

Supplementary Online Content

Chou R, Dana T, Bougatsos C, Grusing S, Blazina I. Screening for impaired visual acuity in older adults: updated evidence report and systematic review for the US Preventive Services Task Force. *JAMA*. doi:10.1001/jama.2016.0783.

eAppendix 1. Search Strategies

eAppendix 2. United States Preventive Services Task Force Quality Rating Criteria

eTable 1. Quality Ratings of Trials of Screening for Impaired Visual Acuity in Older Adults

eTable 2. Quality Ratings of Diagnostic Accuracy Studies for Impaired Visual Acuity in Older Adults

eTable 3. Quality Ratings of Treatment Studies of Antioxidant Vitamins, Minerals, and Other Supplements for Dry AMD Published Since the Prior Review

This supplementary material has been provided by the authors to give readers additional information about their work.

eAppendix 1. Search Strategies

Screening Key Questions

Database: Ovid MEDLINE(R) without Revisions

- 1 Mass Screening/
- 2 exp Vision Tests/
- 3 exp Refractive Errors/
- 4 exp Macular Degeneration/
- 5 exp Vision Disorders/
- 6 exp Vision, Ocular/
- 7 exp Eye Diseases/
- 8 Cataract/
- 9 (presbyop\$ or myop\$ or astigmati\$ or hyperop\$ or cataract\$ or "macular degeneration" or armd).mp.
- 10 or/2-9
- 11 1 and 10
- 12 11 not (adolescen\$ or child\$ or pediatric\$ or toddler or infant\$ or newborn or neonat\$ or prematur\$).mp.
- 13 limit 12 to humans
- 14 limit 13 to english language
- 15 limit 13 to abstracts
- 16 14 or 15

Database: EBM Reviews - Cochrane Central Register of Controlled Trials

- 1 Mass Screening/
- 2 exp Vision Tests/
- 3 exp Refractive Errors/
- 4 exp Macular Degeneration/
- 5 exp Vision Disorders/
- 6 exp Vision, Ocular/
- 7 exp Eye Diseases/
- 8 Cataract/
- 9 (presbyop\$ or myop\$ or astigmati\$ or hyperop\$ or cataract\$ or "macular degeneration" or armd).mp.
- 10 or/2-9
- 11 1 and 10
- 12 11 not (adolescen\$ or child\$ or pediatric\$ or toddler or infant\$ or newborn or neonat\$ or prematur\$).mp.

Database: EBM Reviews - Cochrane Database of Systematic Reviews

- 1 ((vision or visual) adj5 screen\$).mp.
- 2 (presbyop\$ or myop\$ or astigmati\$ or hyperop\$).mp.
- 3 (macula\$ adj3 degenerat\$).mp.
- 4 cataract\$.mp.
- 5 1 and (or/2-4)
- 6 5 not (child\$ or pediater\$ or neonat\$ or prematur\$).mp.

Diagnostic Accuracy Key Question

Database: Ovid MEDLINE(R) without Revisions

- 1 Vision, Ocular/
- 2 Vision Disorders/
- 3 Vision Tests/
- 4 Refractive Errors/
- 5 Macular Degeneration/
- 6 Cataract/

7 Eye Diseases/
 8 (presbyop\$ or myop\$ or astigmati\$ or hyperop\$ or cataract\$ or "macular degeneration" or armd).mp.
 9 vision.mp.
 10 or/1-9
 11 screen\$.mp.
 12 10 and 11
 13 "Sensitivity and Specificity"/
 14 (specificity or accurac\$ or "predictive value").tw.
 15 (sensitiv\$ or diagnostic).mp.
 16 or/13-15
 17 12 and 16
 18 17 not (adolescen\$ or child\$ or pediatric\$ or toddler or infant or neonat\$ or prematur\$).mp.
 19 limit 18 to humans
 20 limit 19 to english language
 21 limit 19 to abstracts
 22 20 or 21

Database: EBM Reviews - Cochrane Central Register of Controlled Trials

1 Vision, Ocular/
 2 Vision Disorders/
 3 Vision Tests/
 4 Refractive Errors/
 5 Macular Degeneration/
 6 Cataract/
 7 Eye Diseases/
 8 (presbyop\$ or myop\$ or astigmati\$ or hyperop\$ or cataract\$ or "macular degeneration" or armd).mp.
 10 or/1-9
 11 screen\$.mp.
 12 10 and 11
 13 "Sensitivity and Specificity"/
 14 (specificity or accurac\$ or "predictive value").tw.
 15 (sensitiv\$ or diagnostic).mp.
 16 or/13-15
 17 12 and 16
 18 17 not (adolescen\$ or child\$ or pediatric\$ or toddler or infant or neonat\$ or prematur\$).mp.

Database: EBM Reviews - Cochrane Database of Systematic Reviews

1 (presbyop\$ or myop\$ or astigmati\$ or hyperop\$).mp.
 2 (macula\$ adj3 degenerat\$).mp.
 3 cataract\$.mp.
 4 visual acuity.mp.
 6 (diagno\$ adj2 accur\$).mp.
 7 5 and 6

Treatment Key Questions

Database: Ovid MEDLINE(R) without Revisions

1 exp Refractive Errors/dt, pc, rt, th
 2 exp Cataract/dh, dt, pc, rt, th
 3 Cataract Extraction/
 4 exp Macular Degeneration/dh, dt, pc, rt, su, th
 5 exp Vision Disorders/dh, dt, pc, rt, su, th
 6 or/1-5
 7 6 not (adolescen\$ or child\$ or pediatric\$ or toddler or infant\$ or newborn or neonat\$ or prematur\$).mp.
 8 limit 7 to humans

- 9 limit 8 to english language
- 10 limit 8 to abstracts
- 11 limit 10 to "all aged (65 and over)"
- 12 limit 11 to yr="2008 - 2014"

Database: EBM Reviews - Cochrane Central Register of Controlled Trials

- 1 exp Refractive Errors/dt, pc, rt, th
- 2 exp Cataract/dh, dt, pc, rt, th
- 3 Cataract Extraction/
- 4 exp Macular Degeneration/dh, dt, pc, rt, su, th
- 5 exp Vision Disorders/dh, dt, pc, rt, su, th
- 6 or/1-5
- 7 6 not (adolescen\$ or child\$ or pediatric\$ or toddler or infant\$ or newborn or neonat\$ or prematur\$).mp.

Database: EBM Reviews - Cochrane Database of Systematic Reviews

- 1 ("age-related macular degeneration" or "age related macular degeneration" or "AMD" or "ARMD").ti,ab.
- 2 ("impaired visual acuity" or "impaired vision" or "visual acuity").ti,ab.
- 3 cataract\$.ti,ab.
- 4 (presbyop\$ or myop\$ or astigmati\$ or hyperop\$).ti,ab.
- 5 treatment.ti,ab.
- 6 (or/1-4) and 5
- 7 6 not (child\$ or pediater\$ or neonat\$ or prematur\$).mp.
- 8 limit 7 to new reviews
- 9 limit 8 to full systematic reviews
- 10 9 not diabet\$.mp.

eAppendix 2. United States Preventive Services Task Force Quality Rating Criteria

Criteria for Assessing Internal Validity of Individual Studies

The Methods Work Group for the US Preventive Services Task Force (USPSTF) developed a set of criteria by which the internal validity of individual studies could be evaluated. The USPSTF accepted the criteria, and the associated definitions of quality categories, that relate to internal validity at its September 1999 meeting.

This appendix describes the criteria relating to internal validity and the procedures that topic teams follow for all updates and new assessments in making these judgments.

All topic teams use initial "filters" to select studies for review that deal most directly with the question at issue and that are applicable to the population at issue. Thus, studies of any design that use outdated technology or that use technology that is not feasible for primary care practice may be filtered out before the abstraction stage, depending on the topic and the decisions of the topic team. The teams justify such exclusion decisions if there could be reasonable disagreement about this step. The criteria below are meant for those studies that pass this initial filter.

Presented below are a set of minimal criteria for each study design and then a general definition of three categories: "good," "fair," and "poor," based on those criteria. These specifications are not meant to be rigid rules but rather are intended to be general guidelines, and individual exceptions, when explicitly explained and justified, can be made. In general, a "good" study is one that meets all criteria well. A "fair" study is one that does not meet (or it is not clear that it meets) at least one criterion but has no known "fatal flaw." "Poor" studies have at least one fatal flaw.

Systematic Reviews

Criteria:

Comprehensiveness of sources considered/search strategy used.

Standard appraisal of included studies.

Validity of conclusions.

Recency and relevance are especially important for systematic reviews.

Definition of ratings from above criteria:

Good: Recent, relevant review with comprehensive sources and search strategies; explicit and relevant selection criteria; standard appraisal of included studies; and valid conclusions.

Fair: Recent, relevant review that is not clearly biased but lacks comprehensive sources and search strategies.

Poor: Outdated, irrelevant, or biased review without systematic search for studies, explicit selection criteria, or standard appraisal of studies.

Randomized Controlled Trials and Cohort Studies

Criteria:

- Initial assembly of comparable groups:
 - For RCTs: adequate randomization, including first concealment and whether potential confounders were distributed equally among groups.
 - For cohort studies: consideration of potential confounders with either restriction or measurement for adjustment in the analysis; consideration of inception cohorts.
- Maintenance of comparable groups (includes attrition, cross-overs, adherence, contamination).
- Important differential loss to follow-up or overall high loss to follow-up.
- Measurements: equal, reliable, and valid (includes masking of outcome assessment).
- Clear definition of interventions.
- All important outcomes considered.
- Analysis: adjustment for potential confounders for cohort studies, or intention to treat analysis for RCTs.

Definition of ratings based on above criteria:

Good: Meets all criteria: Comparable groups are assembled initially and maintained throughout the study (follow-up at least 80 percent); reliable and valid measurement instruments are used and applied equally to the groups; interventions are spelled out clearly; all important outcomes are considered; and appropriate attention to confounders in analysis. In addition, for RCTs, intention to treat analysis is used.

Fair: Studies will be graded "fair" if any or all of the following problems occur, without the fatal flaws noted in the "poor" category below: Generally comparable groups are assembled initially but some question remains whether some (although not major) differences occurred with follow-up; measurement instruments are acceptable (although not the best) and generally applied equally; some but not all important outcomes are considered; and some but not all potential confounders are accounted for. Intention to treat analysis is done for RCTs.

Poor: Studies will be graded "poor" if any of the following fatal flaws exists: Groups assembled initially are not close to being comparable or maintained throughout the study; unreliable or invalid measurement instruments are used or not applied at all equally among groups (including not masking outcome assessment); and key confounders are given little or no attention. For RCTs, intention to treat analysis is lacking.

Case-Control Studies

Criteria:

- Accurate ascertainment of cases
- Nonbiased selection of cases/controls with exclusion criteria applied equally to both.
- Response rate.
- Diagnostic testing procedures applied equally to each group.
- Measurement of exposure accurate and applied equally to each group.
- Appropriate attention to potential confounding variables.

Definition of ratings based on criteria above:

Good: Appropriate ascertainment of cases and nonbiased selection of case and control participants; exclusion criteria applied equally to cases and controls; response rate equal to or greater than 80 percent; diagnostic procedures and measurements accurate and applied equally to cases and controls; and appropriate attention to confounding variables.

Fair: Recent, relevant, without major apparent selection or diagnostic work-up bias but with response rate less than 80 percent or attention to some but not all important confounding variables.

Poor: Major selection or diagnostic work-up biases, response rates less than 50 percent, or inattention to confounding variables.

Diagnostic Accuracy Studies

Criteria:

Screening test relevant, available for primary care, adequately described.

Study uses a credible reference standard, performed regardless of test results.

Reference standard interpreted independently of screening test.

Handles indeterminate results in a reasonable manner.

Spectrum of patients included in study.

Sample size.

Administration of reliable screening test.

Definition of ratings based on above criteria:

Good: Evaluates relevant available screening test; uses a credible reference standard; interprets reference standard independently of screening test; reliability of test assessed; has few or handles indeterminate results in a reasonable manner; includes large number (more than 100) broad-spectrum patients with and without disease.

Fair: Evaluates relevant available screening test; uses reasonable although not best standard; interprets reference standard independent of screening test; moderate sample size (50 to 100 subjects) and a "medium" spectrum of patients.

Poor: Has fatal flaw such as: Uses inappropriate reference standard; screening test improperly administered; biased ascertainment of reference standard; very small sample size or very narrow selected spectrum of patients.

Reference: United States Preventive Services Task Force Procedure Manual.¹

eTable 1. Quality Ratings of Trials of Screening for Impaired Visual Acuity in Older Adults

Study, Year	Random assignment	Allocation concealed	Groups similar at baseline	Eligibility criteria specified	Blinding: outcome assessors or data analysts	Intention-to-treat analysis	Reporting of attrition, contamination, etc.	Differential loss to follow-up or overall high loss to follow-up	Appropriate analysis including cluster correlation	Funding source	Quality
Eekhof, 2000 ²	Yes	Not applicable (cluster)	Yes	Yes	Unclear	No	Yes	Yes	No	Unclear	Fair
Moore, 1997 ³	Yes	Not applicable (cluster)	Yes	Yes	Unclear	No	Yes	Yes	No	Robert Wood Johnson Clinical Scholars Program; National Institute on Aging Geriatric Academic Program	Fair
Smeeth, 2003 ⁴	Yes	Not applicable (cluster)	Yes	Yes	Unclear	No	Yes	Yes	Yes	Medical Research Council/ United Kingdom Department of Health	Good

eTable 2. Quality Ratings of Diagnostic Accuracy Studies for Impaired Visual Acuity in Older Adults

Study, Year	Appropriate spectrum of patients	Adequate sample size (>500)	Credible reference standard used	Reference standard applied to all patients	Screening test adequately described	Reference standard interpreted independently	Quality
<i>Studies from Update</i>							
Jessa, 2012 ⁵	Yes	No	Yes	Yes	Yes	Unclear	Fair
Swanson, 2009 ⁶	Yes	No	Yes	Yes	Yes	Unclear	Fair
<i>Studies from Prior Review</i>							
Ariyasu, 1996 ⁷	Unclear	No	Yes	Unclear	Yes	No	Poor-fair
Eekhof, 2000 ⁸	Unclear	Yes	Yes	Yes	Yes	Unclear	Fair
Hiller, 1983 ⁹	Yes	Yes	Yes	Unclear	Yes	Unclear	Fair
Ivers, 2001 ¹⁰	Unclear	Yes	Yes	Unclear	Yes	Unclear	Poor-fair
McMurdo, 1988 ¹¹	Unclear	No	Yes	Yes	Yes	Yes	Fair
Teh, 2006 ¹²	Unclear	No	Yes	Unclear	Yes	Unclear	Poor-fair
Wang, 1998 ¹³	Unclear	No	Yes	No	Yes	Unclear	Poor-fair
Woods, 1998 ¹⁴	Unclear	Yes	Yes	No	Yes	Unclear	Poor-fair

Note: Studies from prior review not re-rated.

eTable 3. Quality Ratings of Treatment Studies of Antioxidant Vitamins, Minerals, and Other Supplements for Dry AMD Published Since the Prior Review

Systematic Review: Quality = Good

Author, Year	A priori design provided	Duplicate study selection and data abstraction -Study selection -Data abstraction	Comprehensive literature search performed	Non-English language studies considered for inclusion	Conducted searches for unpublished (gray) literature	List of included studies provided	List of excluded studies provided with reasons	Characteristics of the included studies provided	Scientific quality of included studies: Assessed Documented	Sensitivity analyses or stratified analyses conducted according to study quality	Study conclusions supported by the evidence (Was study quality considered in the synthesis?)	Conflict of interest stated: Systematic Review Individual Studies
Evans, 2012 ¹⁵	Yes	- Yes - Yes	Yes	Yes	Yes	Yes	Yes	Yes	-Yes -Yes	No	Yes	-Yes -No

Trials

Author, year	Randomization adequate?	Allocation concealment adequate?	Groups similar at baseline?	Eligibility criteria specified?	Outcome assessors masked?	Care provider masked?	Patient masked?	Attrition and withdrawals reported?	Loss to follow-up: differential/high?	Analyze people in the groups in which they were randomized?	Quality
Chew, 2013 ¹⁶ (AREDS)	Yes	Unclear	Yes	Yes	Yes	Unclear	Yes	Yes	No	Yes	Good
Ma, 2012 ¹⁷	Yes	Unclear	Yes	Yes	Yes	Yes	Yes	Yes	No	Yes	Good
Murray, 2013 ¹⁸	Yes	Unclear	Yes	Yes	Yes	Unclear	Yes	Yes	No	Yes	Good
Souied, 2013 ¹⁹	Yes	Unclear	Yes	Yes	Yes	Yes	Yes	Yes	No	Yes	Good

References

1. Procedure Manual. U. S. Preventive Services Task Force. <http://www.uspreventiveservicestaskforce.org/Page/Name/procedure-manual>. Accessed July 16, 2015.
2. Eekhof J, De Bock G, Schaapveld K, Springer M. Effects of screening for disorders among the elderly: an intervention study in general practice. *Fam Pract*. 2000;17(4):329-333.
3. Moore AA, Siu A, Partridge JM, Hays RD, Adams J. A randomized trial of office-based screening for common problems in older persons. *Am J Med*. 1997;102(4):371-378.
4. Smeeth L, Fletcher AE, Hanciles S, Evans J, Wormald R. Screening older people for impaired vision in primary care: cluster randomised trial. *BMJ*. 2003;327(7422):1027-1031.
5. Jessa Z, Evans BJW, Thomson DW. The development & evaluation of two vision screening tools for correctable visual loss in older people. *Ophthalmic Physiol Opt*. 2012;32(4):332-348.
6. Swanson MW, McGwin G, Jr., Elliott AF, Owsley C. The nursing home minimum data set for vision and its association with visual acuity and contrast sensitivity. *J Am Geriatr Soc*. 2009;57(3):486-491.
7. Ariyasu RG, Lee PP, Linton KP, LaBree LD, Azen SP, Siu AL. Sensitivity, specificity, and predictive values of screening tests for eye conditions in a clinic-based population. *Ophthalmology*. 1996;103(11):1751-1760.
8. Eekhof JA, De Bock GH, Schaapveld K, Springer MP. Screening for hearing and visual loss among elderly with questionnaires and tests: which method is the most convincing for action? *Scand J Prim Health Care*. 2000;18(4):203-207.
9. Hiller R, Krueger DE. Validity of a survey question as a measure of visual acuity impairment. *Am J Public Health*. 1983;73:93-96.
10. Ivers RQ, Optom B, Macaskill P, Cumming RG, Mitchell P. Sensitivity and specificity of tests to detect eye disease in an older population. *Ophthalmology*. 2001;108(5):968-975.
11. McMurdo M, Baines P. The detection of visual disability in the elderly. *Health Bulletin*. 1988;46(6):327-329.
12. Teh RC, Lim WS. Utility of a patient-response screening question for visual impairment. *J Am Geriatr Soc*. 2006;54(2):370-372.
13. Wang F, Tielsch JM, Ford DE, Quigley HA, Whelton PK. Evaluation of screening schemes for eye disease in a primary care setting. *Ophthalmic Epidemiol*. 1998;5(2):69-82.
14. Woods RL, Tregear SJ, Mitchell RA. Screening for ophthalmic disease in older subjects using visual acuity and contrast sensitivity. *Ophthalmology*. 1998;105(12):2318-2326.
15. Evans JR, Lawrenson JG. Antioxidant vitamin and mineral supplements for slowing the progression of age-related macular degeneration. *Cochrane Database Syst Rev*. 2012;11. Art. No.: CD000254. DOI: 10.1002/14651858. CD000254.pub3:CD000254.
16. Chew EY, Clemons TE, Agron E, et al. Long-term effects of vitamins C and E, beta-carotene, and zinc on age-related macular degeneration: AREDS report no. 35. *Ophthalmology*. 2013;120(8):1604-1611.e1604.
17. Ma L, Yan SF, Huang YM, et al. Effect of lutein and zeaxanthin on macular pigment and visual function in patients with early age-related macular degeneration. *Ophthalmology*. 2012;119(11):2290-2297.
18. Murray IJ, Makridaki M, van der Veen RLP, Carden D, Parry NRA, Berendschot TTJM. Lutein supplementation over a one-year period in early AMD might have a mild beneficial effect on visual acuity: the CLEAR study. *Invest Ophthalmol Vis Sci*. 2013;54(3):1781-1788.
19. Souied EH, Delcourt C, Querques G, et al. Oral docosahexaenoic acid in the prevention of exudative age-related macular degeneration: the Nutritional AMD Treatment 2 study. *Ophthalmology*. 2013;120(8):1619-1631.