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ISSN: 2468-6530

### DESCRIPTION

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*Ophthalmology Retina*, a journal of the American Academy of Ophthalmology, serves society by publishing clinical and basic science research and other relevant manuscripts that relate to the sense of sight. Excellence is pursued through unbiased peer-review, the advancement of innovation and discovery, and the promotion of lifelong learning.

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Authorship

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Correspondence (previously Letters to the Editor) allows concise commentary about an article published in the journal within 6 months of its online posting. The text should raise a question for clarification, offer an alternative perspective, or explain a flaw in methodology or a perceived misinterpretation of data. The correspondence should address no more than three points. Correspondence should not be used as an avenue to introduce new material without subjecting it to typical peer review.

Format: Correspondence is limited to 700 words, double-spaced, with no more than 5 references including the article to which the authors are responding. Figures, tables, or graphs are typically not included. The title follows the following format: Re: [insert last name of first author of published article] et al.: [insert title of the published article to which the Correspondence refers.] The correspondence should start with "To the Editor" and the article being commented on should be referenced in the first paragraph and be the first listed reference. Comments such as "... I commend the author for their fine study" or overly critical remarks are neither necessary nor appropriate. Letters should end with the name, degree, and location (city, state or city, country) for each author.

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All correspondence and replies are published online, although the material is listed in the print Table of Contents.

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Do not use drug trade names in titles. Please use the generic name in the abstract, as appropriate, but include the trade name once, in parentheses, after the first use of the generic name. Similarly, in the text, use the generic name, but include the trade name once, in parentheses, after the first use of the generic name.

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Photographs (including those generated electronically from MRI, fluorescein angiography, perimetry, OCT, etc.) must be masked to prevent patient identification. Clinical photographs that permit identification of an individual (those exposing anything more than just the eyes) must be accompanied by a signed statement by the patient or guardian granting permission for publication of the images for educational purposes. All graphics, including composites (such as clinical photographs, fluorescein angiography, CT, MRI, OCT, photomicrographs, etc.) should be submitted at the actual size that they would be presented in the journal, i.e., 100% of their print dimensions to avoid scaling. The width should be no more than 7 inches.

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Double-space the entire manuscript after the title page and add continuous line numbering to the manuscript file. The average published manuscript in *Ophthalmology Retina*, including references, is 6 printed pages or less. This corresponds, depending on font size and printing, to 16-20 pages of double-spaced draft.

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### 2. Abstract – see separate "Abstract" section

**3. Texta.** Introduction: Without a heading, the two- to three-paragraph introduction should explain why the study was done and in particular what hypothesis is being tested. The introduction should refer only to the most pertinent past publications and should not be an extensive review of the literature. b. Methods, Intervention, or Testing: This section should be written with sufficient detail to permit others to duplicate the work. Also required are the following, as appropriate within the methods section:

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The journal welcomes submission of high quality photographs, photomicrographs, radiologic or other imaging studies, or procedural illustrations that depict novel features of clinically important entities. Single images or a related pair of images may be submitted and the accompanying legend should be 100 words or fewer. There is a limit of three authors. If accepted, the submission will be published when space permits in the print journal.

To submit an image for consideration, please log in to <https://www.editorialmanager.com/ORET/default.aspx> as an Author and select "Pictures & Perspectives" as the Article Type for a New Submission. Upload a single or composite high resolution .tif image file and a Word document for the title, author byline, and legend. *Ophthalmology Retina* will need a completed copyright transfer form at acceptance (see [Downloadable Forms](#)). Once the form is accepted and transmitted, the Editorial Office will assign the image for a future issue. Images submitted by photographers and clinicians in this manner are used for the "Pictures & Perspectives" section occasionally, so it may be several months before it appears in print.

### **Précis**

All full-length manuscripts must include a précis of 35 words or less summarizing the main finding/outcome of the study. The précis should not duplicate the abstract conclusion. If the paper is published, the précis will appear under the title in the Table of Contents. The précis is submitted as a separate file and should not be included in the manuscript file. Please refrain from using abbreviations/acronyms in the précis.

### **Prior and Repetitive Publication; Plagiarism**

The journal will not consider manuscripts that have appeared in other journals, in part or in total, in other publications, except in special circumstances approved by the Editor-in-Chief. Likewise, updates of previously published studies that add minimal new information to an existing publication will not be considered. Overlap between patient groups described in serial manuscripts must be acknowledged, and references to previous publications that include the same patients must be provided. Authors uncertain as to whether specific data might be considered prior or repetitive publication should alert the Editor-in-Chief on the cover letter and provide copies of the publications in question.

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### **Precedence**

Authors who claim precedence for an idea, observation, or therapy should describe the literature search methodology used to support their assertion.

### **Reference Format**

Indicate references by (consecutive) superscript arabic numerals in the order in which they appear in the text. The numerals are to be used *outside* periods and commas, *inside* colons and semicolons. For further detail and examples please refer to the AMA Manual of Style, A Guide for Authors and Editors, Tenth Edition, ISBN 0-978-0-19-517633-9.

Number the references in the list in the order in which they appear in the text.

Most manuscripts in *Ophthalmology Retina* are neither intended to be review articles nor require encyclopedic referencing. Twenty or 30 references suffice for the majority of manuscripts and nearly all can be presented with less than 40.

### **Examples:**

Reference to a journal publication:

1. Van der Geer J, Hanraads JAJ, Lupton RA. The art of writing a scientific article. *J Sci Commun*. 2010;163:51-59.

Reference to a book:

2. Strunk W Jr, White EB. *The Elements of Style*. 4th ed. New York, NY: Longman; 2000.

Reference to a chapter in an edited book:



3. Mettam GR, Adams LB. How to prepare an electronic version of your article. In: Jones BS, Smith RZ, eds. *Introduction to the Electronic Age*. New York, NY: E-Publishing Inc; 2009:281-304.

Reference to a website:

4. Cancer Research UK. Cancer statistics reports for the UK. <http://www.cancerresearchuk.org/aboutcancer/statistics/cancerstatsreport/>; 2003 Accessed 13.03.03.

Dataset:

5. Oguro M, Imahiro S, Saito S, Nakashizuka T. Mortality data for Japanese oak wilt disease and surrounding forest compositions, Mendeley Data, v1; 2015. <http://dx.doi.org/10.17632/xwj98nb39r.1>

### **Reporting Refractive Surgery Outcomes and Astigmatism**

[Astigmatism\\_Reporting\\_links\\_to\\_Reporting\\_Refractive\\_Surgery\\_Outcomes\\_and\\_Astigmatism](#) When reporting refractive surgery outcomes, please include 6 graphs to illustrate the following (references 1-3): Uncorrected distance visual acuity Change in corrected distance visual acuity Spherical equivalent (attempted versus achieved) Spherical equivalent refractive accuracy Spherical equivalent refraction stability Refractive astigmatism

Descriptions of astigmatism should adhere to terminology and graphical representations originally described by Alpíns (references 4-6). An editorial by Reinstein et al (reference 7) presents the argument for standardization.

Waring GO III, Reinstein DZ, Dupps WJ, Kohnen T, Mamalis N, Rosen ES, Koch DD, Obstbaum SA, Stulting RD. Standardized graphs and terms for refractive surgery results. *J Refract Surg* 2011;27:7-Erratum in *J Refract Surg* 2011;27:88. Reinstein DZ, Waring GO III. Graphic reporting of outcomes of refractive surgery. *J Refract Surg* 2009;5:975-8. Waring GO III. Standard graphs for reporting refractive surgery. *J Refract Surg* 2000;16:459-66. Erratum in *J Refract Surg* 2001;17:following table of contents. Alpíns N. Astigmatism analysis by the Alpíns method. *J Cataract Refract Surg* 2001;27:31-49. Alpíns NA. Vector analysis of astigmatism changes by flattening, steepening, and torque. *J Cataract Refract Surg* 1997;23:1503-14. Alpíns NA. A new method of analyzing vectors for changes in astigmatism. *J Cataract Refract Surg* 1993;19:524-33. Reinstein DZ, Archer TJ, Randleman JB. JRS standard for reporting astigmatism outcomes of refractive surgery. *J Refract Surg* 2014;30:654-9. Erratum in: *J Refract Surg* 2015;3:129.

### **Reports**

Reports are typically submitted after invitation from the Editorial Board. Specifically, some full-length manuscripts contain noteworthy information that can be presented in a more concise communique. The Editorial Board may invite the authors to abridge their work, taking into consideration suggestions for revision in the initial reviews, and resubmit the paper as a Report. Reports do not exceed 1000 words or include more than 7 references, and may feature one figure, graph, chart, or concise table on the print version. Three additional items can be included as online supplemental material. Please insert "(available at <https://www.opthalmologyretina.org/>)" at relevant point(s) in your manuscript. Please note that online supplemental material must conform to the same requirements regarding legends, abbreviations, etc. as for the print publication. A 35-word unstructured abstract is required for editors'/reviewers' view only and will not publish with the report. The text should be in narrative rather than a structured format. When uploading Reports, please select the "Case Report" submission type, select "Manuscript to Report (Invited)" as the Manuscript Category, and include the manuscript number of the original submission on the cover letter. Please include a point-by-point response to the original reviewer(s)' questions and suggestions. Please note that an acknowledgment section is reserved for grants and funding only.

### **Review Articles**

#### *Systematic Reviews and Meta-analysis*

Systematic reviews seek to collect and critically assess all evidence that fits pre-specified criteria to answer a clinical question pertaining to the cause, diagnosis, prognosis, prevention, or therapy for a condition. A systematic review may contain a meta-analysis, which uses statistical methods to combine results from similar but independent studies.

Features of a systematic review include “a clearly stated set of objectives with pre-defined eligibility criteria for studies; an explicit, reproducible methodology; a systematic search that attempts to identify all studies that would meet the eligibility criteria; an assessment of the validity of the findings of the included studies, for example through the assessment of risk of bias; and a systematic presentation, and synthesis of the characteristics and findings of the included studies (Higgins JPT, Green S (editors). Chapter 1. *Cochrane Handbook for Systematic Reviews of Interventions* Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011).

It is possible to conduct a systematic review and meta-analysis of the evidence supporting any type of research question, whether the question is about intervention effectiveness or harm, etiology, prognosis, diagnostic accuracy, toxicity, incidence, or prevalence. Where intervention effectiveness questions are typically addressed by randomized controlled trials, most other questions are addressed using observational studies. Systematic reviews may be conducted for human or animal studies, in vivo or in vitro.

For standards and classic references in conducting systematic reviews and meta-analyses, please refer to: Institute of Medicine. *Finding what works in health care: standards for systematic reviews*. 2011. Chandler J, Churchill R, Higgins J, Tovey D. *Methodological standards for the conduct of new Cochrane Intervention Reviews*. Version 2.2. 17 December 2012. Higgins JPT, Green S (editors). *Cochrane Handbook for Systematic Reviews of Interventions* Version 5.1.0 [updated March 2011]. The Cochrane Collaboration, 2011. *Handbook for Diagnostic Accuracy Reviews* [Draft] Little J, Higgins JPT (editors). *The HuGENE™ HuGE Review Handbook*, version 1.0. Guidelines for systematic review and meta-analysis of gene disease association studies (see also Systematic Reviews of Genetic Association Studies, PLoS Medicine 2009;6(3):e1000028) *Systematic Reviews*. CRD's guidance for undertaking reviews in health care. Centre for Reviews and Dissemination, University of York, 2009

For reporting systematic reviews and meta-analyses, if you are submitting a report of a systematic review and/or meta-analysis of randomized controlled trials, please follow the PRISMA guidelines for reporting; A systematic review and/or meta-analysis of observational studies, please follow the MOOSE guidelines for reporting.

A complete list of guidelines for reporting systematic reviews and meta-analyses can be found at the Enhancing the Quality and Transparency Of health Research (EQUATOR) network's website. We strongly recommend you visit the EQUATOR's website for reporting guidelines for systematic reviews and meta-analyses of other study designs (e.g., individual participant data, health equity, genetic association studies). The Cochrane Collaboration also has developed Standards for the Reporting of Cochrane Intervention Reviews.

### **Title Page:**

The title should clearly describe the research question and identify the report as a systematic review, meta-analysis, or both in the subtitle. (Example: Anti-vascular endothelial growth factor for neovascular age-related macular degeneration - A systematic review and meta-analysis.)

### **Prcis:**

The prcis should indicate a new insight the article offers or a principal controversy that is addressed.

### **Structured Abstracts:**

Abstracts for systematic reviews and meta-analysis must be limited to 350 words and include five sections following the PRISMA guidelines: Topic: provide an explicit statement of the specific clinical question being addressed with reference to a brief description of the participants, interventions (or exposures), comparators, and outcomes examined. Clinical relevance: characterize the magnitude and importance of the condition; when relevant, define the current standard of care. Methods: describe the key eligibility criteria for including studies in the systematic review, key databases searched and search dates, methods of assessing the risk of bias in the individual studies. Results: summarize the number and type of included studies and participants, and relevant characteristics of studies; describe the results of main outcomes (benefits and harms), preferably indicating the number of studies and participants for each. If a meta-analysis was done, include summary measures and

confidence intervals; report the direction of the effect or association (i.e., which group is favored) and size of the effect using language meaningful to clinicians and patients. Conclusion: summarize the strengths and limitations of the evidence, your general interpretation of the results, and important implications.

Note that the abstract content and conclusions should agree with what is in the manuscript text.

## Manuscript text

The text should use standard journal formatting and be divided into four distinct sections. The brief descriptions below are gathered from the PRISMA, the MOOSE guidelines, and the Standards for the Reporting of Cochrane Intervention Reviews. The text should report institutional review board approval or exemption, financial disclosures and potential conflicts of interest of the authors, and funding sources of the review.

1. **Introduction** (unlabeled) should provide a concise description of the condition or clinical problem addressed by the review question, provide perspectives on the importance of its management to patient well-being and quality of life, and why it is important to do the review. Always end the introduction with a clear and concise statement of the study's main objectives or hypotheses.

2. **Methods:** The methods section should include the following subheadings: *Eligibility criteria for considering studies for this review:* state eligibility criteria for participants, interventions (or exposures) and comparators, and eligible study design(s) if applicable. Define primary and secondary outcomes of the review and state whether an article had to report measurement of at least one of the outcomes to be eligible. If so, provide rationale. *Search methods for identifying studies:* list all information sources searched, including databases, trial registries, websites, difficult-to-access literature (e.g., grey literature, conference proceedings), reference lists of included studies, and whether individuals or organizations were contacted. For all searches, provide the date of the last search and whether there was any time period or language restriction. Present the exact full search strategy (or strategies) used for at least one database in an Appendix with sufficient detail to permit replication. Report which software was used to manage the records identified and eligibility status. *Study selection:* describe the process for selecting studies, how many people were involved at each step of the review, whether any steps were done by more than one person, and if so whether they worked independently and how different opinions were resolved. *Data collection and risk of bias assessment:* List and define data items extracted from the reports of included studies. Describe methods used for assessing risk of bias of included studies (risk of bias is a formal assessment of what is often considered study "quality"), and how this information was used in any data synthesis. Describe the process for data extraction and risk of bias assessment, how many people were involved at each step, whether any steps were done by more than one person, and if so whether they worked independently and how different opinions were resolved. Report the software used for data collection and management. *Data synthesis and analysis:* state the methods for combining results across studies, which include qualitative synthesis (see Chapter 4, section on "Qualitative Synthesis of the Body of Evidence; *Finding what works in health care: standards for systematic reviews*) and quantitative synthesis (i.e., meta-analysis). State the summary measures used to quantify the treatment effect or association such as risk ratio, odds ratio, and difference in means. Describe methods for assessing clinical, methodological, and statistical heterogeneity (e.g., I<sup>2</sup> statistic, tau-squared, statistical test). Describe methods for additional analyses such as meta-regression, subgroup analysis, and sensitivity analysis, if done, indicate which were pre-specified. State the statistical software used for analysis. Indicate whether a systematic review protocol exists, if so, where and how it can be accessed; and if available, provide systematic review registration information including registration number.

3. **Results:** Provide numbers of studies retrieved, screened, assessed in full for eligibility, included in the review, and included in the meta-analysis, with reasons for exclusion at each stage, ideally with a flow diagram. Present characteristics of included studies including information on the study design, participants, interventions (or exposures) and comparators, outcomes, and source of funding, ideally in a table. Present domain-based risk of bias assessment of each study, ideally in a table or a figure. Composite quality scores and scales are discouraged. For all outcomes considered, irrespective of the direction or strength of the results, present, (1) simple summary data for each group, and (2) estimates of treatment effect (or association) between groups with a measure of

statistical uncertainty (e.g., confidence intervals). If meta-analysis was done, report meta-analytical results ideally with a forest plot, number of studies and participants for each meta-analysis, as well as measures of statistical heterogeneity. Present results of any additional analyses (such as meta-regression, subgroup analysis, and sensitivity analysis) if done. Provide a thoughtful qualitative synthesis by analyzing the nature, strengths, and weaknesses of the evidence, and developing a deeper understanding of how an intervention might work (or not), or whether a true association exists, for whom and under what circumstances.

**4. Discussion:** Summarizes the main findings including the strength of evidence for each main outcome. Provide a general interpretation of the evidence considering their relevance to key stakeholders, including patients, healthcare providers, researchers, payers, and policy makers. A Summary of Findings or GRADE table is optional. Discuss limitations at study and outcome level (such as risk of bias), and at review level (such as incomplete retrieval of identified studies, reporting biases). Provide a general interpretation of the results in the context of other evidence, and implications for practice and future research.

In the cover letter to the Editor, please state explicitly (1) whether reporting guidelines have been followed, if so, which reporting guidelines; (2) whether the exact full search strategy (or strategies) used for at least one database was presented in an Appendix with sufficient detail to permit replication. Failure to follow the reporting guidelines or upload the search strategy may result in delay in review or rejection of the manuscript. Please submit a PRISMA worksheet and diagram as separate files.

Checklist: <http://prisma-statement.org/documents/PRISMA%202009%20checklist.doc>

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#### Translational Science Reviews

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The CONSORT Worksheet <http://www.consort-statement.org/Media/Default/Downloads/CONSORT%202010%20Checklist.doc> for randomized controlled trials has been required since 1996 and is available online. The following chart ([https://www.elsevier.com/\\_\\_data/promis\\_misc/OPHTHA\\_STUDY\\_DESIGN.docx](https://www.elsevier.com/__data/promis_misc/OPHTHA_STUDY_DESIGN.docx)) provides basic information regarding study designs.

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Taichman DB, Backus J, Baethge C, et al. Sharing Clinical Trial Data: A Proposal From the International Committee of Medical Journal Editors. *JAMA* 2016;315(5):467-468.

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