Until April 2006, the Canadian Agency for Drugs and Technologies in Health (CADTH) was known as the Canadian Coordinating Office for Health Technology Assessment (CCOHTA).

Cite as: Dunfield L, Keating T. Preschool vision screening [Technology Report number 73]. Ottawa: Canadian Agency for Drugs and Technologies in Health; 2007.

Production of this report is made possible by financial contributions from Health Canada and the governments of Alberta, British Columbia, Manitoba, New Brunswick, Newfoundland and Labrador, Northwest Territories, Nova Scotia, Nunavut, Ontario, Prince Edward Island, Saskatchewan, and Yukon. The Canadian Agency for Drugs and Technologies in Health takes sole responsibility for the final form and content of this report. The views expressed herein do not necessarily represent the views of Health Canada or any provincial or territorial government.

Reproduction of this document for non-commercial purposes is permitted provided appropriate credit is given to CADTH.

CADTH is funded by Canadian federal, provincial, and territorial governments.

Legal Deposit – 2007
National Library of Canada
ISBN: 1-897257-71-6 (online)
I3001 – February 2007

PUBLICATIONS MAIL AGREEMENT NO. 40026386
RETURN UNDELIVERABLE CANADIAN ADDRESSES TO
CANADIAN AGENCY FOR DRUGS AND TECHNOLOGIES IN HEALTH
600-865 CARLING AVENUE
OTTAWA ON K1S 5S8
Preschool Vision Screening

Lesley Dunfield, PhD¹
Tamara Keating, MLIS¹

February 2007

¹ Canadian Agency for Drugs and Technologies in Health, Ottawa ON
Health technology assessment (HTA) agencies face the challenge of providing quality assessments of medical technologies in a timely manner to support decision making. Ideally, all important deliberations would be supported by comprehensive health technology assessment reports, but the urgency of some decisions often requires a more immediate response.

The Health Technology Inquiry Service (HTIS) provides Canadian health care decision makers with health technology assessment information, based on the best available evidence, in a quick and efficient manner. Inquiries related to the assessment of health care technologies (drugs, devices, diagnostic tests, and surgical procedures) are accepted by the service. Information provided by the HTIS is tailored to meet the needs of decision makers, taking into account the urgency, importance, and potential impact of the request.

Consultations with the requestor of this HTIS assessment indicated that a review of the literature would be beneficial. The research question and selection criteria were developed in consultation with the requestor. The literature search was carried out by an information specialist using a standardized search strategy. The review of evidence was conducted by one internal HTIS reviewer. The draft report was internally reviewed and externally peer-reviewed by two or more peer reviewers. All comments were reviewed internally to ensure that they were addressed appropriately.
The Health Technology Inquiry Service (HTIS) is an information service for those involved in planning and providing health care in Canada. HTIS responses are based on a limited literature search and are not comprehensive, systematic reviews. The intent is to provide a list of sources and a summary of the best evidence on the topic that CADTH could identify using all reasonable efforts within the time allowed. HTIS responses should be considered along with other types of information and health care considerations. The information included in this response is not intended to replace professional medical advice, nor should it be construed as a recommendation for or against the use of a particular health technology. Readers are also cautioned that a lack of good quality evidence does not necessarily mean a lack of effectiveness particularly in the case of new and emerging health technologies, for which little information can be found, but which may in future prove to be effective. While CADTH has taken care in the preparation of the report to ensure that its contents are accurate, complete, and up to date, CADTH does not make any guarantee to that effect. CADTH is not liable for any loss or damages resulting from use of the information in the report.

Copyright: This report contains CADTH copyright material. It may be copied and used for non-commercial purposes, provided that attribution is given to CADTH.

Links: This report may contain links to other information on available on the web sites of third parties on the Internet. CADTH does not have control over the content of such sites. Use of third party sites is governed by the owners’ own terms and conditions.
ABBREVIATIONS

AAO   American Academy of Ophthalmology
AAP   American Academy of Pediatrics
AOA   American Optometric Association
CPS   Canadian Paediatric Society
ICER  incremental cost-effectiveness ratio
NCR   non-cycloplegic retinoscopy
NPV   negative predictive value
NR    not reported
NS    not specified
PPV   positive predictive value
QALY  quality-adjusted life-year
STBS  small target binocular suppression test
STRDS small target random-dot stereogram
USPSTF US Preventive Services Task Force
VA    visual acuity
VIP   Vision in Preschoolers
# TABLE OF CONTENTS

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABBREVIATIONS</td>
<td>iii</td>
</tr>
<tr>
<td>1 CONTEXT AND POLICY ISSUES</td>
<td>1</td>
</tr>
<tr>
<td>2 RESEARCH QUESTIONS</td>
<td>2</td>
</tr>
<tr>
<td>3 METHODS</td>
<td>2</td>
</tr>
<tr>
<td>3.1 Literature Search</td>
<td>2</td>
</tr>
<tr>
<td>3.2 Screening Process</td>
<td>2</td>
</tr>
<tr>
<td>4 FINDINGS</td>
<td>2</td>
</tr>
<tr>
<td>4.1 Effectiveness of Preschool Vision Screening</td>
<td>2</td>
</tr>
<tr>
<td>4.1.1 Specific screening tests</td>
<td>3</td>
</tr>
<tr>
<td>4.1.2 Screening effectiveness</td>
<td>4</td>
</tr>
<tr>
<td>4.1.3 Re-screening effectiveness</td>
<td>6</td>
</tr>
<tr>
<td>4.1.4 Personnel administering vision screening</td>
<td>6</td>
</tr>
<tr>
<td>4.1.5 Testability</td>
<td>7</td>
</tr>
<tr>
<td>4.1.6 Limitation of Studies</td>
<td>7</td>
</tr>
<tr>
<td>4.2 Current Practice</td>
<td>8</td>
</tr>
<tr>
<td>4.3 Professional Organizations</td>
<td>8</td>
</tr>
<tr>
<td>4.4 Cost Effectiveness of Preschool Vision Screening</td>
<td>10</td>
</tr>
<tr>
<td>5 CONCLUSIONS AND IMPLICATIONS</td>
<td>11</td>
</tr>
<tr>
<td>6 REFERENCES</td>
<td>11</td>
</tr>
<tr>
<td>APPENDIX 1</td>
<td>17</td>
</tr>
</tbody>
</table>


1 CONTEXT AND POLICY ISSUES

The purpose of preschool vision screening is to detect vision disorders, such as amblyopia, strabismus, and refractive errors, that occur at an early age (younger than six years old) so that they can be corrected. Amblyopia is a loss of vision that cannot be explained by ocular pathology or refractive error. Uncorrected refractive errors, media opacity, and strabismus are the causes of amblyopia. Strabismus, a misalignment of the eyes, is the most common cause of amblyopia. Strabismus includes esotropia (eye turns in), exotropia (eye turns out), and hypertropia (eye turns up). Refractive errors include myopia (nearsightedness), hyperopia (farsightedness), astigmatism, and anisometropia (unequal refractive power in the eyes). Amblyopia affects approximately 2% to 4% of the population, and strabismus affects approximately 4%.

Amblyopia and other vision disorders can be corrected if they are detected in infancy or early childhood. Visual pathways develop until about 10 years of age, and treatment of amblyopia should begin before this age if it is to be effective. Visual pathways will not develop properly in children if they do not see clear images. Amblyopia may affect health, academic, occupational, and social functioning. Left untreated, refractive errors and amblyopia may affect the ability to play sports, development, and school performance. Children with amblyopia have a risk of vision loss in the non-amblyopic eye, resulting in blindness. The early detection of vision disorders allows proper treatment to be undertaken to improve visual acuity.

Referral criteria exist for the diagnosis of amblyopia. A visual acuity (VA) of <20/40 is a common criterion. A VA of <20/30 or a 0.2 difference between the eyes can be used as referral criteria for amblyopia detection. The detection of vision problems at an early age can lead to effective treatments that may increase visual acuity. There are many methods for detecting vision disorders, with different groups reporting various sensitivities and specificities for specific tests. Tests for visual acuity include Snellen, Tumbling E, HOTV, Allen cards, and Lea Symbols. Ocular alignment is tested using the corneal light reflex, red reflex (Brückner reflex), and cross-cover tests, and random-dot stereotests (Frisby, Randot, Random-dot E, Lang). Different screening tests are recommended for different age groups. Red reflex is checked during the newborn medical examination. At this same time, the eyes are examined for any structural abnormalities. To obtain the most accurate results, the most difficult test that a child can perform should be part of the vision screening. The American Academy of Pediatrics (AAP), the American Optometric Association (AOA), the American Academy of Ophthalmology (AAO), and the US Preventive Services Task Force (USPSTF) differ in the tests that they recommend for each age group. The Canadian Paediatric Society (CPS) recommends testing of visual acuity, red reflex, and corneal light reflex in children aged three years to five years.

The effectiveness of preschool vision screening has been debated. There is no agreement about the age at which children should be screened, on the tests that should be used, and on the outcomes that should be measured. Some organizations suggest that vision examinations should only be conducted by licensed eye-care professionals. Others indicate that ophthalmologists are not the only ones who can effectively detect vision defects through preschool vision screening, and that screening tests can be conducted by nurses, primary-care physicians, or other health professionals. Trained lay screeners often administer the tests in public screening programs.
2 RESEARCH QUESTIONS

What is the published evidence on the clinical and cost effectiveness of preschool vision screening (in children under the age of six years) in comparison to a complete diagnostic vision examination by an optometrist or ophthalmologist?

3 METHODS

3.1 Literature Search

Published literature was obtained by cross-searching EMBASE®, MEDLINE®, CINAHL®, and ERIC databases on the OVID search system. Regular alerts were established on EMBASE, MEDLINE, and CINAHL, and information retrieved via alerts was current to July 21, 2006. Parallel searches were performed on PubMed and the Cochrane Library (Issue 2, 2006) databases. Publications were limited to the English language only, with publication dates of 1996 to the present. Filters were applied to limit the retrieval to health technology assessments, systematic reviews, clinical studies, and economic studies.

The web sites of regulatory agencies, and health technology assessment and related agencies were searched, as were specialized databases, such as those of the University of York’s Centre for Reviews and Dissemination. The Google™ search engine was used to search for information on the Internet. These searches were supplemented by hand-searching the bibliographies of selected papers. The British Columbia Association of Optometrists and the British Columbia Ministry of Health also supplied several documents for review.

3.2 Screening Process

Reports were included that assessed vision screening in children <6 years of age. Studies that included older children were excluded if the results were not separated into age groups that included preschool-aged children only. Three external reviewers provided comments on this report.

4 FINDINGS

4.1 Effectiveness of Preschool Vision Screening

There are criteria to consider when determining whether a screening program will be useful. The screening program should do more good than harm, the tests must be able to identify the defect, there must be an appropriate intervention to treat the defect, there must be an advantage in detecting and treating the defect at an earlier age, and the cost must be justified.25 In addition, the prevalence of the defect must be high enough and must cause substantial disability, to justify screening.25

A UK assessment determined that all these criteria were not met with preschool vision screening.5 They found insufficient evidence that treatment of these disorders resulted in visual gains,5,26 or that amblyopia, strabismus, and refractive errors caused disability. This assessment recommended that screening programs not be implemented unless they have been evaluated, because there was no evidence found for the benefits of preschool vision screening.5

A Cochrane Collaboration review on amblyopia screening concluded that insufficient evidence existed to determine the effectiveness of screening programs on amblyopia prevalence. The authors noted that this did not mean that there was no benefit to screening but rather that there was a lack of
good quality evidence (e.g., randomized controlled trials) in this area.\textsuperscript{27}

Kentucky passed a law in 2000 requiring each child entering public school to have a vision examination by an ophthalmologist or optometrist.\textsuperscript{28} A survey of Kentucky Optometric Association members found that 5,316 children were examined between July 2000 and April 2001; 13.92\% were prescribed eyeglasses, 3.4\% were diagnosed with amblyopia, and 2.31\% with strabismus.\textsuperscript{28} The Modified Clinical Technique (which consists of visual acuity tests, a cover test, retinoscopy, and ophthalmoscopy) administered by an ophthalmologist or optometrist was suggested as the only test that can effectively assess preschool vision with an acceptable sensitivity and specificity.\textsuperscript{9} This result was not unexpected, because this technique, which consisted of multiple tests administered by ophthalmologists or optometrists, would be similar to a clinical examination.

A report by C. Green, which was prepared for the BC Ministry of Health Services, suggested that there was a basis for recommending preschool vision screening, even though research evidence was lacking.\textsuperscript{4} Four options were proposed for vision testing in preschoolers: fortify existing vision care, use primary screening by public health nurses with referral to ophthalmology and optometry, use screening by technical or lay examiners, or use comprehensive examinations by optometrists. This report suggested that a combination of these approaches may be the best option.\textsuperscript{4}

### 4.1.1 Specific screening tests

Sensitivity, specificity, PPV, and negative predictive value (NPV) have been assessed for vision screening techniques in preschool children. Appendix 1 summarizes these studies. Screening tests were compared to a complete examination performed by an ophthalmologist, optometrist, or orthoptist (health professionals who assist ophthalmologists), to determine sensitivity and specificity. The studies that are listed in Appendix 1 are arranged by the type of screening test.

In one comparison, it was found that the positive predictive value (PPV) of photoscreening was 73\%, whereas the PPV for traditional screening methods (HOTV, Random-dot E) was 0\%.\textsuperscript{29} A computerized photoscreener was found to have a sensitivity of 94.6\% and a specificity of 90.1\%, compared to 85.7\% sensitivity and 81.0\% specificity for retinoscopy in a study comparing these methods among 300 children aged nine months to 50 months.\textsuperscript{30} This indicates that with the computer photoscreener, 5.4\% of patients will have a negative test result and a vision defect (false negative), and 9.9\% will test positive without a defect (false positive). As a result, 5.4\% of patients will be missed in screening and untreated, and 9.9\% of patients will be referred unnecessarily. In this study, retinoscopy had a 14.3\% false negative rate and a 19\% false positive rate. MTI Photoscreener sensitivity was higher (89\%) than in the examination of the Brückner reflex (64\%), in a case-control study of 10 patients with amblyogenic risk factors and six patients in the control group.\textsuperscript{31}

Numerous comparisons of vision screening tests for preschool children have been published. The Vision in Preschoolers (VIP) study group compared the sensitivities of 11 screening tests among 2,588 children aged three years to five years.\textsuperscript{32} Visual acuity tests included crowded Linear Lea Symbols and crowded linear HOTV. Tests for stereoacuity were the Random-dot E and Stereo Smile II. Retinoscopy, the Retinomax autorefractor, and the SureSight autorefractor were used to test refractive error. The sensitivity of photoscreeners (iScreen, MTI, Power Refractor II video-photorefraction) and the sensitivity of the cover-uncover test were also examined. Non-cycloplegic retinoscopy (NCR), SureSight, and Retinomax were the most sensitive tests for the detection of amblyopia, at 88\%, 80\%, and 78\% respectively, and 94\% specificity. For the detection of strabismus, the MTI
Photoscreener, cover-uncover test, Stereo Smile II test, SureSight, and Retinomax were most sensitive (65%, 60%, 58%, 54%, and 54% respectively). NCR, Retinomax, SureSight, and Lea Symbols were the most sensitive for the detection of significant refractive error (74%, 66%, 63%, and 58% respectively). Lea Symbols, Retinomax, and NCR were most sensitive for detecting reduced visual acuity, at 48%, 39%, and 38% respectively.\textsuperscript{32}

The VIP study group examined the sensitivities of these tests for the detection of any vision disorder. The most sensitive were NCR (64%), Retinomax (63%), SureSight Vision screener (63%), and Lea Symbols (61%). For discovering conditions that should be detected and treated early, NCR was the most sensitive at 90%, followed by Retinomax (88%), SureSight (81%), and Lea Symbols (77%). The specificities were all \( \geq 90\% \).\textsuperscript{33} Licensed eye-care professionals administered these tests.

A comparison of Lea Symbols, MTI Photoscreener, keratometry, and Retinomax was carried out in a sample of 379 preschool children for the detection of astigmatism.\textsuperscript{34} The sensitivity and specificity of the tests were 92% and 56% (Lea symbols), 60% and 86% (MTI photoscreening), 95% and 77% (keratometry), and 93% and 95% (Retinomax).\textsuperscript{34} Visual acuity screening using Lea Symbols was compared to noncycloplegic refraction for the detection of astigmatism in 245 children aged three years to five years.\textsuperscript{35} Lea Symbols had a sensitivity of 90% and a specificity of 44%, whereas refraction had a 91% sensitivity and an 86% specificity.\textsuperscript{35} The sensitivity and specificity of a digital Randodot test were compared to those of the Randot stereoaucity test, the Titmus test, and the Lang test. Similar results were found in preschool children.\textsuperscript{36} Randot Stereocards were found to produce reliable valid results in children up to 24 months old.\textsuperscript{37}

Two hand-held autorefractors (Retinomax and SureSight) were compared to cycloplegic retinoscopy (paralysis to minimize accommodation) in 35 children aged three years to five years.\textsuperscript{38} Moderate agreement was found between the two autorefractors, and between the autorefractors and cycloplegic retinoscopy. The authors concluded that these devices may be useful as screening tools.\textsuperscript{38} A limitation of this study is that these children were rendered cycloplegic before testing. A screening program would likely not use cycloplegia, hence these results may not be generalizable to a screening program.\textsuperscript{38} In another study, 43 children were tested before and after the induction of cycloplegia with autorefractors.\textsuperscript{39} The results of the testing with autorefractors when cycloplegia was not induced were inconsistent. The best device would give a more accurate reading and would not allow accommodation by the child being tested.\textsuperscript{39} Another study found that the Retinomax was a useful screening device for children 46 weeks to 81 weeks old. The accuracy of the results was affected by patients’ cooperation.\textsuperscript{40} Furthermore, these children were rendered cycloplegic.\textsuperscript{40} The results obtained by an ophthalmic nurse using a handheld autorefractor (Retinomax) were compared to those of traditional retinoscopy performed by an ophthalmologist in children under the age of six years.\textsuperscript{41} Results were similar using the methods with cycloplegia, indicating that Retinomax is useful for measuring refractive error. The results obtained without cycloplegia were inaccurate.\textsuperscript{41}

### 4.1.2 Screening effectiveness

A randomized controlled trial examining the effects of preschool vision screening was conducted in the UK.\textsuperscript{42} The methods of randomization and allocation concealment were inadequate, because the date of birth was used to determine which children were assigned to the intervention group. The control group was assessed with visual surveillance (eight- and 18-month examinations by health
visitors and family doctors using the cover test and observing visual behaviour), while those in the intervention group were examined by an orthoptist who tested visual acuity, ocular alignment, stereopsis, and non-cycloplegic refraction. The specificity was 92% for the control group and 95% for the intervention group (p<0.01). More children were found to have amblyopia in the intervention group (1.6%) compared with the control group (0.5%, p<0.01). In a comparison of the tests that were administered as part of the intervention, it was found that photorefraction was the most sensitive component.

An observational study designed to test the validity of a preschool vision screening program was conducted in Canada. Public health nurses administered tests for visual acuity, stereoacuity, and ocular alignment to >1,100 children per year, over three years. The results were compared to those from practitioners’ reports. The sensitivity ranged from 60.4% to 70.9%, and the specificity ranged from 69.6% to 79.9%. The PPV was 21.6% to 32.3%, and the NPV was 92.6% to 95.3%. This study concluded that based on the numbers of children detected with vision defects, the screening program is valid and should be continued.

A comparison of 808 eight-year-old children who had received screening for vision defects in infancy and 782 children who had not received screening found that the prevalence of amblyopia was higher in the children who were not screened (2.6%) compared with those who had received screening (1%) (p=0.0098). Ophthalmologists conducted the screening, which included a corneal reflex test, fixation-and-following test, ductions and versions examination, cover-uncover test, alternate cover test, and retinoscopy. The screening program had a sensitivity of 85.7%, a specificity of 98.6%, a PPV of 62.1%, and an NPV of 99.6%, indicating that it was effective.

A retrospective cohort study of 6,081 children assessed the visual outcomes of children aged 7.5 years who had received preschool vision screening or who had not received screening in the UK. More children were offered preschool vision screening (24.9%) than actually attended the screening (16.7%). Children who received vision screening had a 45% lower prevalence of amblyopia compared to those who did not receive screening. When all children who were offered the screening were included in the analysis, the effects of early detection on amblyopia outcome diminished. The authors concluded that the effectiveness of preschool vision screening had to be improved. This study indicated that although a vision screening program can be efficacious, its effectiveness is limited by the number of children who receive screening. The effectiveness of the screening program will be increased if a greater proportion of children who are offered vision screening receive it.

Screening of 3.5-year-old children by an orthoptist and the results of referrals to the hospital eye service were examined in a retrospective study of 6,794 children conducted in the UK. The screening test resulted in 5.1% of children referred to the hospital eye service, and detection of straight-eyed amblyopia and strabismic amblyopia. Upon treatment, 87.2% of children with straight-eyed amblyopia and 64.3% of children with strabismic amblyopia achieved an improvement in visual acuity. Another retrospective study that followed 3,126 children from birth to age 10 years found that vision screening is effective in detecting defects. Visual acuity was assessed at ages four years, 5.5 years, seven years, and 10 years by nurses at child health care centres or in the schools. Anetropia (any refractive error) was mainly detected at age four years. Most cases of strabismus were detected before the age of four years, while microtropia (small angle heterotropia) was detected at four years. The prevalence of amblyopia was reduced to 0.2% from 2% by screening, and most patients with amblyopia increased their visual acuity with treatment, indicating that
screening and treatment can reduce the prevalence of amblyopia.\textsuperscript{15}

An assessment of the vision screening program in Sweden found that tests (HOTV) for children four years and 5.5 years of age had a sensitivity of 92\% and a specificity of 97\%.\textsuperscript{16} This study found that screening and subsequent treatment decreased the prevalence of amblyopia.\textsuperscript{16}

4.1.3 Re-screening effectiveness

A retrospective study of 1,545 children in the UK examined the effects of a secondary screening program that was led by optometrists and orthoptists. Children were referred to the program by physicians, community medical officers, health visitors, school nurses, and primary orthoptic screeners who had conducted a screening protocol. This method of secondary screening that combines the skills of an optometrist and an orthoptist was found to be effective in detecting most children with vision problems, while avoiding unnecessary referrals to the hospital service.\textsuperscript{48} Upon secondary testing, 43\% of patients were found to be normal and did not require examination by the hospital eye service.\textsuperscript{48}

A preschool vision screening program that was developed by nursing students and faculty in the US assessed vision using Lea Symbols and the Random-dot E stereopsis test.\textsuperscript{49} This study screened 181 children aged three years to five years and found 5.5\% had abnormal results (28/181). Upon re-screening 20 of these children, it was found that nine had an abnormal result and were referred. This approach decreased the number of referrals to primary-care providers.\textsuperscript{49}

4.1.4 Personnel administering vision screening

A study of personnel using the MTI Photoscreener for children <3 years of age found that the sensitivity and specificity of the screening method were unaffected by the screeners’ ophthalmic knowledge. Screening was conducted by ophthalmologists, pediatricians, ophthalmic technicians, health department employees, Prevention of Blindness Society employees, and Lions Club volunteers.\textsuperscript{50} The AAP suggests that photoscreening may result in higher screening rates. Photoscreening will be useful for difficult-to-screen children, and the evaluator should be properly trained.\textsuperscript{51} The AAO recognizes that photoscreening and autorefractive devices may be useful for screening, but state that they should not replace current screening protocols.\textsuperscript{3}

An evaluation of screening programs in Sweden and Canada found that properly trained non-ophthalmologists can reliably screen four-year-old children. It was noted that children who fail screening tests should be referred to an ophthalmologist.\textsuperscript{23} HOTV is the most common test used in Canada.\textsuperscript{23} The VIP study compared the administration of vision screening tests among three- to five-year-olds by nurses and lay screeners. The tests administered were Retinomax, SureSight, crowded linear Lea Symbols, single Lea Symbols (administered by lay screeners only), and Stereo Smile II.\textsuperscript{24} A gold standard examination by an ophthalmologist or an optometrist was used for comparison. The single Lea Symbols test administered by lay screeners resulted in higher sensitivity compared to the crowded linear Lea Symbols administered by lay screeners or nurses. All other screening tests resulted in higher sensitivity when administered by the nurses compared to the lay screeners. Except in the case of the linear Lea Symbols, these differences were small and not statistically significant. This indicates that similar results can be achieved with nurses and lay screeners.\textsuperscript{24}

A UK study comparing vision screening conducted by orthoptists (1,582 children), health visitors (2,081 children), and clinical medical officers (1,701 children) found insufficient evidence to support an orthoptic-conducted preschool vision screening program.\textsuperscript{14} The
prevalence of amblyopia was similar in children who were screened by the examiners.14

A Canadian review of literature from 1983 to 1995 was conducted to assess screening for strabismus.52 This review found that low-risk children should be screened by a primary-care physician, whereas high-risk children (with low birth weight, a family history of strabismus, congenital ocular abnormality, or systemic conditions with vision-threatening ocular manifestations) should be examined by an ophthalmologist.52 A UK study of 2,041 children examined the effectiveness of preschool vision screening.53 This study found that screening by health visitors was as effective as screening by orthoptists.53 Lay screeners were used in a study using the MTI photoscreener for preschool vision screening.54 The PPV was 41.4% for astigmatism, 60.5% for anisometropia, and 84.2% for strabismus. In a study based in Taiwan, trained and certified kindergarten teachers effectively tested preschool children for amblyopia and strabismus.55

4.1.5 Testability

A more direct comparison in the VIP study found that HOTV scores were lower than Lea Symbols scores, likely because of the difficulty in testing younger children (three years) with HOTV numbers.56 Another study found similar results, with testability rates (i.e., children being able to complete the tests) that were higher for Lea Symbols than HOTV for vision screening of three-year-olds.11 Testability rates in yet another study were found to be similar for Lea Symbols and the HOTV chart in the testing of three-year-old children.57 Lea symbols have been shown to be effective in detecting amblyopia in children as young as 23 months.58 The VIP study also examined the testability of three Random-dot stereotests and found a significant difference.59 The testability of the Stereo Smile test was 91%, 81% for the Random-dot E test, and 71% for the Randot preschool test, when assessed in children aged three years to 3.5 years. The authors suggest that tests offering two choices for the children to pick from, such as the Stereo Smile test and the Random-dot E test, offer higher testability.59 Another study found that successful screening in four-year-olds was higher (88% to 98%) compared to three-year-olds (70% to 90%). It suggested that there is insufficient evidence showing that the detection of amblyopia at three years of age results in better treatment outcomes than detection at four years of age.7 Overall, testability is higher in older children, because screening in younger children is more difficult.

4.1.6 Limitation of Studies

There is a lack of rigorous controlled studies examining the effectiveness of preschool vision screening. One randomized controlled study that was identified was not of high quality, so its findings should be viewed accordingly.42 The other types of studies reviewed are observational studies or non-randomized controlled studies.

Another limitation relates to the population screened. Some studies are screening children from a head-start program (which serves children from low-income families24,32-34) who would not reflect the general population. One study includes children from a clinical practice who may have had underlying conditions.60 Another limitation is the small size of the sample undergoing screening that was used in some studies.29,31 The screening age can also be a limitation in some studies. The screening of children <4 years of age can be difficult, and the screening results for this age group will be difficult to apply to an older population.54,60

Agreement in the literature regarding the type of test to use for preschool vision screening is lacking. Most studies measure the reduction in amblyopia as the outcome, but measures such as school performance may be a more relevant outcome to consider. In addition, no long-term studies exist, making it difficult to determine how these screening programs may affect future outcomes for these children.
One limitation of this review is the fact that it was conducted by one reviewer, because of the need to provide this report in a short timeframe. This may introduce bias in terms of selection of studies and interpretation of results. This report, however, was reviewed by several external clinical experts. In addition, the literature search was limited to the English language and excluded studies published before 1996. In some of the studies that are cited, orthoptists administered the vision-screening tests to preschool children. Orthoptists are common in areas such as the UK, but there is a lack of orthoptists in Canada. The generalizability of these findings to a Canadian setting is therefore questionable.

4.2 Current Practice

Throughout Canada, there exist a variety of programs for vision screening in children. Table 1 describes the current preschool vision screening programs in some provinces.

4.3 Professional Organizations

The CPS recommends an examination of external eye structures, red reflex, signs of posterior eye disease, and corneal light reflex for newborn children to age three months. At ages six months to 12 months, ocular alignment, ocular fixation, and following should be tested. Visual acuity testing should be completed for children aged three years to five years. These recommendations are based on those of the AAO. The Canadian Task Force on Preventive Health Care also recommends the visual acuity testing of preschoolers.

The American Public Health Association recommends a comprehensive eye examination for preschool children at ages six months, two years, and four years. The USPSTF assessed vision screening for children under the age of five years and found fair evidence to suggest that screening tests could detect amblyopia, strabismus, and refractive error with “reasonable accuracy.”

Amblyogenic risk factors can be detected by photoscreening in children up to the age of three years. Stereopsis (with Random-dot E, Titmus Fly Stereotest), visual acuity (HOTV, Lea Symbols, Tumbling E) can be used in children >3 years of age. No direct evidence was found to suggest that screening improves visual acuity in preschool children. Furthermore, the USPSTF did not find enough evidence to suggest an optimal screening test. The task force found fair evidence that visual acuity can be improved by the early detection of amblyogenic risk factors. These recommendations are “B level,” which indicates that fair evidence was found that the outcomes of preschool vision screening can outweigh the harms, and that this service should be provided.

The AOA’s recommendations for the screening of infants and toddlers include visual acuity (fixation preference tests, preferential looking visual acuity test), refraction (cycloplegic retinoscopy, near retinoscopy), binocular vision, and ocular motility (cover test, Hirschberg test, Brückner test, versions, near point of convergence). Tests for preschool children should include visual acuity (Lea symbols, Broken Wheel Acuity cards, HOTV), refraction (static retinoscopy, cycloplegic retinoscopy), binocular vision, accommodation, and ocular motility (cover test, positive and negative fusional vergences, near point of convergence, stereopsis, monocular estimate method retinoscopy, versions).
Table 1*: Preschool vision screening programs in Canada

<table>
<thead>
<tr>
<th>Province</th>
<th>Current Practice</th>
<th>Tests</th>
</tr>
</thead>
<tbody>
<tr>
<td>New Brunswick (Claudette Landry, NB Department of Health, Fredericton: personal communication, 2006 July 11)</td>
<td>public health nurses conduct vision screening for children at age 3.5 years</td>
<td>visual inspection of eyes, Randot, HOTV</td>
</tr>
<tr>
<td>Prince Edward Island (Aaron Campbell, Medical Programs Coordinator, Dept. of Health, Charlottetown: personal communication, 2006 September 12)</td>
<td>public health nurses conduct vision screening for children at birth, 2 months, 4 months, 6 months, 12 months, 15 months, 18 months, 4 years to 4.5 years</td>
<td>Frisby stereotest and Lea Symbols</td>
</tr>
<tr>
<td>Newfoundland</td>
<td>public health nurses, children aged 3 years</td>
<td>Sheridan-Gardner, cover-uncover, corneal light reflexes</td>
</tr>
<tr>
<td>Nova Scotia</td>
<td>public health nurses, children ages 4.5 years to 5.5 years</td>
<td></td>
</tr>
<tr>
<td>Quebec</td>
<td>no information provided</td>
<td></td>
</tr>
<tr>
<td>Ontario</td>
<td>screening done by primary care providers as part of 18-month “Well Baby” visit, testing repeated at ages 2 years to 3 years, 4 years to 5 years</td>
<td>red reflex, corneal light reflex, cover-uncover test</td>
</tr>
<tr>
<td>Manitoba</td>
<td>no information provided</td>
<td></td>
</tr>
<tr>
<td>Saskatchewan</td>
<td>no province-wide preschool vision screening program</td>
<td></td>
</tr>
<tr>
<td>Alberta</td>
<td>“Eye see…Eye learn” program provides examinations conducted by optometrist for children before they start school</td>
<td></td>
</tr>
<tr>
<td>British Columbia</td>
<td>no province-wide preschool vision screening program</td>
<td></td>
</tr>
<tr>
<td>Nunavut (Terry Creagh, Department of Health and Social Services, Government of Nunavut, Iqaluit: personal communication, 2006 Aug 21)</td>
<td>screening done primarily by public health nurses, community health nurses, community health representatives for children aged 4 years to 6 years</td>
<td>light reflex and visual acuity</td>
</tr>
<tr>
<td>Yukon</td>
<td>no information provided</td>
<td></td>
</tr>
<tr>
<td>Northwest Territories</td>
<td>conducted by public health nurses</td>
<td>illiterate E test or symbol chart, stereoscopic fly, corneal light reflex, cover-uncover</td>
</tr>
</tbody>
</table>

*Preschool vision screening programs may not be universally conducted across the province. Additional vision screening programs may be available in some regions through the work of charitable organizations.

Recommendations from the AAO for newborn to age three months include red reflex and inspection. Infants aged three months to six months should be tested for their ability to fix and follow, for red reflex, and for inspection. Children aged six months to 12 months should undergo the same tests; testing should also be done for alternate occlusion and corneal light reflex. Visual acuity, corneal light reflex, cover-uncover, red reflex, and inspection should be performed on children aged three years. These tests should be repeated at five years of age. If abnormalities are detected by screening, a comprehensive medical eye examination should be performed. These recommendations are based on the interpretations of an expert panel based on the best available scientific data.
The AAP recommends ocular history, vision assessment, external inspection of the eyes and lids, ocular motility assessment (corneal reflex test, cross cover test, Random-dot E test), pupil examination, and red reflex examination for children up to the age of three years. In addition, age-appropriate visual acuity tests (Lea Symbols, Allen cards, Snellen letters and numbers, Tumbling E, and HOTV) and ophthalmoscopy are recommended for children aged three years to five years.\textsuperscript{17,18}

4.4 Cost Effectiveness of Preschool Vision Screening

A US study was developed to design and test a cost-efficient community-based preschool vision screening program for a population of native American children among whom there was a high prevalence of astigmatism. Autorefraction was found to be the most expensive test, but it had the greatest specificity.\textsuperscript{64} An economic evaluation, based on a field study of 121 German kindergartens, found that vision screening in kindergartens by orthoptists costs approximately €924 for each case of amblyopia detected.\textsuperscript{65} Another study, conducted by the same group, but using a decision analytic model, calculated the cost-effectiveness ratio to be €727 for each case of amblyopia detected.\textsuperscript{66} The cost-effectiveness ratio was found to be influenced by the prevalence rate of the target condition and by test specificity. In the decision analytic model, one orthoptic examination costs €7.87 compared to €36.40 for an examination by an ophthalmologist.\textsuperscript{66} In Canada, there is a lack of orthoptists, so orthoptic screening is unlikely to be an option. Thus, the costs of screening described in these studies are unlikely to be generalizable to the Canadian setting.

The same group of German investigators constructed another decision analytic model to calculate the cost per detected case of amblyopia for five methods of screening among three-year-olds.\textsuperscript{67} In this German scenario, the least expensive option was visual acuity testing with re-screening for children who had inconclusive results, at an average cost of €878 per detected case. The most expensive option was the combination test of visual acuity, cover test, examination of eye motility, and direct referral to an ophthalmologist or to re-screening for inconclusive results.\textsuperscript{67} In yet another German study, a different group of investigators used a decision analytic model and found that the most cost-effective screening strategy for the detection of amblyopia or its risk factors is screening by an ophthalmologist for all children before the age of one year.\textsuperscript{68} This analysis, however, suggested that all four screening scenarios left a significant proportion of undetected children.

A cost-benefit analysis of vision screening methods for preschool and school-aged children found the benefit-to-cost ratio for screening programs was >1.0, indicating that the benefits of screening outweighed the costs.\textsuperscript{69} Another study found that, in 2005, comprehensive eye examinations conducted by an ophthalmologist or optometrist compared to usual care would cost US$12,985 per quality-adjusted life-year (QALY); the authors concluded that it was cost effective.\textsuperscript{70} This may be optimistic because the authors used the cost of monocular blindness to define costs, and not all children with amblyopia will develop monocular blindness. This same study found that when compared to vision screenings, universal eye examinations were cost-effective at US$18,390 per QALY. The cost-effectiveness of the treatment of amblyopia in 2005 was US$1,800 per QALY.\textsuperscript{70} In a separate study, the authors suggest that decision makers should consider orthoptic screening based on the ICER.\textsuperscript{71} Again, in Canada, this strategy would be unfeasible because of the lack of orthoptists to conduct screening.
5 CONCLUSIONS AND IMPLICATIONS

Preschool vision screening aims to detect vision disorders at an early age, with the assumption that early detection leads to earlier treatment and an improvement in outcomes. According to the literature examined in this review, the specific test that is used, the age and underlying health of the children, and the personnel administering the test all influence the effectiveness of preschool vision screening. Some studies suggest that screening should only be conducted by a trained eye-care professional, whereas others state that physicians, nurses, and lay screeners can effectively conduct preschool vision screening. There does not seem to be a consensus as to who should be administering the preschool vision screening test.

The effectiveness of the tests for preschool vision screening varies. For example, with photoscreening, sensitivities ranged from 27.8% to 88%, and specificities ranged from 40% to 98.5% in different studies (Appendix 1). Other preschool vision screening tests have variable sensitivities and specificities. In addition, professional organizations recommend different tests to assess vision in different age groups of preschool children. Although many studies have been published examining different tests to detect vision defects in preschool children, no single test or group of tests has been shown to be superior for preschool vision screening. While no Canadian studies have assessed the cost effectiveness of preschool vision screening, the studies suggest that universal eye examinations for preschool children have a low cost per QALY.

A preschool vision screening program meets most of the general criteria to consider when assessing a screening program. Although the prevalence of vision defects is low, they do cause disability. An earlier age of detection is a benefit in the treatment of amblyopia. No studies showed harms associated with screening. The tests can detect the defects that they are meant to detect, and there are effective treatments for these defects. Additional research is needed to ascertain the utility of preschool vision screening in the Canadian context. Such research should aim to determine who can conduct such tests, what tests should be used for what ages of children, and whether one- or two-stage screening is most effective from a clinical and a cost viewpoint.

6 REFERENCES


17. Eye examination in infants, children, and young adults by pediatricians: organizational principles to guide and define the child health care system and/or improve the health of all children. *Ophthalomology* 2003;110(4):860-5.


## Appendix 1: Sensitivity, specificity, PPV, and NPV for pre-school vision screening tests

<table>
<thead>
<tr>
<th>Study</th>
<th>Number of Children Tested</th>
<th>Age of Children</th>
<th>Screening Test</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>PPV</th>
<th>NPV</th>
<th>Administered By</th>
</tr>
</thead>
<tbody>
<tr>
<td>Donahue et al.</td>
<td>&gt;400,000</td>
<td>preschool age</td>
<td>MTI Photoscreener</td>
<td>NR</td>
<td>NR</td>
<td>80%</td>
<td>NR</td>
<td>technicians, orthoptists, ophthalmologists, optometrists, medical students, program coordinators</td>
</tr>
<tr>
<td>Berry et al.</td>
<td>949</td>
<td>6 months to 59 months</td>
<td>MTI Photoscreener</td>
<td>50.0%</td>
<td>98.5%</td>
<td>57.5%</td>
<td>94.4%</td>
<td>NS</td>
</tr>
<tr>
<td>Tong et al.</td>
<td>392</td>
<td>&lt;4 years</td>
<td>MTI Photoscreener</td>
<td>65%</td>
<td>87%</td>
<td>NR</td>
<td>NR</td>
<td>ophthalmologist, ophthalmic medical technologist</td>
</tr>
<tr>
<td>Weinand et al.</td>
<td>112</td>
<td>6 months to 48 months</td>
<td>MTI Photoscreener</td>
<td>82.8%</td>
<td>61.8%</td>
<td>68.2%</td>
<td>48.1%</td>
<td>pediatrician, orthoptist, ophthalmologist</td>
</tr>
<tr>
<td>Tong et al.</td>
<td>100</td>
<td>&lt;3 years</td>
<td>MTI Photoscreener</td>
<td>37% to 88%</td>
<td>40% to 88%</td>
<td>NR</td>
<td>NR</td>
<td>ophthalmologist, pediatrician, ophthalmic technician, health department employee, Prevention of Blindness Society employee, Lions Club volunteer</td>
</tr>
<tr>
<td>Cooper et al.</td>
<td>105</td>
<td>12 months to 44 months</td>
<td>Fortune Photoscreener,</td>
<td>Fortune 60%</td>
<td>82%, 90%</td>
<td>NR</td>
<td>NR</td>
<td>ophthalmologist</td>
</tr>
<tr>
<td>Cooper et al.</td>
<td>113</td>
<td>11 months to 44 months</td>
<td>Otago and Dortmans Photoscreeners</td>
<td>70%, 70%</td>
<td>75% to 86%</td>
<td>NR</td>
<td>NR</td>
<td>NS</td>
</tr>
<tr>
<td>Fern et al.</td>
<td>220</td>
<td>&lt;6 years</td>
<td>Photoscreener</td>
<td>27.8% to 47.0%</td>
<td>91%</td>
<td>66.7% to 94.4%</td>
<td>NR</td>
<td>NR</td>
</tr>
<tr>
<td>Pott et al.</td>
<td>196</td>
<td>5 years</td>
<td>Polaroid suppression test</td>
<td>60%</td>
<td>91%</td>
<td>NR</td>
<td>NR</td>
<td>NS</td>
</tr>
<tr>
<td>Birch et al.</td>
<td>1,260</td>
<td>2 years to 5 years</td>
<td>Random-dot stereoaucity</td>
<td>91%</td>
<td>91%</td>
<td>NR</td>
<td>NR</td>
<td>NS</td>
</tr>
<tr>
<td>Rasmussen et al.</td>
<td>400</td>
<td>3 years</td>
<td>Lang II Random-dot stereotest</td>
<td>33%</td>
<td>85%</td>
<td>9.5%</td>
<td>96%</td>
<td>nurses</td>
</tr>
<tr>
<td>Simons et al.</td>
<td>112</td>
<td>3 years to 5 years</td>
<td>STRDS and STBS test</td>
<td>NR</td>
<td>STRDS 80%; STBS 96%</td>
<td>NR</td>
<td>NR</td>
<td>NS</td>
</tr>
<tr>
<td>Lim et al.</td>
<td>894</td>
<td>3 years to 5 years</td>
<td>home vision screening (visual acuity testing with picture cards)</td>
<td>NR</td>
<td>NR</td>
<td>77%</td>
<td>NR</td>
<td>NS</td>
</tr>
<tr>
<td>Barry et al.</td>
<td>1,180</td>
<td>3 years</td>
<td>cover test, Lea single optotype</td>
<td>90.9%</td>
<td>93.8%</td>
<td>NR</td>
<td>NR</td>
<td>orthoptist</td>
</tr>
</tbody>
</table>
## Appendix 1: Sensitivity, specificity, PPV, and NPV for pre-school vision screening tests

<table>
<thead>
<tr>
<th>Study</th>
<th>Number of Children Tested</th>
<th>Age of Children</th>
<th>Screening Test</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>PPV</th>
<th>NPV</th>
<th>Administered By</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cordonnier et al. (^{83})</td>
<td>1,218</td>
<td>9 months to 36 months</td>
<td>Retinomax</td>
<td>37% to 87%</td>
<td>93% to 99%</td>
<td>19% to 69%</td>
<td>96% to 100%</td>
<td>orthoptist</td>
</tr>
<tr>
<td>Barry et al. (^{85})</td>
<td>404</td>
<td>3 years</td>
<td>Retinomax–non-cycloplegic retinoscopy</td>
<td>80%</td>
<td>58%</td>
<td>NR</td>
<td>NR</td>
<td>orthoptist</td>
</tr>
<tr>
<td>Cordonnier et al. (^{86})</td>
<td>1,205</td>
<td>9 months to 36 months</td>
<td>Retinomax</td>
<td>51% to 84%</td>
<td>90% to 98%</td>
<td>58% to 84%</td>
<td>90% to 96%</td>
<td>orthoptist</td>
</tr>
<tr>
<td>Cordonnier et al. (^{87})</td>
<td>897</td>
<td>9 months to 36 months</td>
<td>Retinomax</td>
<td>70.2% to 78.7%</td>
<td>79.2% to 94.6%</td>
<td>51.4% to 78.6%</td>
<td>91.7% to 93.0%</td>
<td>orthoptist</td>
</tr>
<tr>
<td>Newman et al. (^{13})</td>
<td>936</td>
<td>3.5 years</td>
<td>Sheridan-Gardiner single optotype test, cover test, ocular movement, 20(^\circ) prism test, TNO stereotest</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>100% (99.6% Sheridan Gardiner test alone)</td>
<td>orthoptist</td>
</tr>
<tr>
<td>Chui et al. (^{58})</td>
<td>178</td>
<td>3 years to 4 years</td>
<td>Lea Symbols, Frisby plates</td>
<td>50% to 75%</td>
<td>68% to 95%</td>
<td>NR</td>
<td>90% to 96%</td>
<td>public health nurses</td>
</tr>
<tr>
<td>Briscoe et al. (^{88})</td>
<td>292</td>
<td>4 years to 6 years</td>
<td>vision screening computer program</td>
<td>50%</td>
<td>98.9%</td>
<td>63%</td>
<td>NR</td>
<td>non-trained personnel</td>
</tr>
<tr>
<td>Simon et al. (^{89})</td>
<td>122</td>
<td>6 months to 5 years</td>
<td>visual evoked potential system</td>
<td>97.3%</td>
<td>80.8%</td>
<td>70.6%</td>
<td>98.4%</td>
<td>NS</td>
</tr>
<tr>
<td>Atilla et al. (^{90})</td>
<td>89</td>
<td>&lt;4 years</td>
<td>fix-follow-maintain</td>
<td>53.1%</td>
<td>38.5%</td>
<td>32.6%</td>
<td>NR</td>
<td>NS</td>
</tr>
<tr>
<td>Kemper et al. (^{91})</td>
<td>170</td>
<td>&lt;5 years</td>
<td>SureSight autorefractor</td>
<td>80% to 88%</td>
<td>41% to 58%</td>
<td>NR</td>
<td>NR</td>
<td>study investigator or ophthalmologist</td>
</tr>
<tr>
<td>Büchner et al. (^{92})</td>
<td>336</td>
<td>3.5 years to 4.5 years</td>
<td>SureSight autorefractor</td>
<td>41% to 95%</td>
<td>73% to 92%</td>
<td>NR</td>
<td>NR</td>
<td>NS</td>
</tr>
</tbody>
</table>

NR=not reported; NS=not specified; NPV=negative predictive value; PPV=positive predictive value; STRDS=small target Random-dot Stereogram; STBS=small target binocular suppression.

*Likelihood ratio of positive test=sensitivity/(1−specificity); likelihood ratio of negative test=(1−sensitivity)/specificity.