

Appraisal of clinical practice guidelines for the management of attention deficit hyperactivity disorder (ADHD) using the AGREE II Instrument: a systematic review

Turki Albatti , Saleh Al Salehi, Fahad Bashiri, Muddathir Hamad, Haya Al-Joudi, Hadeel Daghash, Jeremy Varnham, Yasser Amer

REVIEW TITLE AND BASIC DETAILS

Review title

Appraisal of clinical practice guidelines for the management of attention deficit hyperactivity disorder (ADHD) using the AGREE II Instrument: a systematic review

Review objectives

The aim is to explore the quality of and critically appraise recently published evidence-based clinical practice guidelines for the management of Attention Deficit Hyperactivity Disorder (ADHD)

Keywords

ADHD, AGREE II instrument, Attention Deficit Hyperactivity Disorder, clinical practice guidelines, evidence, neurology, pediatrics, psychiatry, psychology, quality assessment

SEARCHING AND SCREENING

Searches

Data Sources include:-

- Guidelines International Network <http://www.g-i-n.net/library/international-guidelines-library>
- National Guidelines Clearinghouse <http://www.guidelines.gov>
- EBSCO - DynaMed Plus <https://dynamed.ebscohost.com/>
- National Institute for Health and Clinical Excellence <http://www.nice.org.uk/>
- Scottish Intercollegiate Guidelines Network <http://www.sign.ac.uk/index.html>
- US National Library of Medicine, National Institutes of Health (MEDLINE/ PubMed) <http://www.ncbi.nlm.nih.gov/PubMed> (for CPGs published as full-text articles)

- Google Scholar (free) <http://scholar.google.com/> (for CPGs published as full-text articles) • Plus CPGs published by national and international specialized scientific societies/ associations relevant to ADHD (e.g. American Academy of Pediatrics, American Psychiatric Association, European Psychiatric Association, etc.).

Additionally, we will also conduct internet searches for CPGs published online only.

Study design

Clinical Practice Guidelines

ELIGIBILITY CRITERIA

Condition or domain being studied

ADHD in children and adolescents. The inclusion or exclusion of adults will have been at the discretion of the Source CPG developers and age limits, co-morbidities, etc, should have been defined by them. We will assess how this was reported as part of our appraisal using the AGREE II Instrument.

Population

The participants, including inclusion and exclusion criteria, should have been specified by the guideline developers and cannot be known until we begin the appraisal of the guidelines identified by our searches. We will include only CPGs with a section targeting children and adolescents. Relevant items of the AGREE II Instrument will be used to assess how well the scope of the CPG was reported.

Intervention(s) or exposure(s)

The interventions included within the included CPGs will have been pre-defined by the CPG development group to match the individual requirements within the scope. We will only consider CPGs that include the comprehensive management of ADHD. The criteria of the AGREE II Instrument will be used to assess how well the included interventions were reported and presented. Identified key target interventions include diagnosis and assessment (i.e. parent, carer, teacher, or patient complaints, history and physical examination, psychological tools, differential diagnosis, and investigations) and treatment (i.e. psychosocial interventions, pharmacological treatment, comorbidities, parents/ carers and home, monitoring, special cases, transition of care from childhood to adulthood, complementary medicine, and treatment of complications of pharmacological treatment)

Comparator(s) or control(s)

Comparator(s) and control groups with inclusion and exclusion criteria may or may not have been pre-defined by the CPG development group. Comparators would not usually be defined when multiple interventions are included in the recommendations of the CPG.

Context

Primary, Secondary and Tertiary healthcare facilities that provide care for patients with ADHD
In addition to School-based and Home-based healthcare services.

OUTCOMES TO BE ANALYSED

Main outcomes 1 change

(1) To standardize evidence-based assessment and management of ADHD including Early diagnosis and intervention of ADHD and Multi-modal interventions for ADHD.

(2) To improve patient target outcomes and safety (e.g. symptoms of ADHD, academic performance, social relationships, parent-child interactions and family stress, and risk for serious accidental injury).

Measures of effect

Not applicable.

Additional outcomes 1 change

(1) To decrease variation of practice.

(2) To improve patient-related adverse health outcomes that co-occur with ADHD (e.g. substance abuse disorder and smoking, sleep difficulties/disorders, physical injuries, etc.).

Measures of effect

Not applicable.

DATA COLLECTION PROCESS

Data extraction (selection and coding)

Five reviewers including one CPG methodologist will independently screen the titles and abstracts of all searched documents and determine the ones for full-text review and eligible CPGs for ADHD. Disagreements will be resolved through discussion and consensus with the reviewers. The same five reviewers will search the internet for CPGs published online only. Information relevant to rating of each of the 23 items on the AGREE II Instrument will be extracted from included CPGs by utilizing the online tool (MY AGREE Plus) that is freely available and accessible from the AGREE Enterprise website (<http://www.agreetrust.org/>). The authors will then independently score every item on a 1-7 scale based on how well each CPG addresses the listed questions and the AGREE criteria. An additional validation of the included CPGs will be conducted for inclusion of systematic reviews in the evidence-based with their category whether Cochrane systematic reviews or others.

Risk of bias (quality) assessment

The AGREE II Instrument addresses the reporting of CPGs. It does not assess aspects of study design quality such as appropriateness of the methods used or trustworthiness of the recommendations. We are not aware of any validated instrument for assessing risk of bias of CPGs. Some of the items on the AGREE II checklist are relevant to the detection of potential sources of bias (e.g. bias due to under-representation of some stakeholders in the development process - items 4 and 5, reporting bias - items 9-12, bias due to conflict of interest/independence of funding bodies - items 22 and 23).

PLANNED DATA SYNTHESIS

Strategy for data synthesis

Descriptive statistics will be generated and presented in tabular format to summarize the characteristics of the CPGs eligible for inclusion. Once all seven AGREE appraisers have scored each item on the AGREE II, they will hold a meeting to discuss discrepancies between scores and address any obvious discrepancies and errors. For every included CPG, we will use the 'My AGREE PLUS' (<http://www.agreetrust.org/my-agree/>) to record scores and to keep online records of individual appraiser's scores. For every single CPG, a standardized quality score will be calculated for each of the six domains of the AGREE II Instrument, using the equation presented in the AGREE II User's Manual. In summary, standardized domain scores will be calculated by summing up all the appraisers' scores of the individual items in a domain and by scaling the total as a percentage of the maximum possible score for that domain. This calculation is automatically generated on the My AGREE PLUS platform. The overall quality of the included CPGs in each domain of the AGREE II will be presented through the two overall assessments included in the instrument. Since the six domains of the AGREE II are independent, standardized domain quality scores will not be aggregated.

Analysis of subgroups or subsets

The references section of included CPGs will be reviewed and assessed for inclusion of 'Systematic Reviews' in the evidence-base of these CPGs (especially the Cochrane systematic reviews).

REVIEW AFFILIATION, FUNDING AND PEER REVIEW

Review team members

- Dr Turki Albatti , Department of Psychiatry, King Saud University Medical City, King Saud University, Riyadh, Saudi Arabia
- Dr Saleh Al Salehi, Child Development Center, King Abdullah Bin Abdulaziz University Hospital, Princess Noura bint Abdulrahman University, Riyadh, Saudi Arabia
- Dr Fahad Bashiri, Division of Neurology, Department of Pediatrics, College of Medicine, King Saud University Medical City, King Saud University, Riyadh, Saudi Arabia
- Dr Muddathir Hamad, Division of Neurology, Department of Pediatrics, College of Medicine, King Saud University Medical City, King Saud University, Riyadh, Saudi Arabia
- Dr Haya Al-Joudi, Department of Neurosciences, King Faisal Specialist Hospital and Research Center, Riyadh, Saudi Arabia
- Dr Hadeel Daghash, Ada'a Program, Assistant Deputyship for Hospital Services, Ministry of Health, Riyadh
- Mr Jeremy Varnham, Saudi ADHD Society, Riyadh, Saudi Arabia and School of Psychology, University of East London, UK
- Dr Yasser Amer, Quality Management Department and Research Chair for Evidence-Based Health Care and Knowledge Translation, King Saud University Medical City, King Saud University, Riyadh, Saudi Arabia - Alexandria Center for Evidence-Based Clinical Practice Guidelines, Faculty of Medicine, Healthcare Quality Directorate, Alexandria University Hospitals, Alexandria University, Alexandria, Egypt

Review affiliation

The Saudi ADHD Society, Riyadh, Saudi Arabia

Funding source

The unified ADHD Clinical Practice Guidelines Project is the strategic project 7.2 of the Saudi ADHD Society for the period 2017-2019. The Saudi ADHD Society is a registered non-profit under licence 474 from the Saudi Ministry of Labor and Social Development, and the project received the Ministry approval (No. 52476) on 5/8/1438. The project is entirely funded by the Saudi ADHD Society. No funding received from any pharmaceutical or industrial company

Named contact

Dr. Yasser Sami Amer. Dr. Yasser Sami Amer, Quality Management Department, King Khalid University Hospital, Research Chair for Evidence-Based Health Care and Knowledge Translation, King Saud University Medical City, King Saud University, P.O. Box 7805, Riyadh 11472, Kingdom of Saudi Arabia yamer@ksu.edu.sa; yassersamiamer@gmail.com yasser3amer@yahoo.com

TIMELINE OF THE REVIEW

Review timeline

Start date: 04 March 2017. End date: 30 November 2017

Date of first submission to PROSPERO

12 September 2018

Date of registration in PROSPERO

24 November 2017

CURRENT REVIEW STAGE

Publication of review results ^{1 change}

The intention is to publish the review once completed. The review will be published in English

Amer YS, Al-Joudi HF, Varnham JL, Bashiri FA, Hamad MH, Al Salehi SM, et al. (2019) Appraisal of clinical practice guidelines for the management of attention deficit hyperactivity disorder (ADHD) using the AGREE II Instrument: A systematic review. PLoS ONE 14(7): e0219239. <https://doi.org/10.1371/journal.pone.0219239>

<https://journals.plos.org/plosone/article?id=10.1371/journal.pone.0219239#sec006>

Stage of the review at this submission

Review stage	Started	Completed
Pilot work	✓	✓
Formal searching/study identification	✓	✓
Screening search results against inclusion criteria	✓	✓
Data extraction or receipt of IP	✓	✓
Risk of bias/quality assessment	✓	✓
Data synthesis	✓	✓

Review status

The review is completed.

ADDITIONAL INFORMATION

Additional information 1 change

This review is being undertaken as part of the planning for the CPG adaptation project utilizing the ADAPTE methodology for the Unified CPG for ADHD Project of the Saudi ADHD Society.

PROSPERO version history

- Version 1.2 published on 22 Oct 2019
- Version 1.1 published on 24 Oct 2018
- Version 1.0 published on 05 Oct 2017

Review conflict of interest

None known

Country

Saudi Arabia

Medical Subject Headings

Attention Deficit Disorder with Hyperactivity; Humans

Disclaimer

The content of this record displays the information provided by the review team. PROSPERO does not peer review registration records or endorse their content.

PROSPERO accepts and posts the information provided in good faith; responsibility for record content rests with the review team. The owner of this record has affirmed that the information provided is truthful and that they understand that deliberate provision of inaccurate information may be construed as scientific misconduct.

PROSPERO does not accept any liability for the content provided in this record or for its use. Readers use the information provided in this record at their own risk.

Any enquiries about the record should be referred to the named review contact