

Cost-Utility Analysis of Orthoptic Screening in Kindergarten: A Markov Model Based on Data From Germany

Hans-Helmut König, MD, MPH*, and Jean-Cyriaque Barry, MD‡

ABSTRACT. *Objective.* To estimate the long-term cost-effectiveness of a hypothetical screening program for untreated amblyopia in 3-year-old children conducted by orthoptists in all German kindergartens in the year 2000.

Methods. A cost-utility analysis was performed for which a decision tree was combined with a Markov model. Incremental costs and effects during the children's remaining lifetime were estimated. The model took into account the probability of treatment without screening, age-specific treatment success rates, costs of screening and treatment, as well as effects of unilateral and bilateral visual impairment caused by amblyopia and other eye diseases coming along later in life on quality of life (utility). Model parameter values were obtained from a field study of orthoptic screening in kindergarten, from the literature, and from expert interviews. Costs were estimated from a third-party payer perspective. Uncertainty was assessed by univariate and probabilistic sensitivity analysis (Monte Carlo simulation).

Results. The incremental cost-effectiveness ratio (ICER) of orthoptic screening was 7397 Euro (€) per quality-adjusted life year (QALY) when costs and effects were discounted at 5%. In univariate sensitivity analysis, the ICER was sensitive to the uncertainty regarding the utility of unilateral visual impairment and to the discount rate for effects; besides uncertainty regarding the prevalence of untreated amblyopia, the odds ratio of success of treatment when started late, and the probability of treatment without screening had a noticeable but much smaller effect. Monte Carlo simulation yielded a 90% uncertainty interval for the ICER of 3452 €/QALY to 72 637 €/QALY; the probability of an ICER <25 000 €/QALY was 84%.

Conclusions. The ICER of orthoptic screening seems to fall within a range that warrants careful consideration by decision-makers. Much of the uncertainty in results comes from the uncertainty regarding the effect of amblyopia on quality of life. To reduce this uncertainty, the impact of amblyopia on utility should be investigated. *Pediatrics* 2004;113:e95–e108. URL: <http://www.pediatrics.org/cgi/content/full/113/2/e95>; *amblyopia, vision screening, children, preschool, cost-effectiveness, decision modeling, Markov process.*

ABBREVIATIONS. VA, visual acuity; QALY, quality-adjusted life year; ICER, incremental cost-effectiveness ratio; CI, confidence interval; OR, odds ratio; BMES, Australian Blue Mountain Eye Study.

Preschool vision screening mainly aims at preventing amblyopia by early detection and subsequent treatment of amblyogenic factors such as strabismus and certain refractive errors.^{1–3} Recently, there has been concern about the lack of scientific data on the effectiveness of preschool vision-screening programs^{4–6}, and rigorous evaluation has been called for.^{2,5,7,8}

Because of increasing health expenditure, there is a growing interest among decision-makers in the cost-effectiveness of medical interventions.⁹ Many argue that the cost-effectiveness of a service has to be demonstrated before it may be covered by social health insurance or national health service programs. However, little is known about the cost-effectiveness of childhood vision-screening programs.

In a recent field study, we analyzed the effectiveness and cost-effectiveness of vision screening performed by orthoptists in German kindergartens.^{10–12} Orthoptists are specialized medical aides with thorough training in infant and child vision assessment, ocular motility-disorder examinations, and amblyopia management. In this study, screening was performed at age 3 because, in Germany, children may attend kindergarten from age 3 onward and the fourth year of life was considered best for vision screening.^{13,14} From age 3 onward, in most children monocular visual acuity (VA) can be assessed reliably by simple screening methods,¹⁵ and there is evidence that treatment is most effective when administered as early as possible.^{16–18}

In the field study, the clinical endpoint of analysis was the ophthalmologic diagnosis, with the number of newly detected cases of amblyopia being the measure of effects. However, diagnosis of amblyopia is only an intermediate endpoint, especially with respect to the cost-effectiveness of screening. A more comprehensive analysis should also take into account the costs and effects of subsequent treatment and the effect of a visual deficit on quality of life throughout the remaining lifetime. Such an analysis would require either many years of follow up or the use of economic modeling. Modeling allows one to combine different sources of evidence, to transfer study results to different contexts and extrapolate them to the longer term, and to analyze uncertainty of results in detail.^{19–21}

From the *Health Economics Research Unit, Department of Psychiatry, University of Leipzig, Leipzig, Germany; and ‡Department of Ophthalmology II, University Eye Hospital Tübingen, Tübingen, Germany.

Received for publication May 15, 2003; accepted Oct 20, 2003.

Address correspondence to Hans-Helmut König, MD, MPH, University of Leipzig, Health Economics Research Unit, Department of Psychiatry, Johannisallee 20, D-04317 Leipzig, Germany. E-mail: koenig@aya.yale.edu
PEDIATRICS (ISSN 0031 4005). Copyright © 2004 by the American Academy of Pediatrics.

Building on the results of the field study, the purpose of this analysis was to model the long-term cost-effectiveness of a hypothetical orthoptic screening program for children at age 3 conducted in all German kindergartens in the year 2000. Incremental costs and effects that would occur during the children's remaining lifetime were estimated. The uncertainty inherent in the model was analyzed in detail to assess the precision of results and to identify those variables with the greatest potential impact on cost-effectiveness.

METHODS

This section is organized as follows: First, the study design is described in general and then the decision-analytic model is explained in detail, followed by the description of the model parameters used.

Study Design

Based on a decision-analytic model, a cost-utility analysis was performed in which quality-adjusted life years (QALYs) were used as the measure of effects. QALYs are calculated by weighting the duration of health states by a preference-based score of health-related quality of life (utility), measured on a scale from 0 (dead) to 1 (perfect health).²²

In the model, the strategy "orthoptic screening at age 3 in kindergarten" ("screening") was compared with the strategy "usual care." Although usual care involves no orthoptic screening in kindergarten, amblyopia still may be detected when children see an ophthalmologist with or without referral (eg, by their pediatrician). The main outcome measure was the incremental cost-effectiveness ratio (ICER), ie, the ratio of the difference in mean costs \bar{C} and the difference in mean effects \bar{E} between the screening and usual-care strategies:

$$ICER = \frac{\bar{C}_{screening} - \bar{C}_{usual_care}}{\bar{E}_{screening} - \bar{E}_{usual_care}} = \frac{\Delta\bar{C}}{\Delta\bar{E}} \quad (1)$$

Hence, for the comparison, a purely incremental approach was chosen, ie, only the differences in costs and QALYs between the 2 strategies were considered. This means that, when calculating QALYs, only health states were considered with utility losses set off by

- unilateral visual impairment caused by a not successfully treated target disease; or
- visual impairment caused by any other unilateral eye disease coming along later in life. In a proportion of patients already affected by unilateral visual impairment caused by a target disease, this would cause bilateral visual impairment; in all other persons, this would result in unilateral visual impairment.

Visual impairment caused by any bilateral eye disease was not considered in the model, because it would cause bilateral visual impairment in both patients with the target disease as well as persons without and hence would not set off differences in QALYs between the strategies compared.

For the model, a threshold for the presence of visual impairment had to be defined. In accordance with various therapeutic,^{16,17} epidemiologic,^{23,24} and disability studies,²⁵ this threshold was set at a corrected VA of <0.5 (20/40): Monocular visual impairment was defined as VA <0.5 in the worse eye and ≥ 0.5 in the better eye, and bilateral visual impairment was defined as VA <0.5 in both eyes.

The target disease was defined as untreated unilateral amblyopia, which, without treatment, would cause a lasting visual impairment.

Decision-Analytic Model

A decision tree was combined with a Markov model. Analysis was performed by using the DATA software package (version 4.0, TreeAge Software, Inc, Williamstown, MA).

Decision Tree

The decision tree (Fig 1) has 1 decision node that distinguishes the strategies screening and usual care. In the screening strategy, the model takes into account, by respective chance nodes, that only a proportion of children attend kindergarten at age 3 (labeled

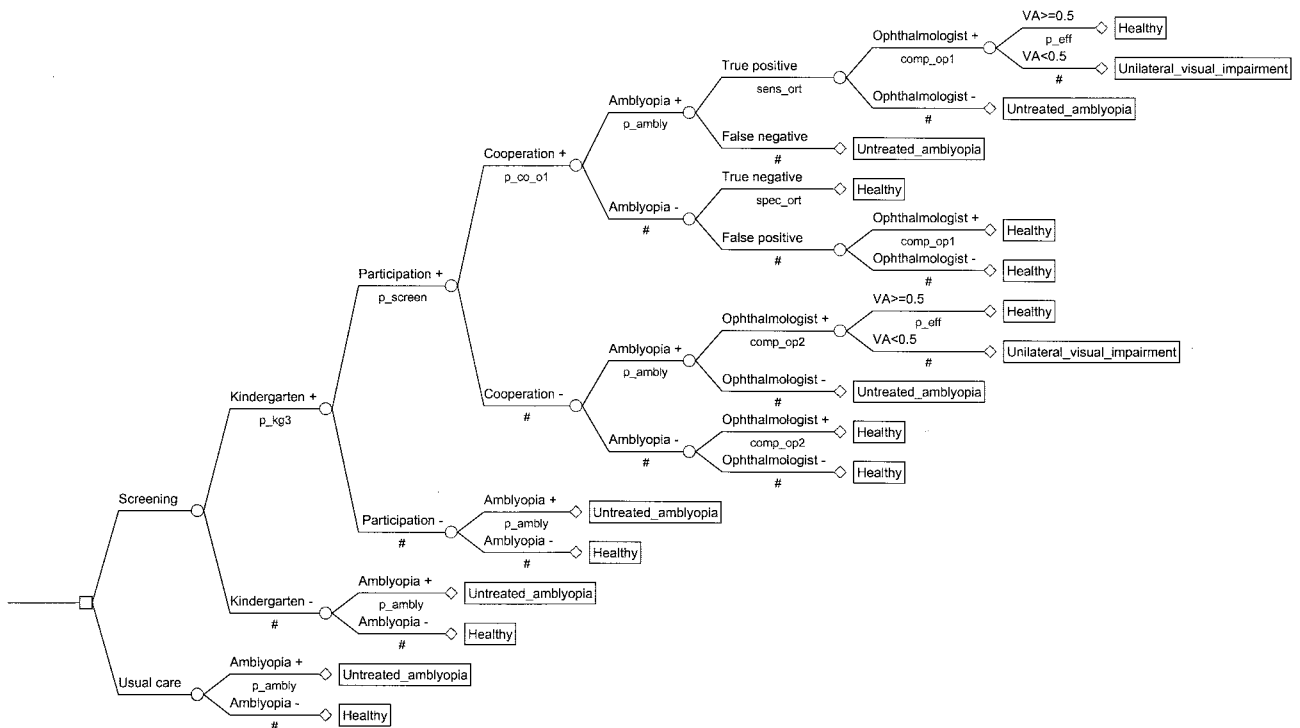


Fig 1. Decision tree for comparison of screening strategy with usual-care strategy. □, decision node; ○, chance node; ◇, end node of decision tree and initial health state of respective Markov process; below the branches of the decision tree, the labels of model parameters representing probabilities (proportions) are stated (see Table 1); #, 1 minus probability of other branch; +, yes; -, no.

“kindergarten +/-”) and would be present and participate in screening on the scheduled examination day (participation +/-). It also takes into account that a proportion of participating children would not cooperate sufficiently with the screening examination (cooperation +/-). Children with a positive screening result as well as noncooperative children would be referred to an ophthalmologist. At this point, the model takes into account that a proportion of parents would not comply with the referral (ophthalmologist +/-). Children in whom amblyopia was newly diagnosed by the ophthalmologist would be treated and, by a certain probability, successfully achieve VA ≥ 0.5 in the amblyopic eye. Therefore, at the end nodes of the screening strategy, 3 different health states can be reached:

1. “Healthy”: if a target disease was not present or was treated successfully.
2. “Unilateral visual impairment (amblyopia)”: if a target disease was detected by the screening but treatment was not successful.
3. “Untreated amblyopia”: if a target disease was not treated yet.

At the end nodes of the usual-care strategy, only the healthy (if no target disease was present) or untreated-amblyopia (if a target disease was present) health states can be reached.

Markov Model

At each end node of the decision tree, a Markov process starts, ie, the end node of the decision tree corresponds to the initial health state of a Markov process. In a Markov process the course of a disease is divided into distinct states, and transition probabilities are assigned for movement between them over a discrete time period called the Markov cycle.²⁶ By attaching estimates of resource use and health effects to the states and transitions in the

model and then running the model over a large number of cycles, long-term costs and effects can be estimated.

According to the 3 different initial health states, 3 Markov processes (A–C) were distinguished. Markov process A starts with the initial untreated-amblyopia health state (Fig 2), which corresponds, eg, to the third end node from the top of the decision tree displayed in Fig 1. Besides, there are 5 additional health states defined in this Markov process. Transition to the healthy state takes place when the child is seen by an ophthalmologist and is treated successfully before age 11; transition to the unilateral-visual-impairment (amblyopia) health state occurs if the child is not treated successfully before age 11. From the healthy state, transition to the unilateral-visual-impairment health state takes place when a unilateral nonamblyopic eye disease comes along and causes unilateral visual impairment. From the unilateral-visual-impairment (amblyopia) health state, transition to the bilateral-visual-impairment (with amblyopia) health state occurs when the nonamblyopic eye of an amblyopic patient is affected by a unilateral eye disease, resulting in bilateral visual impairment. The state “death” can be reached from all other health states.

Markov processes B and C are subsets of Markov process A. Markov process B starts with the initial healthy state, and Markov process C starts with the initial unilateral-visual-impairment (amblyopia) health state. Possible transitions correspond to those described within Markov process A.

In the model presented here, the cycle length was 1 year. Simulation started in cycle 0, which corresponds to the 4th year of life (age 3), and was completed in cycle 86, which corresponds to the 90th year of life. To account for gender differences in mortality and incidence of visual impairment, all Markov processes were made gender specific.

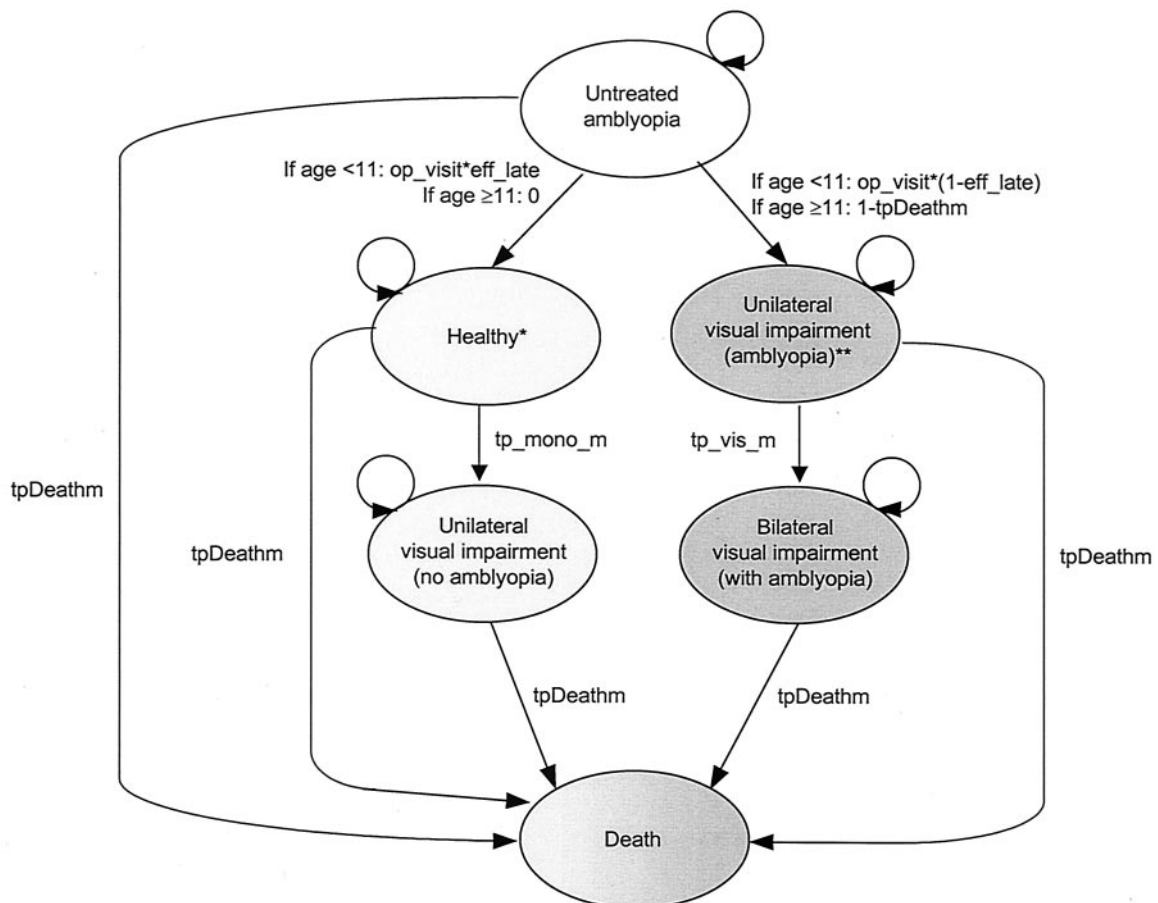


Fig 2. Markov process A for course of disease with an initial untreated-amblyopia health state. Health states are represented by ovals, and possible transitions between those states are shown by arrows. Variable names adjacent to the arrows are the transition probabilities of the model (see Table 1). Variable names refer to men; for women tp_mono_m , tp_vis_m , and $tpDeathm$ are replaced by tp_mono_f , tp_vis_f , and $tpDeathf$, respectively. *, Initial health state of Markov process B (shaded in bright gray); **, initial health state of Markov process C (shaded in dark gray).

TABLE 1. Parameter Values for Decision Tree and Markov Model With Range for Univariate Sensitivity Analysis and Distribution of Monte Carlo Simulation

Model Parameter	Source*	Name Used in Decision Tree or Markov Model	Parameter Value (Base Value)	Range Used for Univariate Sensitivity Analysis	Distribution Used in Monte Carlo Simulation
Screening Population					
Proportion of 3-year-olds in kindergarten	L	p_kg3	0.563	(None)	(None)
Proportion male	L	p_male	0.513	(None)	(None)
Participation in screening program	L	p_screen	0.800	0.800–1.000	Beta (30.22; 3.36)†
Prevalence of target disease	F	p_ambly	0.022	0.010–0.050	Beta (27; 1155)†
Compliance with referral to ophthalmologist					
After positive screening	F	comp_op1	0.972	0.935–0.990	Beta (107; 4)†
After noncooperative screening	L	comp_op2	0.900	0.800–1.000	Uniform (0.8; 1.0)†
Test Characteristics of Orthoptic Screening					
Sensitivity	F	sens_ort	0.909	0.772–0.971	Beta (21; 3)†
Specificity	F	spec_ort	0.924	0.907–0.940	Beta (898; 75)†
Cooperation	F	p_co_o1	0.887	0.869–0.905	Beta (1048; 134)†
Effectiveness of Treatment					
Probability of success of treatment started at age 3	L	p_eff	0.750	0.600–0.900	Beta (23.26; 7.75)†
Utilities					
Healthy	L	u_health	1.000	(None)	(None)
Unilateral visual impairment	L	u_mono	0.960	0.920–1.000	Uniform (0.92; 1.00)†
Bilateral visual impairment	L	u_impair	0.780	0.710–0.850	Beta (104.14; 29.38)†
Costs					
Organization per screening examination, €	F	c_org	5.24	2.62–7.86	Normal (5.24 × 0.9/p_screen; 1.34 × 0.9/p_screen)§
Orthoptic screening examination excluding organization, €					
Fixed costs, €	F	c_fix	4.18	3.28–5.08	Normal (4.18 × 0.9/p_screen; 0.46 × [0.9/p_screen] ^{1.5})§
Variable costs, €	F	c_var	3.58	3.27–3.89	Normal (3.58; 0.16)§
Ophthalmologic examination, €	F	c_op	36.40	27.30–45.50	Normal (36.40; 4.64)§
Costs of treatment (€), by age at beginning of treatment	E	c_tx			
3 years			1611	898–2324	Student's t (6; 1611; 291)¶
4 years			1536	818–2255	Student's t (6; 1536; 294)¶
5 years			1454	735–2174	Student's t (6; 1454; 294)¶
6 years			1363	641–2085	Student's t (6; 1363; 295)¶
7 years			1260	525–1995	Student's t (6; 1260; 300)¶
8 years			1089	448–1729	Student's t (6; 1089; 262)¶
9 years			913	344–1482	Student's t (6; 913; 232)¶
10 years			665	232–1098	Student's t (6; 665; 177)¶
Transition Probabilities					
Mortality for men	L	tp_Deathm	#	(None)	(None)
Mortality for women	L	tp_Deathf	**	(None)	(None)
Probability of visit to ophthalmologist per year, by age	L	op_visit			p_visit × op_visit, distribution of p_visit: Uniform [0.5; 1.5]†
3 years			0.230	0.115–0.345	
4 years			0.270	0.135–0.406	
5 years			0.349	0.175–0.524	
6 years			0.332	0.166–0.498	
7 years			0.344	0.172–0.516	
8 years			0.325	0.162–0.487	
9 years			0.278	0.139–0.417	
10 years			0.305	0.152–0.457	
Probability of treatment success when started late:	L	eff_late			
eff_late _k = [p_eff/(1 - p_eff) × OR _k]/[1 + p_eff/(1 - p_eff) × OR _k] with OR _k for treatment success, by age class (k)					
3 years			OR = 1.000	(None)	(None)
4–5 years			OR = 0.468	0.298–0.734	OR = exp(b); distribution of b: Normal (-0.76; 0.23)§
6–10 years			OR = 0.317	0.187–0.536	OR = exp(b); distribution of b: Normal (-1.15; 0.27)§

Model Parameter	Source*	Name Used in Decision Tree or Markov Model	Parameter Value (Base Value)	Range Used for Univariate Sensitivity Analysis	Distribution Used in Monte Carlo Simulation
Transition probability per year from unilateral visual impairment (amblyopia) to bilateral visual impairment, by gender and age					
Men	L	tp_vis_m	0.00166 0.00005 0.00204 0.00248 0.00325	0.00083–0.00248 0.00003–0.00008 0.00102–0.00306 0.00124–0.00372 0.00163–0.00488	p_trans × tp_vis_m, distribution of p_trans: Uniform [0.5; 1.5]†
Women	L	tp_vis_f	0.00046 0.00026 0.00088 0.00324 0.00201	0.00023–0.00069 0.00013–0.00039 0.00044–0.00132 0.00162–0.00486 0.00100–0.00301	p_trans × tp_vis_f, distribution of p_trans: Uniform [0.5; 1.5]†
Transition probability per year from healthy to unilateral visual impairment (no amblyopia), by gender and age					
Men	L	tp_mono_m	0.00435 0.00012 0.00541 0.00629 0.00729	0.00217–0.00652 0.00006–0.00018 0.00270–0.00811 0.00315–0.00944 0.00365–0.01094	p_trans × tp_vis_f, distribution of p_trans: Uniform [0.5; 1.5]†
Women	L	tp_mono_f	0.00121 0.00068 0.00232 0.00862 0.00431	0.00061–0.00182 0.00034–0.00103 0.00116–0.00347 0.00431–0.01293 0.00215–0.00646	p_trans × tp_mono_f, distribution of p_trans: Uniform [0.5; 1.5]†
Discount Rate	L	cDR	0.050	0.000–0.050	(None)
Discount rate for costs	L	oDR	0.050	0.000–0.050	(None)

* Source of parameter value: L, literature; E, field study; E, expert opinion (see text for details).

† Beta distribution (α ; β); values of α and β with decimal digits stand for fitted Beta distributions.

‡ Uniform distribution (minimum; maximum).

§ Normal distribution (mean; standard deviation).

|| Discounted by 5% discount rate to year when treatment is started.

¶ Student's *t* distribution (degrees of freedom; mean; observed standard deviation).

Mortality rate for men in Germany.

** Mortality rate for women in Germany.

Model Parameters

Model parameter values were obtained from a field study of orthoptic screening in kindergarten, from the literature, and from expert interviews (Table 1). For each parameter, a (mean) value, which was used for the base analysis, was defined as well as a possible value range, according to which parameter values were varied in 1-way sensitivity analysis. Furthermore, for parameters that, in principle, could be sampled, distributions were specified for probabilistic sensitivity analysis (Monte Carlo simulation) as follows: for proportions of the form $a/(a + b)$, such as test characteristics and utilities, which are bound to a 0–1 interval, Beta distributions were specified [Beta (α, β) with $\alpha = a + 1$ and $\beta = b + 1$].^{26,27} Normal or Student's t distributions were specified for the mean of continuous parameter values. If there were only data on the mean and the likely range of the 95% confidence interval (CI) of a continuous parameter but not the variance, the standard error was estimated by the formula

$$s \approx \frac{u - l}{2 \times 1.96} \quad (2)$$

where u and l are the upper and lower limits of the range, respectively.²⁷ If there was only information on the range but not on the mean of a parameter, then a uniform distribution was used.

Screening Population

In 2000, the number of children aged 3 in Germany was 814 700; 51.3% were male. Of these children, 458 700 (56.3%) were attending 1 of 43 728 kindergartens.²⁸ Thus, the mean number of children aged 3 per kindergarten was 10.49. According to other field studies of vision screening in German kindergartens in which the participation rate ranged between 88%²⁹ and 92%,³⁰ the participation rate was assumed to be 90%, which would yield a mean of 9.44 participating children per kindergarten. For univariate sensitivity analysis, the range of the participation rate was set at 80% to 100%. This range was assumed to represent the 95% CI, and the standard error was estimated according to formula 2. Based on the estimated mean and standard error, a β distribution was fitted by using the method of moments estimation.^{27,31}

Prevalence of Target Disease and Test Characteristics of Orthoptic Screening

Data on the prevalence of untreated amblyopia, on the screening test characteristics (sensitivity, specificity, and proportion of inconclusive results), and on the compliance with ophthalmologic examinations were obtained from a field study in which 1180 3-year-old children were examined by orthoptists in 121 German kindergartens.¹⁰

In the field study, the orthoptic screening examination consisted of cover tests, examination of eye motility and head posture, and uncorrected monocular VA testing using the Lea single optotype test (required test distance 10 ft [3 m], Precision Vision, Villa Park, IL) with pass threshold set at ≥ 0.8 (20/25) monocular VA in both eyes; or ≥ 0.5 (20/40) in both eyes and ≤ 1 line difference between the VA of the right and left eyes (L. Hyvärinen, written communication, 1998). The ophthalmologic criteria for a target disease (positive gold standard) were: any newly administered patching therapy; or any newly administered spectacle therapy, if the corrected VA was ≤ 0.4 (20/50) in either eye or if the difference of VA between the right and left eyes was ≥ 3 lines.¹⁰

For the base analysis it was assumed that, without treatment, children with a positive gold standard (2.2%) would be affected by a lasting unilateral visual impairment. For univariate sensitivity analysis, the range of prevalence was set at 1.0% to 5.0%, which includes the range of prevalence of amblyogenic factors stated in a recent review (2.7%–4.4%)⁵ and takes into account that more than half of all cases may already be detected at the time of screening. For the Monte Carlo simulation, a Beta distribution was defined based on the proportion of gold-standard-positive children found in the field study, $a/(a + b)$, where a equals the number of children with positive gold standard, and b equals the number of children without positive gold standard.

For the test characteristics and compliance rates, values measured in the field study were used as base values. The limits of their 95% CIs were used for univariate sensitivity analysis, and Beta distributions were specified for the Monte Carlo simulation.

Because data on the compliance with direct referral to an ophthalmologist after inconclusive screening could not be obtained from the field study due to its design,¹⁰ this value had to be estimated based on other studies^{14,32} and was set at 90%; in univariate sensitivity analysis, this value was varied from 80% to 100%, and for Monte Carlo simulation a uniform distribution on this interval was used.

Success Rate of Treatment

In studies that used a VA ≥ 0.5 as the threshold for treatment success for amblyopia, reported success rates range from 50% to 100%.^{14,33–36} In a meta-analysis based on 23 studies with 689 patients included, Flynn et al¹⁶ found an overall success rate of 74.3%. The authors developed a multiple logistic regression model to analyze the effect of age at the beginning of treatment on the success rate, controlling for type of amblyopia and VA at beginning of treatment. In a second study,¹⁷ in which 589 patients were included and the overall success rate was 59.9%, Flynn et al validated the regression model and again found a significant influence of age on treatment success: Compared with age group 0 to 3, the odds ratio (OR) for the success of treatment started at age group 4 to 5 was 0.47, and at age group 6 to 10 it was 0.32. They also reported the estimated regression coefficients b_k^* for the age groups k and their standard errors s_k , from which the 95% CIs of OR _{k} could be calculated by using the formula³⁷

$$\exp(b_k^* \pm 1.96 \times s_k), \quad (3)$$

which yielded [0.30; 0.73] for age group 4 to 5 and [0.19; 0.54] for age group 6 to 10.

In the model, the mean treatment success rate at age 3 (p_eff) was set at 75% with the range of 60% to 90% used for univariate sensitivity analysis. For the Monte Carlo simulation, a Beta distribution was fitted for (p_eff), assuming that the mentioned range corresponds to the 95% CI and deriving the standard error according to formula 2.

Success rates of treatment started after age 3 (eff_late) were calculated based on the mentioned OR _{k} for older age groups k using the formula

$$\text{eff_late}_k = \frac{p_eff / (1 - p_eff) \times \text{OR}_k}{1 + p_eff / (1 - p_eff) \times \text{OR}_k} \quad (4)$$

Table 2 shows the resulting success rates of late treatment (eff_late) for various success rates of treatment started at age 3. In univariate sensitivity analysis, OR _{k} values were jointly varied according to their 95% CI, mentioned above. In the Monte Carlo simulation, OR _{k} values were calculated by using the coefficients for age groups b_k , which were estimated in the logistic regression model, in the formula³⁷

$$\text{OR}_k = \exp(b_k), \quad (5)$$

assuming a normal distribution for b_k , with the estimated b_k^* as the mean and the standard deviation s_k . Thus, b_k was calculated according to the formula³⁷

$$b_k = b_k^* + z \times s_k, \quad (6)$$

where z follows the standard normal distribution.

TABLE 2. Success Rate of Amblyopia Treatment Started After Age 3 by Success Rate of Amblyopia Treatment Started at Age 3

Success Rate Age 3, % (p_eff)	Success Rate Ages 4–5, % (OR = 0.47*)	Success Rate Ages 6–10, % (OR = 0.32*)
90	81	74
85	73	64
80	65	56
75	58	49
70	52	42
65	46	37
60	41	32

* OR for success rate of treatment started after age 3 (eff_late) compared to success rate of treatment started at age 3; the success rate (eff_late) is calculated according to the formula: $\text{eff_late} = [p_eff / (1 - p_eff) \times \text{OR}] / [1 + p_eff / (1 - p_eff) \times \text{OR}]$.

Utilities

The utility of the bilateral-visual-impairment health state was derived from a recent study of a large series of patients with visual loss from various ocular diseases using the time trade-off method.³⁸ In this study, a formula for converting VA of the better eye to a mean utility value was derived via regression analysis:

$$U(\text{VA}) = 0.374 \times \text{VA} + 0.514. \quad (7)$$

For the incremental approach, only the reduction of utility caused by visual impairment was relevant. Therefore, the utility of VA = 1.0 was set at $U = 1.00$, ie, it was increased by 0.11 compared with the utility predicted by formula 7. For a conservative estimate of the utility of the bilateral-visual-impairment health state, the utility of VA = 0.4 was calculated by using formula 7 and then also increased by 0.11, resulting in $U = 0.78$. For the health states without any visual impairment, the utility was constantly set at $U = 1.00$. Thus, compared with no visual impairment, utility of bilateral visual impairment was reduced by 0.22. Based on utilities reported by the same working group,³⁹ this reduction of utility was varied by $\pm 1/3$, ie, from 0.15 to 0.29 for univariate sensitivity analysis, corresponding to utilities from 0.71 to 0.85. This interval was considered to reflect the 95% CI, and a Beta distribution was fitted for the Monte Carlo simulation.

The utility associated with unilateral visual impairment caused by amblyopia has not been investigated specifically thus far. In another recent study of the same working group, unilateral impairment (defined as VA ≤ 0.5 in the worse eye and VA ≥ 0.8 in the better eye) from various ocular diseases was found to cause a mean reduction of utility of 0.08, using the time trade-off method.⁴⁰ Because individuals with only 1 sound eye due to amblyopia since childhood may develop compensatory visual mechanisms,⁵ the reported utility reduction was considered a maximum. It thus was varied from 0.00 to 0.08 in univariate sensitivity analysis, ie, from no disutility at all to the disutility stated in the empirical study,⁴⁰ which corresponds to utilities ranging from 0.92 to 1.00. Having no additional information, for the base analysis a utility of 0.96 was used, ie, the middle of the interval. For the Monte Carlo simulation, a uniform distribution on the interval [0.92;1.00] was used.

Transition Probabilities of Markov Model

Probability of Successful Treatment Under Usual Care

Data on the age-specific probability for an amblyopic child to receive treatment under usual care (ie, without orthoptic screening) were not available. It was assumed that untreated amblyopia was definitely detected and treated when an affected child was seen by an ophthalmologist for whatever reason. The probability for a child to be seen by an ophthalmologist (op_visit) was estimated based on a representative survey of ophthalmologic patients in Germany,⁴¹ from which the age-specific number of ophthalmologic patients in 1997 was estimated.⁴² The age-specific probability (op_visit) was derived by dividing the age-specific number of patients by the age-specific number of persons in the German population in 1997.⁴³ For the model it was assumed that the age-specific probability (op_visit) was independent of former visits and of the presence of amblyopia. Table 1 shows the calculated probabilities. For example, without screening, at age 3, 23% of children with untreated amblyopia would be seen by an ophthalmologist and receive treatment; before age 8, a cumulated 84% would have received treatment, and 95% would have received treatment before age 11. In univariate sensitivity analysis, the age-specific probabilities were reduced and increased by the factors 0.5 and 1.5, respectively. For the Monte Carlo simulation, a uniform distribution on the interval [0.5; 1.5] was used for this factor.

In Markov process A, before age 11 (cycles 0–7) the probability of transition from the untreated-amblyopia health state to the healthy state (ie, the probability of successful treatment under usual care) was obtained by multiplying the age-specific probability (op_visit) with the age-group-specific success rate of treatment started later (eff_late). From age 11 onward, the probability of this transition was assumed to be 0.

Incidence of Unilateral Visual Impairment Other Than Amblyopia

Because data on age- and gender-specific incidence of unilateral visual impairment were not available in the literature, incidence

had to be derived from prevalence studies. Because such studies were not available for Germany, data from the Australian Blue Mountain Eye Study (BMES)^{23,44} were used, assuming Australian population and health care characteristics comparable to those in Germany. The BMES provides data on the prevalence of unilateral visual impairment by age group, gender, and severity as well as on the prevalence of amblyopia for a representative population sample aged ≥ 49 years. To estimate the prevalence of unilateral visual impairment not caused by amblyopia, age- and gender-specific prevalence data of unilateral visual impairment²³ were reduced by prevalence data of amblyopia.⁴⁴

It was assumed that the reported age-group-specific prevalence is present in the middle of the age classes. Thus, new age classes were defined, the middle of the original age classes being the limits of the new age classes. Gender-specific transition probabilities $q_{t,j}$ for the new age classes of the width t years and upper limit j years were derived by using the formula⁴⁵

$$q_{t,j} = \frac{P_j - P_{j-t}}{1 - P_{j-t}}, \quad (8)$$

where P_j and P_{j-t} is the prevalence at the upper and lower limit of the new age class, respectively. One-year probabilities of the transition from the healthy state to the unilateral-visual-impairment (no amblyopia) health state q_j were derived by using the formula⁴⁵

$$q_j = 1 - (1 - q_{t,j})^{1/t}. \quad (9)$$

In the Markov model, this transition probability was labeled tp_mono.m for males and tp_mono.f for females.

Incidence of Bilateral Visual Impairment Caused by Unilateral Eye Disease Affecting the Nonamblyopic Eye of Amblyopic Persons

It was assumed that the prevalence of an eye being affected by unilateral, visually-impairing eye disease other than amblyopia is the same for amblyopic and nonamblyopic eyes. Thus, in persons with unilateral amblyopia, the prevalence of the nonamblyopic eye being affected, causing bilateral visual impairment, was estimated to be half of the prevalence of unilateral visual impairment in persons without amblyopia. Because VA in the amblyopic eye may improve when the other eye becomes visually impaired, the estimated prevalence was reduced by 23.6%, based on the results of a study of 144 amblyopic persons with visual impairment in the nonamblyopic eye, of whom 34 (23.6%) improved to VA ≥ 0.5 in the amblyopic eye.⁴⁶ Based on these estimated prevalence data, age-group- and gender-specific incidence, ie, 1-year probability of transition from the unilateral-visual-impairment (amblyopia) health state to the bilateral-visual-impairment (with amblyopia) health state was derived as described above and labeled tp_vis.m for males and tp_vis.f for females.

In univariate sensitivity analysis, the transition probabilities tp_mono.m, tp_mono.f, tp_vis.m and tp_vis.f were jointly reduced and increased by the factors 0.5 and 1.5, respectively. For the Monte Carlo simulation, a uniform distribution on the interval [0.5; 1.5] was used for this factor.

Mortality

Age- and gender-specific death rates were obtained from the most recent life table 1997/1999 of the German Federal Statistical Office.⁴⁷

Costs

Costs were calculated in Euro (€) from a third-party payer perspective, assuming no copayments by the patients, for the year 2000. In that year, the average Euro to US-dollar exchange rate was \$0.92 per 1 €, and the average purchasing power adjusted conversion rate was \$0.99 per 1 € (ie, close to parity).⁴⁸

Costs of Orthoptic Screening Examination

Data on costs of orthoptic screening examination were obtained from the field study mentioned above; the methodology of cost measurement performed in the field study has been described elsewhere.¹¹ Costs of orthoptic screening were divided into costs for organizing the screening program (which were assumed to be fixed per kindergarten) and costs for conducting the screening examination in kindergarten, the latter being subdivided into fixed costs (eg, for traveling) and variable costs, which depended on the number of children examined per kindergarten (eg, labor

TABLE 3. Costs of Treatment of Amblyopia by Age at Beginning of Treatment

Age j at Beginning of Treatment, y	Costs Without Discounting		Costs Discounted at 5% Discount Rate*	
	Mean m_j , €	Standard Error s_j , €	Mean m_j , €	Standard Error s_j , €
3	1844	316	1611	291
4	1734	318	1536	294
5	1619	319	1454	294
6	1496	320	1363	295
7	1365	327	1260	300
8	1156	278	1089	262
9	953	243	913	232
10	680	180	665	177

* Discounted to year at beginning of treatment.

costs for examination time). Vice versa, when calculating the costs per single examination, the share of fixed costs depended on the number of children per kindergarten. Because in the field study the average number of children examined per kindergarten was 9.88, fixed costs per screening examination were multiplied by the factor 9.88/9.44 to adjust for the smaller average number of children (9.44) to be examined per kindergarten in the model.

For the fixed and variable costs per single examination in kindergarten, standard errors of the mean could be estimated in the field study. Thus, these 2 cost categories were varied according to their 95% CI in univariate sensitivity analysis, and normal distribution was used for Monte Carlo simulation. With respect to organization, costs were varied by $\pm 50\%$ in univariate sensitivity analysis. This interval was considered to reflect the 95% CI, and for Monte Carlo simulation, a normal distribution was fitted by estimating the standard error according to formula 2. To account for changes in participation rates that would lead to changes in the average number of children examined per kindergarten, the mean and the standard error of fixed costs were adjusted accordingly.

Costs of Diagnostic Ophthalmologic Examination

Costs of a standard ophthalmologic examination of children referred from screening were based on the German social health insurance's relative value scale for outpatient physician services, which defines individual physician services and states point volumes for them. For the conversion factor (point value), which varies by region, a German average of 0.041 € was used. For univariate sensitivity analysis, these costs were varied by $\pm 25\%$, which corresponded to the range that the conversion factor was likely to vary by region. This interval was considered to reflect the 95% CI. For Monte Carlo simulation, a normal distribution was fitted by estimating the standard error according to formula 2.

Treatment Costs

Because there were no published data on the cost of amblyopia treatment, costs were estimated based on expert opinion. Seven experts of amblyopia treatment (4 ophthalmologists and 3 orthoptists) from different German treatment centers filled in a standardized questionnaire. In the questionnaire, medical services and items (eg, glasses, patches, etc) possibly used for amblyopia treatment were listed. Respondents were asked to estimate the mean number of services and items used per year during up to 9 years of treatment. Services and items were valued monetarily by using "administrative" prices paid by the German statutory health insurance. The methodology of estimating treatment costs has been described in detail elsewhere.⁴⁹

It was assumed that treatment was completed before age 12. To estimate the costs of treatment started later than age 3, treatment costs were reduced by excluding the costs of years that would correspond to age ≥ 12 .

Costs of treatment started at age 3 to 10 were calculated based on each expert's estimate of resource utilization. From these figures, mean estimated costs m_j^* of treatment started at age j were calculated (and used for the base analysis) as well as the standard errors s_j , which reflect the uncertainty of the expert with regard to average costs (Table 3). In univariate sensitivity analysis, the treatment costs were jointly varied according to their 95% CIs. For the Monte Carlo simulation, age-specific treatment costs k_j were calculated by using the formula⁵⁰

$$k_j = m_j^* + t \times s_j \quad (10)$$

where t follows a standardized Student's t distribution with 6 degrees of freedom.

Discounting

Costs and effects were discounted at 5%. In sensitivity analysis, discount rates of 0% and 3% were also used.

Sensitivity Analysis

One-Way Sensitivity Analysis

To analyze the effect of uncertainty in single model parameters on the ICER, univariate sensitivity analysis was performed by varying single-parameter values according to the ranges described above and repeating the analytical solution of the model.

Probabilistic Sensitivity Analysis (Monte Carlo Simulation)

In the Monte Carlo simulation, values for all model parameters were sampled from their respective specified distributions, and the analytical solution of the model was repeated.²⁷ This process of resampling from each of the distributions and recalculating the incremental costs and effects from the model was repeated 10 000 times to generate a distribution of the estimated ICER. Uncertainty intervals were estimated from the simulated data by taking the 2.5th and 97.5th percentile values to represent the endpoints for a 95% interval,²⁷ and cost-effectiveness acceptability curves^{51–53} were constructed.

RESULTS

Base Result

The screening strategy was associated with incremental costs of 13.34 € per child and incremental effects of 1.803×10^{-3} QALYs per child. Thus, for the total population of 814 700 3-year-old children, of whom 50.7% (90% \times 56.3%) would participate in screening, incremental costs were 10.87 million €, and incremental effects were 1469 QALYs. The ICER was 7397 €/QALY.

Univariate Sensitivity Analysis

Figure 3 shows the effect of varying single-parameter values on the ICER. Uncertainty with respect to the utility of unilateral visual impairment had the strongest potential impact on the ICER. If this utility was 0.92, the ICER decreased to 3706 €/QALY; if it was 1.00 (ie, if only bilateral visual impairment but not unilateral visual impairment affected utility), the ICER increased to 1.904 million €/QALY. Figure 4 shows the association between the ICER and the utility of unilateral visual impairment for the range of 0.92 to 1.00: If the utility was >0.988 , the ICER increased to $>25\,000$ €/QALY; if it was >0.994 , the ICER increased to $>50\,000$ €/QALY.

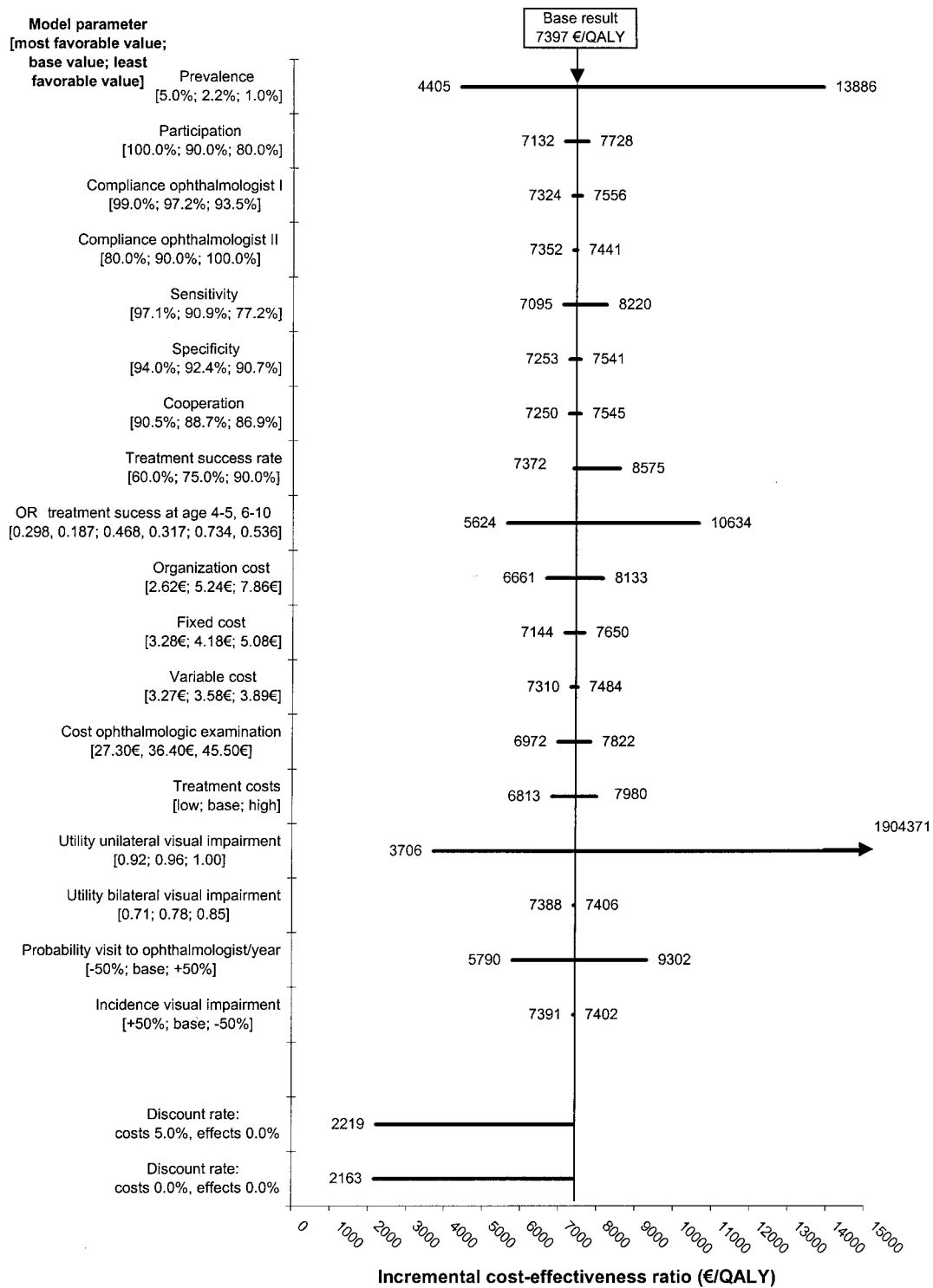


Fig 3. Results of univariate sensitivity analysis.

Besides uncertainty with respect to the prevalence of the target disease, the OR of treatment success when started later and the probability for a child to be seen by an ophthalmologist (op-visit) had a marked, but much smaller impact on the ICER. All other parameters had only little impact, among those being the treatment costs. Almost no impact had the uncertainty with respect to the incidence of visual impairment caused by other eye disease and the utility of bilateral visual impairment.

If only costs but not effects (QALYs) were discounted, the ICER was 2219 €/QALY. If neither costs nor effects were discounted, the ICER was 2163 €/QALY. If costs and effects were both discounted at 3%, the ICER was 5138 €/QALY.

Monte Carlo Simulation

Uncertainty Intervals

Figure 5 shows the joint distribution of incremental costs and effects generated in 10 000 simulations

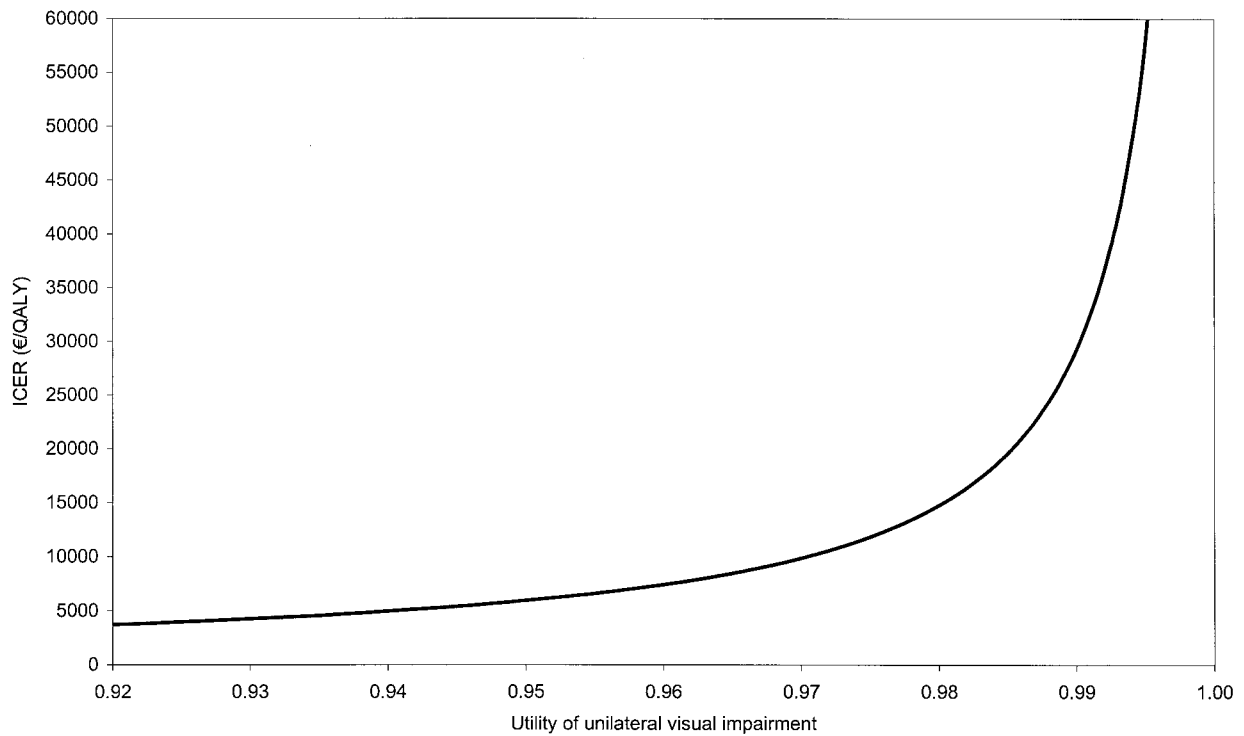


Fig 4. Effect of utility of unilateral visual impairment on ICER of the screening strategy (when base values are used for all other model parameters).

on the cost-effectiveness plane. The 95% uncertainty interval for the incremental costs per child ranged from 10.14 to 18.63 €, and for incremental effects from 8.607×10^{-5} to 5.005×10^{-3} QALYs. For the total population of 814 700 3-year-old children, 95%

of incremental costs lay between 8.26 million and 15.18 million €, and 95% of the effects lay between 70 and 4078 QALYs.

The 95% uncertainty interval of the ICER was 3138 to 148 228 €/QALY, which corresponds to the slope

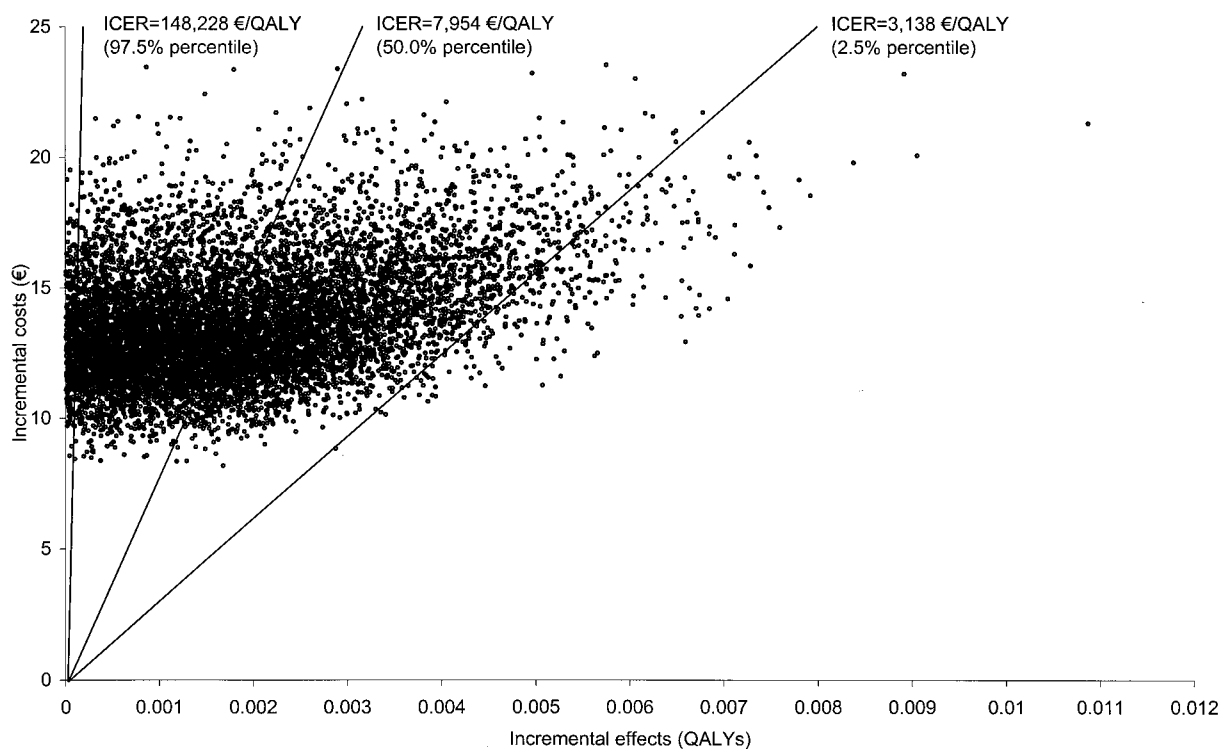


Fig 5. Joint distribution of incremental costs and effects of the screening strategy plotted on the cost-effectiveness plane. Results of 10 000 Monte Carlo simulations; lines represent 2.5th, 50th, and 97.5th percentiles of ICER.

of the lines representing the interval endpoints in Fig 5. The 90% uncertainty interval was 3452 to 72 637 €/QALY. Without discounting effects, the 95% uncertainty interval was 903 to 30 053 €/QALY.

Cost-Effectiveness Acceptability Curves

When costs and effects were discounted at 5%, Monte Carlo simulation yielded an ICER <25 000 €/QALY in 84% and an ICER <50 000 €/QALY in 92%. Without discounting effects, an ICER <25 000 was yielded in 97%, and an ICER <50 000 in 99%. This is displayed by the cost-effectiveness acceptability curves in Fig 6, which give the proportion of observed simulation results lying below varying threshold values for the ICER.

Because much of the variability of the results was due to the uncertainty with respect to the utility of unilateral visual impairment, Monte Carlo simulations were also performed keeping this utility constant at different levels and varying all other parameters according to their distributions (Fig 7). If this utility was 0.92, 0.94, 0.96, 0.98, or 0.99, in 95% of the simulations the ICER was below 6375, 8491, 12 561, 25 296, and 50 789 €, respectively.

DISCUSSION

The Markov model aimed at estimating the life-long gain in health-related quality of life due to prevention of amblyopia through screening and subsequent treatment, as well as the associated costs. The model took into account that children with untreated amblyopia may be detected and successfully treated without screening.

The ICER of orthoptic screening at age 3 in German kindergarten was found to be ~7400 €/QALY

when costs and effects were discounted at 5%. Thus, the ICER was more favorable than that of many routinely provided health care interventions⁵⁴ and similar to the ICER found for diabetic retinopathy screening.⁵⁵

Because the analysis included effects occurring during the remaining lifetime of up to 86 years, discounting effects had substantial impact on the ICER. Without discounting, the ICER was just about one third of the base result. Discounting costs had only little impact on the ICER, because all considered costs occurred during the first 9 years after screening.

Univariate sensitivity analysis showed that uncertainty regarding the utility of unilateral visual impairment had great potential impact on the ICER. If unilateral visual impairment due to amblyopia was not associated with any loss in utility, then orthoptic screening would very likely not be cost-effective. In other words: Just the risk of developing bilateral visual impairment later in life would most likely not justify preschool vision screening. Yet, if unilateral visual impairment was associated with a utility loss of only 0.01, then the ICER fell below 30 000 €/QALY. ICERs obtained in Monte Carlo simulation showed a wide range but were mostly within limits likely to be acceptable to decision-makers. For example, the probability for the ICER being below 25 000 €/QALY was 84%.

Model Structure

For modeling, the course of disease had to be divided into distinct states, and hence thresholds for the presence or absence of visual impairment had to be defined. Because in reality no such thresholds

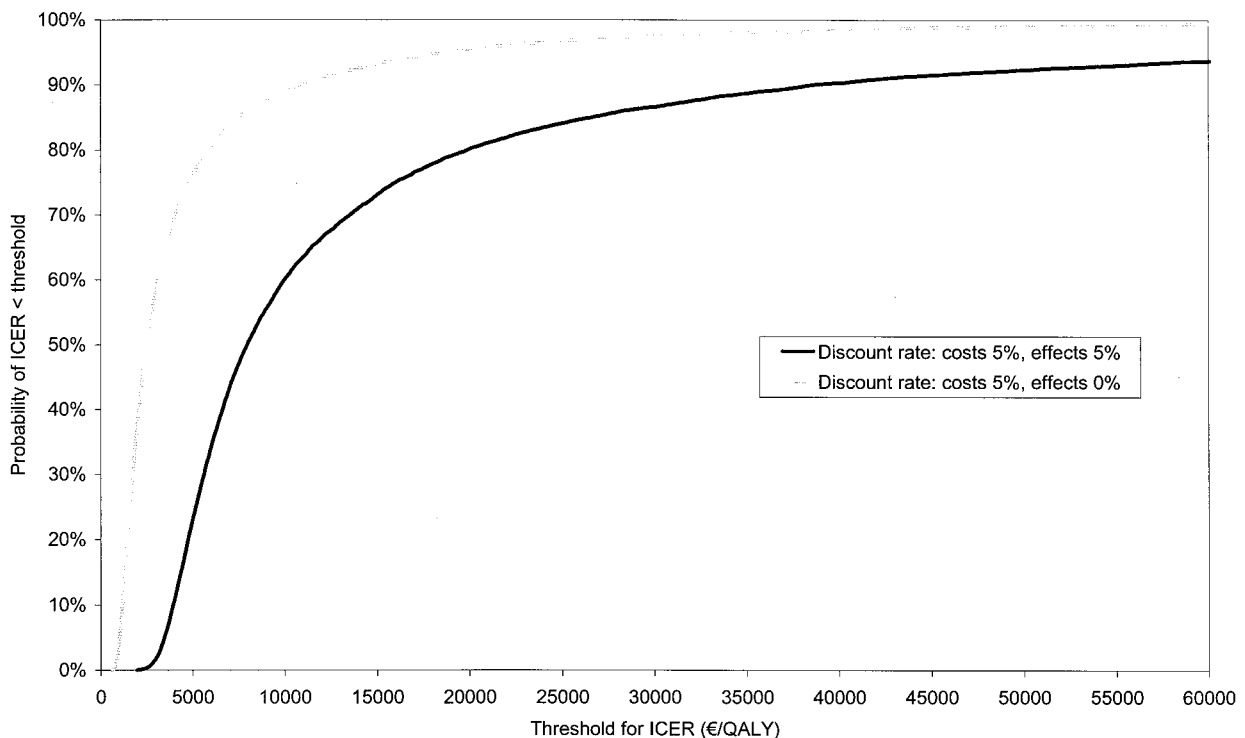


Fig 6. Cost-effectiveness acceptability curves of the screening strategy for various discount rates

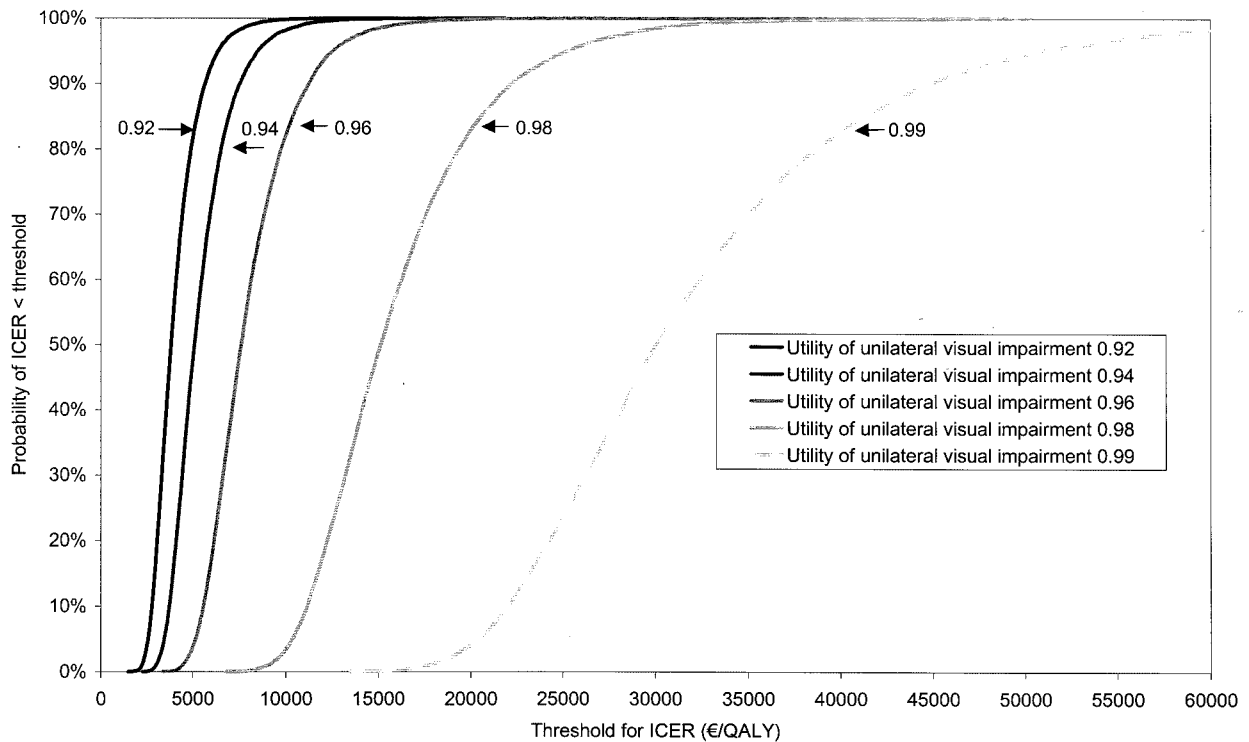


Fig 7. Cost-effectiveness acceptability curves of the screening strategy for various utilities of unilateral visual impairment held constant.

exist, this necessarily entails some simplification. Theoretically, more health states could have been defined, representing different degrees of visual impairment but also making the model more complicated. However, for this purpose, respective data were not available. Uncertainty with respect to the definition of thresholds was addressed by varying model parameters widely in sensitivity analyses.

The model did not distinguish between different causes of amblyopia. This simplification had to be made, because data on age-specific treatment success rates were not available in more detail. However, in 2 studies based on large patient samples,^{16,17} no significant effect of the cause of amblyopia on treatment success was identified after controlling for age and VA at the beginning of treatment, which both had a significant effect. Although the effect of age was explicitly modeled, the effect of VA on treatment success was accounted for by varying treatment success rate at age 3 widely in sensitivity analysis.

The model did not allow any return to preceding health states, eg, from unilateral visual impairment (without amblyopia) to healthy (eg, after treatment of other eye diseases) states. Although such a return may be possible in reality (eg, after treatment), the error introduced by this restriction is considered small because most transition probabilities were estimated from prevalence data. For improvement of unilateral visual impairment caused by amblyopia without treatment, no evidence was found in the literature.⁵⁶

The Markov process was restricted to 86 cycles. Thus, possible QALY gains accumulated from age 90 onward were not considered. When discounted,

these QALY gains would be very small. Without discounting, however, they would make the ICER more favorable, as would a future increase in life expectancy.

In the model, only direct medical costs for detection and treatment of the target disease were considered from the perspective of a third-party payer. Thus, administrative prices (eg, fees) were used where applicable. Because these prices are likely to be comparable to opportunity costs, and no cost sharing by patients was assumed, the calculated costs are probably close to direct medical costs from the societal perspective. Direct nonmedical costs (eg, for traveling to an ophthalmologist) were not considered but probably were small. Direct costs caused by visual impairment coming along later in life were not considered either, but it would have had only little impact on the results due to discounting: If, eg, bilateral visual impairment caused direct costs of 10 000 € per year, the ICER would drop by only 2% to 7220 €/QALY. Indirect costs, caused by losses in productivity due to visual impairment, were not considered either. Including indirect costs would most likely make the ICER more favorable. However, according to the Panel on Cost-Effectiveness in Health and Medicine,⁵⁷ effects of disease on productivity or income may already be reflected by the preference-based valuation of health-related quality of life (utility), and hence there would be no need to include indirect costs in cost-utility analysis.

Critical Model Parameter Values

Values for model parameters were obtained from a field study, the literature, and a survey of experts.

Uncertainty in parameter values was addressed by defining a base model and conducting extensive univariate sensitivity analysis. To assess uncertainty in various model parameters simultaneously, probabilistic sensitivity analysis (Monte Carlo simulation) was performed.

Regarding the prevalence of the target disease, it was assumed that all children detected in the field study would have been affected by lasting visual impairment without treatment. By this assumption, prevalence may be overestimated. However, in the field study, the combined prevalence of the target disease and of amblyopia already treated was ~4%,¹⁰ which falls within the prevalence range for unscreened populations given in the literature.^{1,44,58}

The age-specific probability that a child with untreated amblyopia is seen by an ophthalmologist under usual care was derived from a representative cross-sectional survey of ophthalmologic patients in Germany. This procedure entailed considerable uncertainty, which was addressed by varying these parameter values widely in sensitivity analysis and assuming uniform distribution in Monte Carlo simulation. Under the extreme assumption that, without screening, none of the target diseases would be detected, the ICER would drop to 4528 €/QALY.

The incidence of unilateral visual impairment not caused by amblyopia was estimated based on a large epidemiologic study from Australia (BMES). Because of comparable ethnicity and level of medical care, comparability with the German population was assumed. Yet, in the BMES, only adults aged ≥49 were included. Thus, visual impairment before age 49 was not considered. However, its prevalence in the youngest age group (49–54 years) of the BMES was rather low. If the prevalence rate in the age group 49–54 years resulted from a constant incidence from birth onward, the ICER would be reduced only slightly to 7392 €/QALY.

For valuation of health-related quality of life, utilities were used that had been elicited for visual impairment caused by various ocular diseases. In the underlying studies, neither the cause of visual impairment nor age, gender, education, ethnicity, comorbidity, or duration had a significant impact on the utility. Thus, the use of nondisease-specific utilities seems to be justified, because specific data on the utility associated with unilateral visual impairment caused by amblyopia were not available in the literature. However, the utility loss caused by amblyopia starting very early in life could be comparatively smaller due to adaptation. Hence, in sensitivity analysis, the respective utility loss was varied widely, even to no loss at all.

The best estimate available for the costs of amblyopia treatment was based on expert opinion, which showed only little variability. For treatment started after age 3, lower treatment costs were assumed because of shorter duration. However, it may be possible that late treatment would be more intense and thus more costly, which would make the ICER more favorable.

CONCLUSIONS

The ICER of orthoptic screening seems to fall within a range that warrants careful consideration by decision-makers. The presented uncertainty intervals and cost-effectiveness acceptability curves should enable decision-makers to appraise the results based on their own risk aversion. Much of the uncertainty in results comes from the uncertainty regarding the effect of amblyopia on health-related quality of life. To reduce this uncertainty, additional studies should investigate the impact of amblyopia on utility in more detail.

ACKNOWLEDGMENTS

This study was supported by the Fortune program of the Medical Faculty of the University of Tübingen (grant 846-0-0) and by the Federal Ministry of Education and Research (grant NBL-14-001).

We thank Bernd Schweikert, MSc, and Robert Welte, MSc MPH, Department of Health Economics, University of Ulm, for comments on an earlier version of the article.

REFERENCES

1. Lennerstrand G, Jakobsson P, Kvarnstrom G. Screening for ocular dysfunction in children: approaching a common program. *Acta Ophthalmol Scand.* 1995;73(suppl 214):26–38
2. Simons K. Preschool vision screening: rationale, methodology and outcome. *Surv Ophthalmol.* 1996;41:3–30
3. Sjostrand J, Abrahamsson M. Prevention of amblyopia and the concept of cure. *Eur J Ophthalmol.* 1997;7:121–129
4. Ingram RM, Holland WW, Walker C, Wilson JM, Arnold PE, Dally S. Screening for visual defects in preschool children. *Br J Ophthalmol.* 1986;70:16–21
5. Snowdon SK, Stewart-Brown SL. Preschool vision screening. *Health Technol Assess.* 1997;1:1–83
6. Wright MC, Colville DJ, Oberklaid F. Is community screening for amblyopia possible, or appropriate? *Arch Dis Child.* 1995;73:192–195
7. Hartmann EE, Dobson V, Hainline L, et al. Preschool vision screening: summary of a Task Force report. *Pediatrics.* 2000;106:1105–1116
8. Kemper AR, Margolis PA, Downs SM, Bordley WC. A systematic review of vision screening tests for the detection of amblyopia. *Pediatrics.* 1999;104:1220–1222
9. Russell LB, Gold MR, Siegel JE, Daniels N, Weinstein MC. The role of cost-effectiveness analysis in health and medicine. Panel on Cost-Effectiveness in Health and Medicine. *JAMA.* 1996;276:1172–1172
10. Barry JC, König HH. Test characteristics of orthoptic screening examination in three-year-old kindergarten children. *Br J Ophthalmol.* 2003;87:909–916
11. König HH, Barry JC, Leidl R, Zrenner E. Economic evaluation of orthoptic screening: results of a field study in 121 German kindergartens. *Invest Ophthalmol Vis Sci.* 2002;43:3209–3215
12. König HH, Barry JC. Economic evaluation of different methods of screening for amblyopia in kindergarten. *Pediatrics.* 2002;109(4). Available at: <http://pediatrics.aappublications.org/cgi/content/full/109/4/e59>
13. American Academy of Pediatrics, Committee on Practice and Ambulatory Medicine. Vision screening and eye examination in children. *Pediatrics.* 1986;77:918–919
14. Newman DK, Hitchcock A, McCarthy H, Keast-Butler J, Moore AT. Preschool vision screening: outcome of children referred to the hospital eye service. *Br J Ophthalmol.* 1996;80:1077–1082
15. Egan DF, Brown R. Vision testing of young children in the age range 18 months to 4 1/2 years. *Child Care Health Dev.* 1984;10:381–390
16. Flynn JT, Schiffman J, Feuer W, Corona A. The therapy of amblyopia: an analysis of the results of amblyopia therapy utilizing the pooled data of published studies. *Trans Am Ophthalmol Soc.* 1998;96:431–450
17. Flynn JT, Woodruff G, Thompson JR, et al. The therapy of amblyopia: an analysis comparing the results of amblyopia therapy utilizing two pooled data sets. *Trans Am Ophthalmol Soc.* 1999;97:373–390
18. Fulton AB, Mayer DL. Esotropic children with amblyopia: effects of patching on acuity. *Graefes Arch Clin Exp Ophthalmol.* 1988;226:309–312
19. Buxton MJ, Drummond MF, Van Hout BA, et al. Modelling in economic evaluation: an unavoidable fact of life. *Health Econ.* 1997;6:217–227
20. Consensus Conference on Guidelines on Economic Modelling in Health

- Technology Assessment. Decision analytic modelling in the economic evaluation of health technologies. A consensus statement. *Pharmacoeconomics*. 2000;17:443–444
21. Brennan A, Akehurst R. Modelling in health economic evaluation. What is its place? What is its value? *Pharmacoeconomics*. 2000;17:445–459
 22. Gold M, Siegel J, Russel L, Weinstein M. *Cost-Effectiveness in Health and Medicine*. New York, NY: Oxford University Press; 1996
 23. Attebo K, Mitchell P, Smith W. Visual acuity and the causes of visual loss in Australia. The Blue Mountain Eye Study. *Ophthalmology*. 1996; 103:357–364
 24. Klein R, Klein BE, Linton KL, De Mets DL. The Beaver Dam Eye Study: visual acuity. *Ophthalmology*. 1991;98:1310–1315
 25. West SK, Munoz B, Rubin GS, et al. Function and visual impairment in a population-based study of older adults. The SEE project. Salisbury Eye Evaluation. *Invest Ophthalmol Vis Sci*. 1997;38:72–82
 26. Briggs A, Sculpher M. An introduction to Markov modelling for economic evaluation. *Pharmacoeconomics*. 1998;13:397–409
 27. Briggs AH. Handling uncertainty in cost-effectiveness models. *Pharmacoeconomics*. 2000;17:479–500
 28. Statistisches Bundesamt. *Statistisches Jahrbuch 2002* [English translation: ●●●●]. Stuttgart, Germany: Metzler-Poeschel; 2002
 29. Rüssmann W, König U, Schlimbach K, Pawlowska-Seyda D, Wirbacz B. Brechungsfehler, Schielen und schwachsichtigkeit im vorschulscreening-erfahrungen mit sehstests im kindergarten [English translation: Refractive errors, strabismus and amblyopia in pre-school screening-experiences using a vision test in kindergarten]. *Offentl Gesundheitswes*. 1990;52:77–84
 30. Käsmann-Kellner B, Heine M, Pfau B, Singer A, Ruprecht KW. Screening-untersuchung auf amblyopie, strabismus und refraktionsanomalie bei 1030 kindergartenkindern [English translation: Screening for amblyopia, strabismus and refractive abnormalities in 1,030 kindergarten children]. *Klin Monatsbl Augenheilkd*. 1998;213:166–173
 31. Pratt JW, Raiffa H, Schlaifer R. *Introduction to Statistical Decision Theory*. Cambridge, MA: MIT Press; 1995
 32. Williamson TH, Andrews R, Dutton GN, Murray G, Graham N. Assessment of an inner city visual screening programme for preschool children. *Br J Ophthalmol*. 1995;79:1068–1073
 33. Beardsell R, Clarke S, Hill M. Outcome of occlusion treatment for amblyopia. *J Pediatr Ophthalmol Strabismus*. 1999;36:19–24
 34. Latvala ML, Paloheimo M, Karma A. Screening of amblyopic children and long-term follow-up. *Acta Ophthalmol Scand*. 1996;74:488–492
 35. Leiba H, Shimshoni M, Oliver M, Gottesman N, Levartovsky S. Long-term follow-up of occlusion therapy in amblyopia. *Ophthalmology*. 2001; 108:1552–1555
 36. The Pediatric Eye Disease Investigator Group. A randomized trial of atropine vs. patching for treatment of moderate amblyopia in children. *Arch Ophthalmol*. 2002;120:268–278
 37. Kleinbaum D, Kupper L, Morgenstern H. *Epidemiologic Research. Principles and Quantitative Methods*. New York, NY: Van Nostrand Reinhold; 1982
 38. Sharma S, Brown GC, Brown MM, et al. Converting visual acuity to utilities. *Can J Ophthalmol*. 2000;35:267–272
 39. Brown GC. Vision and quality-of-life. *Trans Am Ophthalmol Soc*. 1999; 97:473–511
 40. Brown MM, Brown GC, Sharma S, Busbee B, Brown H. Quality of life associated with unilateral and bilateral good vision. *Ophthalmology*. 2001;108:643–764
 41. Bertram B. Patienten in augenärztlichen praxen in Deutschland. Teil 1-Alter, geschlecht und diagnosen [English translation: Patients treated in offices of ophthalmologists in Germany. Part 1-Age, gender and diagnosis]. *Der Augenarzt*. 1999;1:23–28
 42. Bertram B. Patienten in augenärztlichen Praxen in Deutschland. Teil 5-spezialisierung in orthoptik, glaukom- und diabetikerbehandlung [English translation: Patients treated in offices of ophthalmologists in Germany. Part 5-Specialization in orthoptic treatment, treatment of glaucoma, and treatment of diabetic patients]. *Der Augenarzt*. 1999;5: 280–282
 43. Statistisches Bundesamt. *Statistisches Jahrbuch 1999* [English translation: Statistical yearbook]. Stuttgart, Germany: Metzler-Poeschel; 1999
 44. Attebo K, Mitchell P, Cumming R, Smith W, Jolly N, Sparkes R. Prevalence and causes of amblyopia in an adult population. *Ophthalmology*. 1998;105:154–159
 45. Miller DK, Homan SM. Determining transition probabilities: confusion and suggestions. *Med Decis Making*. 1994;14:52–58
 46. Vereecken EP, Brabant P. Prognosis for vision in amblyopia after the loss of the good eye. *Arch Ophthalmol*. 1984;102:220–224
 47. Statistisches Bundesamt. *Abgekürzte Sterbetafel 1997/99, Deutschland*. [English translation: Abbreviated life table for Germany 1997/1999]. Wiesbaden, Germany: Statistisches Bundesamt; 2002
 48. Organisation for Economic Co-operation and Development. *OECD Health Data 2001. A Comparative Analysis of 29 Countries*. CD-Rom. Paris, France: Organisation for Economic Co-operation and Development; 2001
 49. König HH, Walter HS, Barry JC. Ressourcenverbrauch und kosten der amblyopiebehandlung [English translation: Resource utilisation and cost of amblyopia treatment]. *Klin Monatsbl Augenheilkd*. 2003;220: 486–491
 50. Kleiter GD. *Bayes Statistik. Grundlagen und Anwendungen* [English translation: *Bayes Statistics. Basic Principles and Applications*]. Berlin, Germany: Walter de Gruyter; 1980
 51. Briggs A, Fenn P. Confidence intervals or surfaces? Uncertainty on the cost-effectiveness plane. *Health Econ*. 1998;7:723–740
 52. Briggs AH, O'Brien BJ, Blackhouse G. Thinking outside the box: advances in the analysis and presentation of uncertainty in cost-effectiveness studies. *Annu Rev Public Health*. 2002;23:377–401
 53. van Hout BA, Al MJ, Gordon GS, Rutten FF. Costs, effects and C/E-ratios alongside a clinical trial. *Health Econ*. 1994;3:309–319
 54. Drummond MF, O'Brien B, Stoddart GL, Torrance GW. *Methods for the Economic Evaluation of Health Care Programmes*. Oxford, United Kingdom: Oxford University Press; 1997
 55. Javitt JC, Aiello LP. Cost-effectiveness of detecting and treating diabetic retinopathy. *Ann Intern Med*. 1996;124:164–169
 56. Simons K, Preslan M. Natural history of amblyopia untreated owing to lack of compliance. *Br J Ophthalmol*. 1999;83:582–587
 57. Weinstein MC, Siegel JE, Gold MR, Kamlet MS, Russell LB. Recommendations of the Panel on Cost-Effectiveness in Health and Medicine. *JAMA*. 1996;276:1253–1258
 58. Preslan MW, Novak A. Baltimore Vision Screening Project. *Ophthalmology*. 1996;103:105–109

Cost-Utility Analysis of Orthoptic Screening in Kindergarten: A Markov Model Based on Data From Germany

Hans-Helmut König and Jean-Cyriaque Barry

Pediatrics 2004;113:e95

DOI: 10.1542/peds.113.2.e95

Updated Information & Services

including high resolution figures, can be found at:
<http://pediatrics.aappublications.org/content/113/2/e95>

References

This article cites 48 articles, 10 of which you can access for free at:
<http://pediatrics.aappublications.org/content/113/2/e95#BIBL>

Subspecialty Collections

This article, along with others on similar topics, appears in the following collection(s):
Ophthalmology
http://www.aappublications.org/cgi/collection/ophthalmology_sub

Permissions & Licensing

Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at:
<http://www.aappublications.org/site/misc/Permissions.xhtml>

Reprints

Information about ordering reprints can be found online:
<http://www.aappublications.org/site/misc/reprints.xhtml>

American Academy of Pediatrics

DEDICATED TO THE HEALTH OF ALL CHILDREN™



PEDIATRICS®

OFFICIAL JOURNAL OF THE AMERICAN ACADEMY OF PEDIATRICS

Cost-Utility Analysis of Orthoptic Screening in Kindergarten: A Markov Model Based on Data From Germany

Hans-Helmut König and Jean-Cyriaque Barry

Pediatrics 2004;113:e95

DOI: 10.1542/peds.113.2.e95

The online version of this article, along with updated information and services, is located on the World Wide Web at:

<http://pediatrics.aappublications.org/content/113/2/e95>

Pediatrics is the official journal of the American Academy of Pediatrics. A monthly publication, it has been published continuously since 1948. Pediatrics is owned, published, and trademarked by the American Academy of Pediatrics, 141 Northwest Point Boulevard, Elk Grove Village, Illinois, 60007. Copyright © 2004 by the American Academy of Pediatrics. All rights reserved. Print ISSN: 1073-0397.

American Academy of Pediatrics

DEDICATED TO THE HEALTH OF ALL CHILDREN™

