

# Systematic review of clinical practice guidelines for uveitis

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## ABSTRACT

To facilitate the integration of eye care into universal health coverage, the WHO is developing a Package of Eye Care Interventions (PECI). Development of the PECI involves the identification of evidence-based interventions from relevant clinical practice guidelines (CPGs) for uveitis. A systematic review of CPGs published on uveitis between 2010 and March 2020 was conducted. CPGs passing title and abstract and full-text screening were evaluated using the Appraisal of Guidelines for Research and Evaluation II (AGREE II) tool and data on recommended interventions extracted using a standard data extraction sheet. Of 56 CPGs identified as potentially relevant from the systematic literature search, 3 CPGs underwent data extraction following the screening stages and appraisal with the AGREE II tool. These CPGs covered screening for, monitoring and treating juvenile idiopathic arthritis (JIA)-associated uveitis, the use of adalimumab and dexamethasone in treating non-infectious uveitis, and a top-level summary of assessment, differential diagnosis and referral recommendations for uveitis, aimed at primary care practitioners. Many of the recommendations were based on expert opinion, though some incorporated clinical study and randomised controlled trial data.

There is currently sparse coverage of the spectrum of disease caused by uveitis within CPGs. This may partially be due to the large number of conditions with diverse causes and clinical presentations covered by the umbrella term uveitis, which makes numerous sets of guidelines necessary. The limited pool of CPGs to select from has implications for clinicians seeking guidance on clinical care strategies for uveitis.

## INTRODUCTION

A key recommendation of the WHO's 2019 World report on vision<sup>1</sup> was to embed eye care within universal health coverage (UHC) in order to address inequities in access to eye care services and contribute to Sustainable Development Goals.<sup>2</sup> To enable countries to plan, budget and integrate eye care into UHC, WHO is developing a priority package of eye care interventions (PECI), in collaboration with Cochrane Eyes and Vision (CEV).<sup>3</sup> A crucial step in the development of the PECI is the systematic identification of evidence-based eye care interventions for selected priority eye conditions using high-quality clinical practice guidelines (CPGs) and, where

needed, systematic reviews. Uveitis has been identified as 1 of 15 priority eye conditions for inclusion in the PECI.

Uveitis is a common, sight-threatening group of disorders, all of which are characterised by inflammation of the uveal tract of the eye (iris, ciliary body and choroid). Population-based studies have suggested an overall incidence of 17–52 people in 100 000 each year, and that uveitis causes 9%–15% of blindness in the Western World.<sup>4</sup> Classifying this heterogeneous disease spectrum, which spans numerous infectious and autoimmune diseases, has been a unique challenge for ophthalmic clinicians. The standardisation of uveitis nomenclature (SUN) group sought to standardise the descriptions of the diseases and syndromes,<sup>5</sup> mapping at least 28 uveitis diseases in 2005. This variability in disease features, both across and within distinct disease groups, has impacted clinical trials in uveitis.

Nevertheless, interventions for uveitis are continually emerging. These have included medications applied to the eye such as topical steroid<sup>6</sup> and non-steroidal eye-drops, local injection treatments to the eye and intraocular steroid implants,<sup>7</sup> systemic immunosuppressants<sup>8</sup> and immunomodulatory therapies which include newer biologic treatments.<sup>9</sup> The purpose of this paper is to present the results, including the quality and current state of evidence, of a systematic review of CPGs for uveitis.

## METHODS

This systematic review of CPGs was conducted in compliance with the methodology presented in the introductory PECI paper.<sup>3</sup> Exclusion criteria for each stage of screening below are provided in [table 1](#). The stages in the review are as follows:

### Systematic literature search

A single, systematic literature search of selected academic (MEDLINE, Embase, CINAHL, Global Health, Global Index Medicus) and guideline databases (online

**Table 1** Exclusion criteria for screening of clinical practice guidelines (CPGs)

Title and abstract screening	Full-text screening	Quality appraisal
<ol style="list-style-type: none"> <li>1. The identified literature was not a CPG</li> <li>2. The CPG was not published in the last 10 years</li> <li>3. The CPG was not in English</li> <li>4. The CPG was not developed for the selected eye conditions</li> </ol>	<ol style="list-style-type: none"> <li>1. There was commercial funding or unmanaged conflicts of interest present</li> <li>2. Absence of affiliation of authors</li> </ol>	<ol style="list-style-type: none"> <li>1. The average score of the two researchers for items 4, 7, 8, 12 or 22 is below 3</li> <li>2. The sum of the average score of the two researchers for all nine items is less than 45</li> </ol>

supplemental Appendix 1) was carried out by a CEV information specialist in March 2020. MeSH terms were used where applicable. In addition, the websites of professional ophthalmology and optometry associations' were searched for relevant guidelines (online supplemental Appendix 1). All searches were limited to the last 10 years, to ensure evidence was current, and to English language, to ensure timely completion of the PECL. The search strategy for academic databases can be found in online supplemental Appendix 2.

### Title and abstract screening

Two members of the technical working group independently screened the titles and abstracts of articles identified from the systematic literature searches. Title and abstracts were presented, and responses tracked, using Abstrackr, a semiautomated online citation screening programme.<sup>10</sup> Citations were excluded if they were not in the English language, not a CPG or were not relevant to a priority eye condition. A secondary title and abstract screening was then conducted to identify CPGs potentially relevant to uveitis. All disagreements were resolved by discussion between a CEV representative and the WHO.

### Full-text screening

Following title and abstract screening, independent full-text screening of the CPGs relevant to uveitis was conducted by two authors (NG and IRR). CPGs without author affiliations listed, or with undisclosed or unmanaged conflicts of interest, were excluded. Discrepancies were resolved through discussion between the two authors or, in the event a consensus couldn't be reached, by discussion with a third author.

### Quality appraisal

The same two authors who conducted full-text screening went on to independently evaluate the quality of CPGs using the 'Appraisal of Guidelines for Research and Evaluation' (AGREE II) tool.<sup>11</sup> Nine items (4, 7, 8, 10, 12, 13, 15, 22 and 23) were specifically used for the appraisal of CPGs (online supplemental Appendix 3). These items were selected based on a consensus finding process among three researchers<sup>12</sup> and in consultation with WHO Guideline Review Committee Secretariat. If the rating of an item differed by more than two points between the two researchers, the results were discussed

between the two authors, and a representative of WHO or CEV, if necessary, to reach consensus.

Following evaluation with the AGREE II tool, guidelines were excluded if (1) the average score of the two researchers for items 4, 7, 8, 12 or 22 was below 3 or (2) the sum of the average score of the two researchers for all nine items is less than 45.

### Data extraction

Data on recommendations were extracted from selected CPGs by one author using a standardised form that captured information on the recommendation (type of recommendation, dosage, target group, etc), the strength of recommendation and the quality of the evidence used to inform the recommendation. The data extraction was then independently checked by a second author. In the event of disagreement, a third author was involved and agreement reached by discussion. The process was repeated for all the CPGs until agreement on the recommended eye care interventions was reached.

With respect to the published protocol, no changes were made. The quality check and the methodological support for this study have been provided by the WHO and CEV.

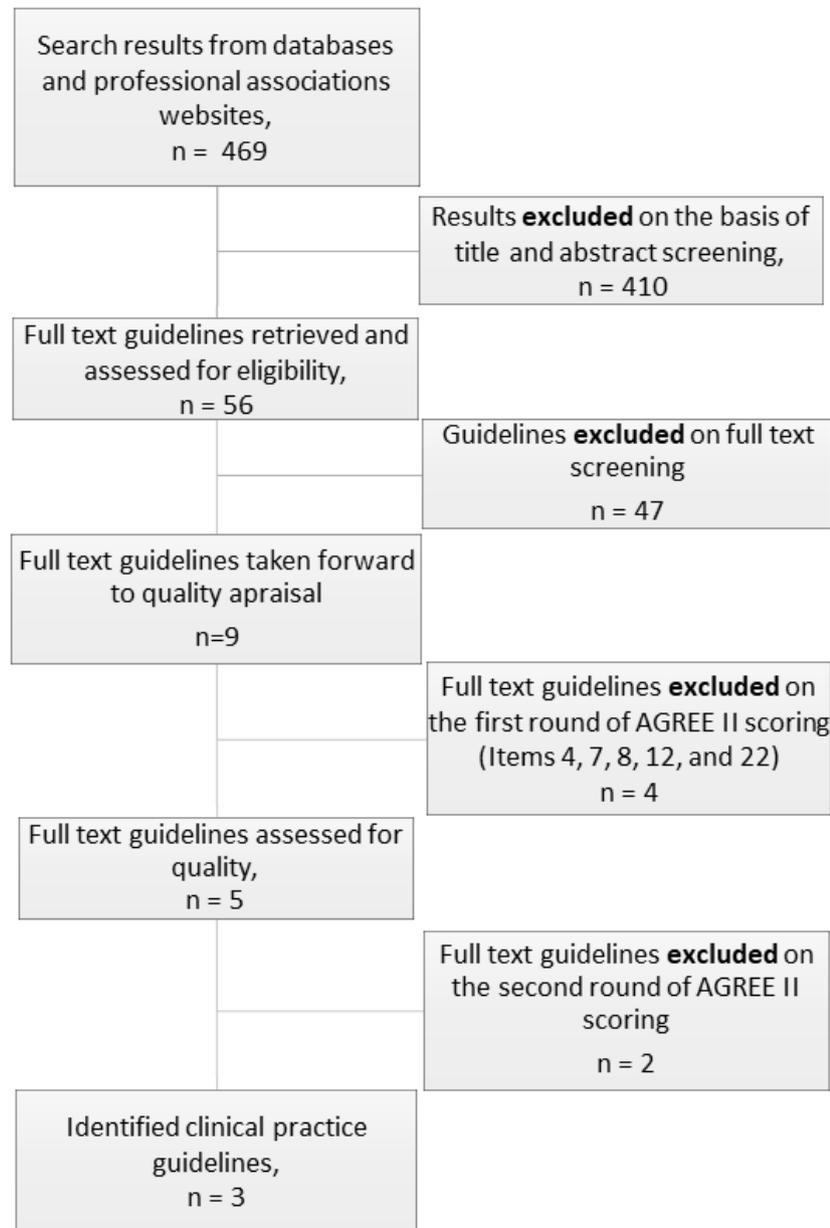
## RESULTS

### Title and abstract screening

The results of the selection process are summarised in figure 1. After combining all searches from academic and guideline databases and professional association webpages, 469 CPGs were identified and underwent independent title and abstract screening. Fifty-six of these were judged to be potentially relevant to uveitis after title and abstract screening, and went on to full-text screening.

### Full-text screening and quality appraisal

Of the 56 potentially CPGs with potential relevance to uveitis, 47 CPGs were excluded. Forty were judged not relevant to uveitis. In a further two cases, exclusion was based on industry funding of the guideline, or because the presence or absence of possible conflicts of interest were not stated, as well as the author affiliations not being stated.<sup>13 14</sup> A further four were excluded based on the potential conflict of interest criteria alone,<sup>15–18</sup> and a final one because the author affiliations were not listed.<sup>19</sup>



**Figure 1** Results of the screening process (diagram).AGREE II, Appraisal of Guidelines for Research and Evaluation II.

A total of nine CPGs were considered potentially relevant and passed to the quality appraisal phase. Of these nine, there were no concerns over conflicts of interest, and author affiliations were unambiguous. Following quality appraisal with the AGREE II tool, six guidelines were excluded for the following reasons (figure 1): four because the average score of the two researchers for items 4, 7, 8, 12 or 22 was less than 3,<sup>20–23</sup> and two because the sum of the average score of the two researchers for all nine items was less than 45<sup>24 25</sup> (table 2).

The CPGs identified for data extraction were as follows<sup>26–28</sup>:

- ▶ Adalimumab and dexamethasone for treating non-infectious uveitis, National Institute for Health and Care Excellence (NICE) (UK) Technology appraisal guidance, July 2017.

- ▶ 2019 American College of Rheumatology/Arthritis Foundation Guideline for the Screening, Monitoring and Treatment of Juvenile Idiopathic Arthritis-Associated Uveitis.<sup>21</sup>

- ▶ Uveitis Clinical Knowledge Summary, NICE (UK), November 2019.

The AGREE II ratings for the three selected CPGs are presented in table 2.

#### Data extraction

An overall view of the strength of recommendation and quality of evidence given in each selected guideline is reported in table 3. Specific recommendations extracted from each CPG are given in online supplemental table, along with the strength of the recommendation as

**Table 2** Guidelines found and selected, and their score in respect of the criteria used to reach the final choice

Guideline	AGREE II ratings											Comprehensiveness (applicability to different settings)
	Average of key items											
	4	7	8	12	22	4, 7, 8, 10, 12, 13, 15, 22, 23	Topic	Publication Date				
NICE — adalimumab <sup>26</sup>	4	5	3.5	6	4	45.5	Posterior uveitis	2017	Low			
ACR — JIA uveitis <sup>27</sup>	4	6.5	5	3	4	45	JIA uveitis	2019	High			
NICE — uveitis CKS <sup>28</sup>	3.5	7	5.5	6	4	46.5	Anterior uveitis	2019	High			
Cincinnati Children's Hospital Medical Centre. BEST Screening for Uveitis in Children with JIA <sup>20</sup>	3	6	2	5.5	3.5	-	JIA uveitis	2021				
Childhood Arthritis and Rheumatology Research Alliance Consensus Treatment Plans for JIA-Associated and Idiopathic Chronic Anterior Uveitis <sup>21</sup>	3	5	2	1.5	3.5	-	JIA uveitis	2019				
Evidence-based, interdisciplinary guidelines for anti-inflammatory treatment of uveitis associated with JIA. <sup>22</sup>	4	5	2	4.5	4	-	JIA uveitis	2012				
College of Optometrists Clinical Management Guidelines Uveitis (Anterior) <sup>23</sup>	3	1.5	1	2	3	-	Anterior uveitis	2020				
Document of treatment recommendations for non-infectious anterior uveitis. <sup>24</sup>	3	3.5	5.5	5.5	4	43	Anterior uveitis	2017				
NICE — fluciclonolone acetamide intravitreal implant for treating recurrent non-infectious uveitis-guidance (TA590). <sup>25</sup>	4	5	3	6	4	44.5	Non-infectious uveitis	2019				

ACR, American College of Rheumatology; AGREE II, Appraisal of Guidelines for Research and Evaluation II; CKS, clinical knowledge summary; JIA, Juvenile Idiopathic Arthritis; NICE, National Institute for Health and Care Excellence.

assessed by the guideline writers, and the type of evidence used to make the recommendation.

The three selected guidelines covered various areas of clinical practice and types of uveitis. The 2019 American College of Rheumatology/Arthritis Foundation Guideline for the Screening, Monitoring and Treatment of JIA-Associated Uveitis<sup>27</sup> covered a specific type of uveitis which is associated with JIA. The document included recommendations for screening, monitoring and treatment of the condition as well as patient education, which were drawn up by an expert panel using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) methodology. General recommendations of note in this guideline centre on tight monitoring of children with uveitis; the authors suggest every 3 months for stable, treated uveitis and monthly for patients tapering treatment. Specific treatment recommendations include using prednisolone acetate 1% over alternative topical preparations, and using topical corticosteroids for short-term control of inflammation. Should systemic therapy be required, then subcutaneous methotrexate is recommended over oral methotrexate in the first instance. The large majority of the recommendations within this guideline are based on either the expert opinion of the panel or on clinical studies. The only randomised controlled trial data available to support a recommendation is on the use of antitumour necrosis factor (TNF) biological therapies, where the use of a monoclonal antibody biologic is advised over etanercept, which works by acting as a decoy receptor.

Adalimumab and Dexamethasone for Treating Non-Infectious Uveitis<sup>26</sup> is a guideline produced by the UK NICE, which is most concerned with providing guidance for medical practitioners working within the UK National Health Service. It specifically deals with the use of the biological treatment adalimumab, and with dexamethasone intravitreal implants, in cases of posterior uveitis. Within this narrow scope, its recommendations are clear and concise and were decided on by an appraisal committee, which heard evidence from clinical experts and received evidence from pharmaceutical companies. Included in this were results from randomised controlled trials.<sup>29–31</sup> The guidance advised the use of adalimumab in cases of non-infectious posterior uveitis unresponsive to corticosteroid therapy, and dexamethasone intravitreal implant as an option if there is active posterior uveitis causing worsening vision with a risk of blindness. Expert opinion also advised that adalimumab therapy be halted if new choroidal or retinal inflammatory lesions developed, or if there was an increase in intraocular inflammation or a decrease in visual acuity by 3 or more lines or 15 letters while on treatment.

The Clinical Knowledge Summary on Uveitis<sup>28</sup> produced by the NICE provides general recommendations on uveitis covering clinical assessment, differential diagnosis and referral. It is explicitly meant to support practitioners in primary care, and recommendations

**Table 3** Strength of recommendation and quality of the evidence of the retrieved CPGs

Guideline	Body of evidence			Strength of recommendation		
	RCTs, systematic reviews or meta-analyses*	Clinical studies	Expert opinion	Strong*	Intermediate	Weak
NICE—adalimumab <sup>26</sup>	Yes	Yes	Yes	X		
ACR—JIA uveitis <sup>27</sup>	Yes	Yes	Yes	X		
NICE—uveitis CKS <sup>28</sup>	No	No	Yes			X

Since the reference scales adopted by each guideline are not directly comparable, we propose here the recommendations according to two 3-point Likert scales, as assessed by two authors (NG and IRR).

\*At least one RCT or one systematic review are required to classify in this column.

ACR, American College of Rheumatology; CKS, clinical knowledge summary; CPGs, clinical practice guidelines; NICE, National Institute for Health and Care Excellence; RCT, randomised controlled trial.

were largely drawn from review articles and medical textbooks.

## DISCUSSION

The CPGs available to help inform the WHO Peci for uveitis formed a relatively small pool. Of 496 CPGs identified by literature search, and 56 passing title and abstract screening, the large majority were not relevant to uveitis on full-text screening. Of those identified to be of relevance to uveitis, only three guidelines were of sufficient quality (based on key AGREE II item scores) to merit inclusion. This highlights an important and concerning deficiency in high-quality CPGs in the field of uveitis and may reflect the heterogeneity of the uveitis disease spectrum and the fact that most individual uveitides are relatively uncommon. Some guidelines<sup>15–18</sup> were excluded at the screening stage owing to possible conflicts of interest as they were funded by industry partners. Industry funding of research is a common and necessary practice, but, given the intended large reach of the Peci particularly among developing nations, it was crucial to avoid any actual or potential commercial bias towards particular interventions. At the point of AGREE II scoring, the most common item for a guideline to fall short on was item 8—‘The criteria for selecting the evidence are clearly described’. Four guidelines had an average score of less than three on this point, which excluded them from inclusion (table 2). This highlights the importance of clarity on how clinical guidelines select the evidence which informs their guidance.

The diverse spectrum of disease under the umbrella of uveitis means that a single, or even a handful, of guidelines cannot comprehensively cover all disease subtypes. Although the most recent SUN working group continue to systematise the approach to disease diagnosis, classification and outcomes,<sup>26</sup> there remain significant gaps in terms of guidelines for investigation and treatment of uveitis, particularly in adult patients. Paediatric uveitis is commonly associated with JIA and this spectrum of disease is well covered by one guideline.<sup>27</sup> Cases of paediatric uveitis occurring without a background of JIA are, however, not covered.

Owing to the relative paucity of guidelines for uveitis, there can be little cross-checking between guidelines from different sources to assess concordance in

recommendations. However, one consistent theme of two of the three included guidelines is that treatment with biologic anti-TNF agents (eg, adalimumab) is appropriate in cases with a high risk of ‘blindness’<sup>26</sup> or ‘sight-threatening complications’.<sup>27</sup> This guidance is largely based on expert opinion, though there are randomised controlled trials comparing Adalimumab to placebo in adult patients (VISUAL I and VISUAL II trials),<sup>30 31</sup> which showed beneficial results. These trials did not, however, compare adalimumab to any conventional immunosuppressive therapy. The SYCAMORE trial<sup>32</sup> did provide this comparison for paediatric uveitis associated with JIA, demonstrating a significantly lower treatment failure rate in the methotrexate plus adalimumab group compared with the methotrexate plus placebo group. This is reflected in the JIA-associated uveitis guideline chosen for inclusion here, which recommends starting methotrexate and a monoclonal antibody TNF inhibitor immediately over methotrexate as monotherapy for severe active uveitis with sight threatening complications.<sup>27</sup>

Owing to the relative rarity of many forms of uveitis, a key challenge for the development of high-quality guidelines on the treatment of these conditions is to generate the primary research on which guidelines are built. This requires high-quality randomised controlled trials, which often need to be conducted across several countries in order to meet recruitment targets. Preceding any trials of therapeutics, consensus must be reached on the definition of the condition being treated. The recent SUN criteria are a vital development in this area.<sup>33</sup> Well-defined endpoints for trials are also needed, and standardisation of these would make cross-comparisons of new therapeutics being tested more straightforward. There is clear scope for improvement in the availability of clinical guidelines on uveitis diagnosis and management and well as the number of uveitides which are covered by guidelines. It is also apparent that separate guidelines may be appropriate for different care settings. Many patients may present first to a primary care setting, where the practitioner may have very limited training in recognising and treating eye disease. It is necessary therefore to be aware of the target audience for guidelines.

Some weaknesses remain in the design of this systematic review; the initial literature search was restricted

to English language guidelines and this does create an anglocentric bias to the selected guidelines. However, this search method does keep this review in line with others completed to inform the Peci. A further limitation is that the initial search was focused on specific eye-related guidelines, which may have missed some guidelines on uveitis related to systemic inflammatory diseases (eg, the spondyloarthropathies) or infectious diseases (eg, tuberculosis and syphilis), though the search strategy did catch guidelines for uveitis associated with JIA.

## CONCLUSION

Uveitis is an important clinical problem, with high prevalence worldwide,<sup>34</sup> and with high visual morbidity.<sup>35</sup> It is an umbrella term for a range of clinical entities, and the range of different aetiologies for uveitis, including infectious, autoimmune and autoinflammatory, traumatic and ischaemic causes, means that single overarching guidelines are difficult to produce. Guidance exists on the use of newer treatments such as the monoclonal antibody anti-TNF treatment adalimumab and slow release dexamethasone intravitreal implants in non-infectious posterior uveitis,<sup>26</sup> but this covers only a small proportion of cases of uveitis, and anti-TNF treatment is only recommended for cases refractory to first line (corticosteroid) and second line (conventional immunosuppressive) therapies. However, no guidance exists on when to initiate these first- and second-line treatments. There is also a marked lack of guidance on the more common presentation of anterior uveitis; and on infectious uveitis, which can be devastating to vision.<sup>36</sup>

Designing specific guidelines for each subtype of uveitis is likely impractical, as is a single CPG to cover all uveitis. However, an achievable way of increasing coverage of uveitis by CPGs, to the benefit of patients worldwide, may be to aim for guidance on uveitides based on aetiology and/or anatomical location. Broad guidelines could cover diagnosis, treatment strategies and monitoring of several uveitis subtypes under this arrangement.

Regrettably, evidence derived from CPGs for uveitis to support the development of the WHO Peci form a small pool, and cover only a minority of the clinical presentations of uveitis in sufficient detail to help clinical decision making. Thus, there will be a greater reliance on Cochrane Systematic Reviews and expert opinion in deciding on interventions to include in the Peci. There is an urgent requirement for high-quality, evidence-based guidance on a wider range of uveitides. The COVID-19 pandemic has precipitated a necessary collaboration between clinicians around the world, including uveitis specialists, due to the need for shared experience during a period of uncertainty. This era of collaboration may accelerate larger and higher-quality studies and reviews, which would support measures such as the Peci in the future.

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  - 33 Standardization of Uveitis Nomenclature (SUN) Working Group. Development of classification criteria for the Uveitides. *Am J Ophthalmol* 2021;228:96–105.
  - 34 Joltikov KA, Lobo-Chan A-M. Epidemiology and risk factors in non-infectious uveitis: a systematic review. *Front Med* 2021;8:695904.
  - 35 Durrani OMet al. Degree, duration, and causes of visual loss in uveitis. *Br J Ophthalmol* 2004;88:1159–62.
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## Appendices

### Appendix 1

#### **Guideline databases**

1. Guidelines International Network
2. National Institute for Clinical Excellence ([UK](#))
3. eGuidelines
4. Guideline Central
5. ECRI Guidelines Trust Database
6. Australian National Health and Medical Research Council clinical practice guidelines
7. Canadian Medical Association Infobase of Clinical Practice Guidelines
8. Trip Database
9. WHO guidelines
10. US Preventive Services Task Force Guideline
11. Scottish Intercollegiate Guidelines Network (SIGN)

#### **Professional association websites**

1. College of Optometrists (UK)
2. Pan American Association of Ophthalmology
3. Eastern Mediterranean Council of Optometry
4. European Society of Ophthalmology
5. World Council of Optometry
6. American Academy of Ophthalmology
7. American Optometric Association
8. Asia Pacific Council of Optometry
9. International Council of Ophthalmology
10. Royal Australian and New Zealand College of Ophthalmologists
11. Royal College of Ophthalmologists
12. European Society of Cataract and Refractive Surgeons
13. American Society of Cataract and Refractive Surgeons
14. European Vitreoretinal Society
15. African Ophthalmology Council
16. Asia Pacific Academy of Ophthalmology

## Appendix 2

### Search strategy for academic databases

#### MEDLINE

- 1 exp eye diseases/ (557476)
- 2 visually impaired persons/ (2391)
- 3 or/1-2 (558065)
- 4 Practice Guidelines as Topic/ (115227)
- 5 Guidelines as Topic/ (39163)
- 6 Practice Guideline/ (26462)
- 7 ((clinical or practice) adj2 guideline\$.tw. (41838)
- 8 or/4-7 (202728)
- 9 3 and 8 (2180)
- 10 exp case reports/ (2078744)
- 11 case report\$.tw. (356889)
- 12 or/10-11 (2160660)
- 13 9 not 12 (2112)
- 14 limit 13 to english language (1676)
- 15 limit 14 to yr="2009 -Current" (949)
- 16 limit 15 to (address or autobiography or bibliography or biography or classical article or clinical trial, veterinary or clinical trials, veterinary as topic or comment or directory or editorial or "expression of concern" or festschrift or historical article or interactive tutorial or interview or letter or news or newspaper article or observational study, veterinary or personal narrative or portrait or video-audio media or webcast) (113)
- 17 15 not 16 (836)

#### Embase

- 1 exp eye disease/cn, di, dm, ep, et, pc, rh, su, th [Congenital Disorder, Diagnosis, Disease Management, Epidemiology, Etiology, Prevention, Rehabilitation, Surgery, Therapy] (314642)
- 2 visual impairment/cn, di, dm, ep, et, pc, rh, su, th [Congenital Disorder, Diagnosis, Disease Management, Epidemiology, Etiology, Prevention, Rehabilitation, Surgery, Therapy] (6819)
- 3 or/1-2 (314642)
- 4 practice guidelines/ (405820)
- 5 ((clinical or practice) adj2 guideline\$.tw. (60447)
- 6 or/4-5 (428306)
- 7 3 and 6 (3082)
- 8 exp case report/ (2347870)
- 9 case report\$.tw. (464572)
- 10 or/8-9 (2402846)
- 11 7 not 10 (2932)
- 12 limit 11 to english language (2574)
- 13 limit 12 to yr="2009 -Current" (1346)
- 14 limit 13 to (conference abstract or conference paper or editorial or letter or note or short survey or tombstone) (257)
- 15 13 not 14 (1089)

### Global Health

- 1 exp eye diseases/ (32535)
- 2 people with visual impairment/ (170)
- 3 or/1-2 (32658)
- 4 exp guideline/ (47207)
- 5 ((clinical or practice) adj2 guideline\$.tw. (4484)
- 6 or/4-5 (49229)
- 7 3 and 6 (333)
- 8 case reports/ (86862)
- 9 case report\$.tw. (94202)
- 10 8 or 9 (94202)
- 11 7 not 10 (329)
- 12 limit 11 to english language (279)
- 13 limit 12 to yr="2009 -Current" (197)

### CINAHL

- S5 S1 AND S4 Limiters - Published Date: 20090101-20200231; English Language;  
Exclude MEDLINE records
- S4 S2 OR S3
- S3 (TX clinical OR TX practice) N2 (guideline\*)
- S2 MM "Practice Guidelines"
- S1 MH "Eye Diseases+"

### WHO Global Index Medicus

#### SEARCH 1

(ab:(practice guideline)) AND (ab:(eye OR vision OR visual OR blindness ))

#### SEARCH 2

(mh:(eye diseases)) AND (mh:(practice guideline))

## Appendix 3

**AGREE II items used for appraisal of clinical practice guidelines**

<b>AGREE II Section</b>	<b>Statement</b>
4	The guideline development group includes individuals from all relevant professional groups.
7	Systematic methods were used to search for evidence.
8	The criteria for selecting the evidence are clearly described.
10	The methods for formulating the recommendations are clearly described.
12	There is an explicit link between the recommendations and the supporting evidence.
13	The guideline has been externally reviewed by experts prior to its publication.
15	The recommendations are specific and unambiguous.
22	The views of the funding body have not influenced the content of the guideline.
23	Competing interests of guideline development group members have been recorded and addressed.

## Supplementary Table

Recommended eye interventions for Uveitis and strength of recommendation and quality of evidence

Description of Intervention	Relevant Guidelines*	Strength of recommendation (strong, intermediate, weak)	Quality of evidence (RCT, systematic review, meta-analysis, clinical studies, expert opinion)
Adalimumab for treating non-infectious uveitis in the posterior segment of the eye in adults with inadequate response to corticosteroid	1	Strong	RCT
Stop adalimumab for non-infectious uveitis in the posterior segment of the eye in adults with inadequate response to corticosteroids if there is 1 of the following: new active inflammatory chorioretinal or inflammatory retinal vascular lesions, or both, or a 2-step increase in vitreous haze or anterior chamber cell grade or worsening of best corrected visual acuity by 3 or more lines or 15 letters.	1	Strong	Expert Opinion
Dexamethasone intravitreal implant is recommended as an option for treating non-infectious uveitis in the posterior segment of the eye in adults, only if there is: active disease (that is, current inflammation in the eye) and worsening vision with a risk of blindness.	1	Strong	RCT
In children and adolescents with JIA at high risk of developing uveitis, ophthalmic screening every 3 months is conditionally recommended over screening at a different frequency.	2	Weak	Clinical Studies
In children and adolescents with JIA and controlled uveitis who are tapering or discontinuing topical glucocorticoids, ophthalmic monitoring within 1 month after each change of topical glucocorticoids is strongly recommended over monitoring less frequently.	2	Strong	Expert Opinion

In children and adolescents with JIA and controlled uveitis on stable therapy, ophthalmic monitoring no less frequently than every 3 months is strongly recommended over monitoring less frequently.	2	Strong	Clinical Studies
In children and adolescents with JIA and controlled uveitis who are tapering or discontinuing systemic therapy, ophthalmic monitoring within 2 months of changing systemic therapy is strongly recommended over monitoring less frequently.	2	Strong	Clinical Studies
In children and adolescents with JIA and active CAU, using prednisolone acetate 1% topical drops is conditionally recommended over difluprednate topical drops.	2	Weak	Expert Opinion
In children and adolescents with JIA and active CAU, adding or increasing topical glucocorticoids for short-term control is conditionally recommended over adding systemic glucocorticoids	2	Weak	Clinical Studies
In children and adolescents with JIA who develop new CAU activity despite stable systemic therapy, topical glucocorticoids prior to changing/escalating systemic therapy is conditionally recommended over changing/escalating systemic therapy immediately.	2	Weak	Expert Opinion
In children and adolescents with JIA and CAU still requiring 1–2 drops/day of prednisolone acetate 1% (or equivalent) for uveitis control, and not on systemic therapy, adding systemic therapy in order to taper topical glucocorticoids is conditionally recommended over not adding systemic therapy and maintaining on topical glucocorticoids only.	2	Weak	Clinical Studies
In children and adolescents with JIA and CAU still requiring 1–2	2	Weak	Clinical Studies

drops/day of prednisolone acetate 1% (or equivalent) for at least 3 months and on systemic therapy for uveitis control, changing or escalating systemic therapy is conditionally recommended over maintaining current systemic therapy.			
In children and adolescents with JIA and CAU who are starting systemic treatment for uveitis, using subcutaneous methotrexate is conditionally recommended over oral methotrexate.	2	Weak	Expert Opinion
In children and adolescents with JIA with severe active CAU and sight-threatening complications, starting methotrexate and a monoclonal antibody TNFi immediately is conditionally recommended over ethotrexate as monotherapy.	2	Weak	Expert Opinion
In children and adolescents with JIA and active CAU starting a TNFi, starting a monoclonal antibody TNFi is conditionally recommended over etanercept.	2	Weak	RCT
In children and adolescents with JIA and active CAU who have an inadequate response to 1 monoclonal antibody TNFi at standard JIA dose, escalating the dose and/or frequency to above standard is conditionally recommended over switching to another monoclonal antibody TNFi.	2	Weak	Expert Opinion
In children and adolescents with JIA and active CAU who have failed a first monoclonal antibody TNFi at above-standard dose and/or frequency, changing to another monoclonal antibody TNFi is conditionally recommended over a biologic in another category.	2	Weak	Expert Opinion
In children and adolescents with JIA and active CAU who have failed methotrexate and 2 monoclonal antibody TNFi at	2	Weak	Clinical Studies

above-standard dose and/or frequency, the use of abatacept or tocilizumab as biologic DMARD options, and mycophenolate, leflunomide, or cyclosporine as alternative nonbiologic DMARD options is conditionally recommended.			
In children and adolescents with spondyloarthritis, strongly recommend education regarding the warning signs of AAU for the purpose of decreasing delay in treatment, duration of symptoms, or complications of iritis.	2	Strong	Expert Opinion
In children and adolescents with spondyloarthritis otherwise well controlled with systemic immunosuppressive therapy (DMARDs, biologics) who develop AAU, conditionally recommend against switching systemic immunosuppressive therapy immediately in favor of treating with topical glucocorticoids first.	2	Weak	Expert Opinion
In children and adolescents with JIA and CAU that is controlled on systemic therapy but who remain on 1–2 drops/day of prednisolone acetate 1% (or equivalent), tapering topical glucocorticoids first is strongly recommended over systemic therapy.	2	Strong	Clinical Studies
In children and adolescents with uveitis that is well controlled on DMARD and biologic systemic therapy only, conditionally recommend that there be at least 2 years of well-controlled disease before tapering therapy.	2	Weak	Expert Opinion
Assessing a patient with possible uveitis <ul style="list-style-type: none"> <li>• Take a medical history, and ask: About the signs and symptoms, including the onset and duration.</li> <li>• If symptoms are unilateral or bilateral.</li> </ul>	3	Strong	Expert Opinion

<ul style="list-style-type: none"> <li>• If vision is affected, or there is an increase in floaters.</li> <li>• If there is any pain – the type of pain (for example, dull, or throbbing), the intensity, and the location (ophthalmic, or non-ophthalmic eye pain).</li> <li>• About a foreign body sensation</li> <li>• If eyes are watering more than normal, or if there is any discharge</li> <li>• If there is photophobia.</li> <li>• Previous illnesses, including previous eye problems.</li> <li>• Medication</li> <li>• Whether the person wears contact lenses — if they do, ask about the hygiene routine.</li> <li>• If there is any history of chemical exposure, trauma or surgery.</li> <li>• Family history</li> </ul> <p>Conduct a physical examination:</p> <ul style="list-style-type: none"> <li>• If perforation of the globe is suspected (for instance in ocular trauma or as a complication of scleritis), do not palpate the eye — arrange for urgent ophthalmology assessment.</li> <li>• Assess for evidence of facial trauma.</li> <li>• Examine the eyelids for inflammation and erythema, or any abnormalities (for example, trichiasis, entropion or ectropion).</li> <li>• Examine the eyelids and surrounding area for rashes and vesicles.</li> <li>• Examine the conjunctiva, including the tarsal surface.</li> </ul>			
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<ul style="list-style-type: none"> <li>• If foreign body is a possibility, invert the upper lid to check for a sub-tarsal foreign body.</li> <li>• Check the pattern of redness (if present)</li> <li>• Perform fluorescein examination</li> <li>• Check for any discharge from the eye(s).</li> <li>• Check the person's visual acuity using a Snellen chart.</li> </ul>			
Refer people with severe eye pain and a significant reduction in vision immediately for same-day assessment by an ophthalmologist.	3	Strong	Expert Opinion
Refer people with suspected uveitis (new presentations, and recurrent) for assessment within 24 hours by an ophthalmologist.	3	Strong	Expert Opinion
Uveitis should be managed by an ophthalmologist	3	Strong	Expert Opinion
Do not initiate treatment for uveitis in primary care, unless asked to do so by an ophthalmologist	3	Strong	Expert Opinion

\*1: NICE – Adalimumab, 2: ACR – JIA Uveitis, 3: NICE – Uveitis CKS

AAU: Acute Anterior Uveitis, CAU: Chronic Anterior Uveitis, DMARD: Disease Modifying Anti-Rheumatic Drug, TNFi: Tumour Necrosis Factor Inhibitor