifically researched the association of interest</td>
</tr>
<tr>
<td>Extrapolation</td>
<td>Inference drawn from studies that researched a different but related key question or researched the same key question but under artificial conditions (e.g., some lab studies)</td>
<td></td>
</tr>
</tbody>
</table>

Tools for summarizing the evidence

The second phase in the critical appraisal process involves summarizing the results of the critical appraisal of individual studies. Reviewers are instructed to complete a template evidence summary table, with key details about each study and its ratings. Studies are listed in descending order of strength in the table. The table simplifies looking across all studies that make up the body of evidence informing a recommendation and allows for easy comparison of participants, sample size, methods, interventions, magnitude and consistency of results, outcome measures and individual study quality as determined by the critical appraisal. These evidence summary tables are reviewed by the working group to determine the rating for the quality of the overall body of evidence and to facilitate development of recommendations based on evidence.

Rating the quality of the overall body of evidence

The third phase in the critical appraisal process is rating the quality of the overall body of evidence. The overall rating depends on the five items summarized in Table 2: strength of study designs, quality of studies, number of studies, consistency of results and directness of the evidence. The various combinations of these factors lead to an overall rating of the strength of the body of evidence as strong, moderate or weak as summarized in Table 3.

A unique aspect of this toolkit is that recommendations are not graded but are formulated based on the graded body of evidence. Actions are either recommended or not recommended; it is the strength of the available evidence that varies, not the strength of the recommendation. The toolkit does highlight, however, the need to re-evaluate new evidence as it becomes available especially when recommendations are based on weak evidence.
Table 3: Criteria for rating evidence on which recommendations are based

<table>
<thead>
<tr>
<th>Strength of Evidence</th>
<th>Grades</th>
<th>Criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td>Strong</td>
<td>AI</td>
<td>Direct evidence from meta-analysis or multiple strong design studies of high quality, with consistency of results</td>
</tr>
<tr>
<td></td>
<td>BI</td>
<td>Direct evidence from multiple moderate design studies of high quality with consistency of results OR Extrapolation from multiple strong design studies of high quality, with consistency of results</td>
</tr>
<tr>
<td></td>
<td>CI</td>
<td>Direct evidence from any combination of strong or moderate design studies of high/medium quality, with a clear trend but some inconsistency of results OR Extrapolation from multiple strong design studies of medium quality or moderate design studies of high/medium quality, with consistency of results OR One strong design study with support from multiple weak design studies of high/medium quality, with consistency of results</td>
</tr>
<tr>
<td>Moderate</td>
<td>BII</td>
<td>Direct evidence from multiple weak design studies of high/medium quality, with consistency of results OR Extrapolation from any combination of strong/moderate design studies of high/medium quality, with inconsistency of results</td>
</tr>
<tr>
<td>Weak</td>
<td>CII</td>
<td>Studies of low quality regardless of study design OR Contradictory results regardless of study design OR Case series/case reports OR Expert opinion</td>
</tr>
</tbody>
</table>

Pilot test of the CAT

Of 34 individuals who indicated an interest in completing the pilot test, 17 completed it. Multiple peer-reviewed studies were selected representing analytic studies, descriptive studies and literature reviews. The same studies were assigned to participants with similar content expertise. Each participant was asked to appraise three analytic studies, two descriptive studies and one literature review, using the appropriate critical appraisal tool as identified by the participant. For each study appraised, one critical appraisal tool and the associated tool-specific feedback form were completed. Each participant also completed a single general feedback form. A total of 101 of 102 critical appraisals were conducted and returned, with 81 tool-specific feedback forms and 14 general feedback forms returned.

The majority of participants (>85%) found the flow of each tool was logical and the length acceptable but noted they still had difficulty identifying the study designs (Table 4).

Table 4: Pilot test feedback on user friendliness

<table>
<thead>
<tr>
<th>Items</th>
<th>Analytic Critical Appraisal Tool (%)</th>
<th>Descriptive Critical Appraisal Tool (%)</th>
<th>Literature review Critical Appraisal Tool (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Logical flow</td>
<td>n=39 of 51*</td>
<td>89.7</td>
<td>96.4</td>
</tr>
<tr>
<td>Acceptable length</td>
<td></td>
<td>97.4</td>
<td>100</td>
</tr>
<tr>
<td>Clear phrasing and explanations</td>
<td></td>
<td>72.2</td>
<td>88.5</td>
</tr>
<tr>
<td>Tool was helpful for critical appraisal process</td>
<td></td>
<td>92.3</td>
<td>85.7</td>
</tr>
</tbody>
</table>

*Number of tool-specific forms returned for total number of critical appraisals conducted

The vast majority of the feedback forms (86–93%) indicated that the different tools facilitated the critical appraisal process. In the assessment of consistency, however, only four of ten analytic studies appraised (40%), had complete agreement on the rating of overall study quality by participants, the other six studies had differences noted as mismatches. Four of the six studies with mismatches were observational studies. The differences were minor. None of the mismatches included a study that was rated as both high and low quality by different participants. Based on the comments provided by participants, most mismatches could likely have been resolved through discussion with peers. Mismatched ratings were not an issue for the descriptive studies and literature reviews. In summary, the pilot test provided useful feedback on different aspects of the toolkit. Revision were made to address the issues identified from the pilot test and thus strengthen the CAT.

Discussion

The Infection Prevention and Control Guidelines Critical Appraisal Tool Kit was developed in response to the needs of infection control professionals reviewing literature that generally did not include clinical trial evidence. The toolkit was designed to meet the identified needs for training in critical appraisal with extensive instructions and dictionaries, and tools applicable to all three types of studies (analytic studies, descriptive studies and literature reviews). The toolkit provided a method to progress from assessing individual studies to summarizing and assessing the strength of a body of evidence and assigning a grade. Recommendations are then developed based on the graded
body of evidence. This grading system has been used by the Public Health Agency of Canada in the development of recent infection prevention and control guidelines (5,7). The toolkit has also been used for conducting critical appraisal for other purposes, such as addressing a practice problem and serving as an educational tool (8,9).

The CAT has a number of strengths. It is applicable to a wide variety of study designs. The criteria that are assessed allow for a comprehensive appraisal of individual studies and facilitates critical appraisal of a body of evidence. The dictionaries provide reviewers with a common language and criteria for discussion and decision making.

The CAT also has a number of limitations. The tools do not address all study designs (e.g., modelling studies) and the toolkit provides limited information on types of bias. Like the majority of critical appraisal tools (10,11), these tools have not been tested for validity and reliability. Nonetheless, the criteria assessed are those indicated as important in textbooks and in the literature (12,13). The grading scale used in this toolkit does not allow for comparison of evidence grading across organizations or internationally, but most reviewers do not need such comparability. It is more important that strong evidence be rated higher than weak evidence, and that reviewers provide rationales for their conclusions; the toolkit enables them to do so.

Overall, the pilot test reinforced that the CAT can help with critical appraisal training and can increase comfort levels for those with limited experience. Further evaluation of the toolkit could assess the effectiveness of revisions made and test its validity and reliability.

A frequent question regarding this toolkit is how it differs from GRADE as both distinguish stronger evidence from weaker evidence and use similar concepts and terminology. The main differences between GRADE and the CAT are presented in Table 5. Key differences include the focus of the CAT on rating the quality of individual studies, and the detailed instructions and supporting tools that assist those with limited experience in critical appraisal. When clinical trials and well controlled intervention studies are or become available, GRADE and related tools from Cochrane would be more appropriate (2,3). When descriptive studies are all that is available, the CAT is very useful.

Table 5: Features of the Critical Appraisal Toolkit (CAT) and GRADE (continued)

<table>
<thead>
<tr>
<th>Feature</th>
<th>CAT</th>
<th>GRADE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Assessment of individual studies</td>
<td>Tools are provided for the critical appraisal of individual studies and a quality rating given per study.</td>
<td>Each study is individually assessed, but no quality rating is provided per study.</td>
</tr>
<tr>
<td>Assessment of body of evidence</td>
<td>Overall body of evidence is graded based on criteria provided.</td>
<td>Overall body of evidence is graded on criteria provided.</td>
</tr>
<tr>
<td>Scoring and criteria</td>
<td>A qualitative assessment is made based on strength of study designs, the quality of studies, number of studies, consistency of results, and directness of the evidence. A grade is assigned based on the assessment.</td>
<td>A numeric score is calculated based on whether the evidence is randomized or non-randomized, risk of bias, inconsistency, indirectness, imprecision and publication bias. The score is translated to a grade.</td>
</tr>
<tr>
<td>Grade of evidence</td>
<td>Evidence is graded as strong, moderate or weak quality.</td>
<td>Evidence is graded as high, moderate, low or very low certainty.</td>
</tr>
<tr>
<td>Grade of recommendations</td>
<td>Recommendations are not graded, actions are either recommended or not.</td>
<td>Recommendations are graded as strong or weak/conditional.</td>
</tr>
<tr>
<td>Guidance for reviewers</td>
<td>Detailed criteria and explanations for use are provided in a single toolkit.</td>
<td>Detailed criteria and instructions provided in multiple documents and training available.</td>
</tr>
</tbody>
</table>

Abbreviation: GRADE, Grading of Recommendations Assessment, Development and Evaluation

### Conclusion

The Infection Prevention and Control Guidelines Critical Appraisal Tool Kit was developed in response to needs for training in critical appraisal, assessing evidence from a wide variety of research designs, and a method for going from assessing individual studies to characterizing the strength of a body of evidence. Clinician researchers, policy makers and students can use these tools for critical appraisal of studies whether they are trying to develop policies, find a potential solution to a practice problem or critique an article for a journal club. The toolkit adds to the arsenal of critical appraisal tools currently available and is especially useful in assessing evidence from a wide variety of research designs.
Author's Statement

DM – Conceptualization, methodology, investigation, data collection and curation and writing – original draft, review and editing

TO – Conceptualization, methodology, investigation, data collection and curation and writing – original draft, review and editing

KD – Conceptualization, review and editing, supervision and project administration

Conflict of interest
None.

Contributor
Jennifer Kruse, Public Health Agency of Canada – Conceptualization and Project administration

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