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*Cochrane Database of Systematic Reviews* 2012, Issue 6. Art. No.: CD003433.

DOI: 10.1002/14651858.CD003433.pub3.

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Beta radiation for glaucoma surgery (Review)

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[Intervention Review]

# Beta radiation for glaucoma surgery

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**Editorial group:** Cochrane Eyes and Vision Group.

**Publication status and date:** New search for studies and content updated (no change to conclusions), published in Issue 6, 2012.

**Review content assessed as up-to-date:** 26 March 2012.

**Citation:** Kirwan JF, Rennie C, Evans JR. Beta radiation for glaucoma surgery. *Cochrane Database of Systematic Reviews* 2012, Issue 6. Art. No.: CD003433. DOI: 10.1002/14651858.CD003433.pub3.

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## ABSTRACT

### Background

The outcome of glaucoma surgery can be affected by the rate at which the surgical wound heals. Beta radiation has been proposed as a rapid and simple treatment to slow down the healing response.

### Objectives

To assess the effectiveness of beta radiation during glaucoma surgery (trabeculectomy).

### Search methods

We searched CENTRAL (which contains the Cochrane Eyes and Vision Group Trials Register) (*The Cochrane Library* 2012, Issue 3), MEDLINE (January 1950 to March 2012), EMBASE (January 1980 to March 2012), the *metaRegister* of Controlled Trials (*mRCT*) ([www.controlled-trials.com](http://www.controlled-trials.com)), ClinicalTrials.gov ([www.clinicaltrials.gov](http://www.clinicaltrials.gov)) and the WHO International Clinical Trials Registry Platform (ICTRP) ([www.who.int/ictrp/search/en](http://www.who.int/ictrp/search/en)). There were no date or language restrictions in the electronic searches for trials. The electronic databases were last searched on 26 March 2012.

### Selection criteria

We included randomised controlled trials comparing trabeculectomy with beta radiation to trabeculectomy without beta radiation.

### Data collection and analysis

We collected data on surgical failure (intraocular pressure > 21 mmHg), intraocular pressure and adverse effects of glaucoma surgery. We pooled data using a fixed-effect model.

### Main results

We found four trials that randomised 551 people to trabeculectomy with beta irradiation versus trabeculectomy alone. Two trials were in Caucasian people (126 people), one trial in black African people (320 people) and one trial in Chinese people (105 people). People who had trabeculectomy with beta irradiation had a lower risk of surgical failure compared to people who had trabeculectomy alone (pooled risk ratio (RR) 0.23 (95% CI 0.14 to 0.40)). Beta irradiation was associated with an increased risk of cataract (RR 2.89, 95% CI 1.39 to 6.0).

### **Authors' conclusions**

Trabeculectomy with beta irradiation has a lower risk of surgical failure compared to trabeculectomy alone. A trial of beta irradiation versus anti-metabolite is warranted.

## **PLAIN LANGUAGE SUMMARY**

### **Beta radiation in glaucoma surgery**

The aim of glaucoma surgery is to lower the pressure in the eye. The outcome of glaucoma surgery can be affected by the rate at which the surgical wound heals. Beta radiation has been proposed as a rapid and simple treatment to slow down the healing response. It is applied during the operation using a radioactive applicator which emits beta rays which have only a very local penetration to a depth of less than one millimetre. The intensity of the emission from the applicator (usually Strontium-90) determines the duration it is applied to the surgical site in order to deliver the required dose of radiation which effectively prevents scar tissue formation.

We found four trials that randomised 551 people to trabeculectomy with beta irradiation versus trabeculectomy alone. People who had trabeculectomy with beta irradiation were less likely to have an eye pressure that was too high one year after surgery compared to people who had trabeculectomy alone. However, people who had beta irradiation had an increased risk of cataract after surgery.

## SUMMARY OF FINDINGS FOR THE MAIN COMPARISON *[Explanation]*

Beta radiation for glaucoma surgery						
<b>Patient or population:</b> patients with glaucoma surgery <b>Settings:</b> hospitals in low, middle or high income countries <b>Intervention:</b> beta radiation						
Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Control	beta radiation				
Surgical failure (intraocular pressure >21mmHg) (follow-up: 12 months)	Low risk population <sup>1</sup>		RR 0.23 (0.14 to 0.4)	561 (4)	⊕⊕⊕○ moderate <sup>2</sup>	
	100 per 1000	23 per 1000 (14 to 40)				
	High risk population <sup>1</sup>					
	300 per 1000	69 per 1000 (42 to 120)				
Intraocular pressure (follow-up: 12 months)	The mean intraocular pressure in the control groups was <b>13.85 mmHg</b>	The mean Intraocular pressure in the intervention groups was <b>0.97 lower</b> (2.56 lower to 0.62 higher)		123 (2)	⊕⊕○○ low <sup>3,4</sup>	
Adverse effect: cataract	Medium risk population		RR 1.79 (0.62 to 5.14)	423 (3)	⊕⊕⊕○ moderate <sup>5,6</sup>	
	50 per 1000	89 per 1000 (31 to 257)				

\*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; RR: Risk ratio;

GRADE Working Group grades of evidence

**High quality:** Further research is very unlikely to change our confidence in the estimate of effect.

**Moderate quality:** Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

**Low quality:** Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

**Very low quality:** We are very uncertain about the estimate.

<sup>1</sup> Based on control events in included studies. Higher risks of surgical failure seen in black African populations.

<sup>2</sup> Majority of evidence from larger high quality study in black African people. However, relative risks for smaller studies in Caucasian and Chinese people were of a similar order but imprecise (pooled relative risk 0.15 (95% CI 0.02 to 1.28)).

<sup>3</sup> Data for Caucasian patients only.

<sup>4</sup> Two studies contributed data to this outcome, n=123 people. Pooled mean difference was -0.97 (-2.56, 0.62). i.e. beta radiation compatible with a reduction of approximately 2.5mmHg intraocular pressure but also with an increased pressure of less than 1mmHg.

<sup>5</sup> Barnes 2000 and Kirwan 2006 had similar estimates of the risk of cataract associated with beta irradiation of approximately 3 to 3.5. In contrast Rehman 2002 had a relative risk reduction of 50%. However, there were small numbers of events (2 cataracts in each group) in Rehman 2002.

<sup>6</sup> The pooled analysis had a total of 9 events in the treatment group and 4 in the control group.

## BACKGROUND

Beta radiation inhibits wound healing, and has been shown to improve survival of glaucoma filtration surgery in vitro and in vivo (Khaw 1991). Beta radiation causes growth-arrest, primarily due to its effects on P450 (a controller of cell cycling), rather than producing destruction of cells (Constable 1999).

### Description of the condition

The term glaucoma defines a group of conditions in which there is characteristic optic neuropathy associated with progressive visual field loss (loss of peripheral and then central vision). Raised intraocular pressure (IOP) is recognised to be one of the principal risk factors for glaucoma (Buhmann 2000; Hollows 1966; Leske 1995). Glaucoma is the commonest cause of irreversible blindness worldwide, with at least 70 million people affected and over seven million blind from the condition (Quigley 1996). With an ageing world population, the prevalence is likely to increase considerably in the coming years.

### Description of the intervention

Lowering IOP has long been established as a means of treating glaucoma, despite a lack of good data for its effectiveness in preventing visual field loss (Rossetti 1993; Vass 2007). What information we do have suggests that surgery is currently the most effective means of lowering IOP (Burr 2004; Migdal 1986, Lichter 2001). Trabeculectomy is the most widely used surgical procedure for glaucoma. In the developed world, surgery is used where other modalities (medical or laser therapy or both) are not sufficiently effective. In poorer countries trabeculectomy may be used as a primary treatment for glaucoma.

The aim of the trabeculectomy operation is to create a fistula (channel) between the anterior chamber and the subconjunctival space to allow controlled release of fluid from the eye. A partial-thickness scleral flap is fashioned at the limbus, with a sclerostomy created under this flap forming a so-called guarded fistula. Fluid from the anterior chamber drains out of the eye through this fistula to the subconjunctival space forming a conjunctival bleb. Fluid in this bleb is thought to be carried away by vessels within the conjunctiva and also by passing through the conjunctival wall. Final IOP may be determined by the size of the bleb, downstream drainage from the conjunctival vessels and the healing response of the eye. The most common cause for failure of trabeculectomy is subconjunctival fibrosis (Hitchings 1983) resulting in occlusion of the fistula with loss of IOP lowering effect.

In elderly white people with glaucoma who have not had prior eye surgery, the success rate of surgery is considered to be in the region of 80%. However, the success rate is lower in certain groups including black people, people with ocular inflammation or neovascularisation, those who have had previous ocular surgery and

children (Ritch 1996). It is also apparent that prior topical medical treatment may prejudice the overall success of surgery (Broadway 1994; Lavin 1990).

Various agents have been used to reduce the excessive healing that leads to failure of trabeculectomy. The most widely used agents are Mitomycin C and 5-Fluorouracil. Mitomycin C was introduced in 1983 although widespread use did not occur until the mid 1990s (Chen 1983). 5-Fluorouracil was introduced in 1984, after positive reports from a randomised controlled trial (RCT), and was used widely until use of Mitomycin C became widespread (FFSG 1996; Heuer 1984; Singh 2000). Systematic reviews of postoperative 5-Fluorouracil (Wormald 2001) and of intraoperative Mitomycin C (Wilkins 2005) are already published in *The Cochrane Library*.

Although both of these antimetabolites have proven effective clinically they are associated with complications. Both Mitomycin C and 5-Fluorouracil are used in liquid form, delivered by placing microsurgical sponges soaked in the drug directly onto the operative site. Because the antimetabolites are liquids they carry the risk of leakage away from the treatment site. This leakage can lead to extraocular or, more seriously, intraocular toxicity (Franks 1991). Furthermore, the variability of the delivery of the drug between the impregnated sponge and the subconjunctival tissues means that accurate dosimetry has proven difficult. Both treatments have been associated with the development of thin, avascular filtration blebs, and these are in turn associated with an increased risk of sight threatening complications including hypotony (a very soft eye) and endophthalmitis (ocular infection) (Wolner 1991; Yaldo 1993; Zacharia 1993).

One alternative to the use of antimetabolites is beta radiation. Precise control of dose and area of treatment may be easily achieved as it is applied using a Strontium-90 applicator at the completion of surgery. The doses used are typically 750 cGy or 1000 cGy and single application is used without fractionation. Beta radiation was first used for glaucoma surgery in the 1940s (Liff 1944) and has been used at Moorfields Eye Hospital for many years in the management of paediatric glaucoma. Good results were reported with a non-randomised retrospective study in a cohort of these patients (Miller 1991). Beta radiation may be especially suitable for use in developing countries due to the technical simplicity of application and as no running costs are required.

### Why it is important to do this review

Recently, several workers have described the use of beta radiation with trabeculectomy. The overall effectiveness of this mode of therapy is unknown and a systematic review is needed to assess the effects both in people where a high success rate is expected and in those where a lower success rate would be expected.

## OBJECTIVES

The aims of this review were to assess the effects of beta radiation with trabeculectomy on surgical failure due to postoperative scarring in people with glaucoma. Mean IOP was an additional primary outcome. The review also aimed to gather evidence on whether beta radiation is associated with increased complications after surgery as compared to standard trabeculectomy.

## METHODS

### Criteria for considering studies for this review

#### Types of studies

We included randomised controlled trials (RCTs).

#### Types of participants

We included trials in which participants were people with glaucoma undergoing trabeculectomy. All glaucoma diagnosis categories were included. We included people undergoing a first surgical procedure without any perceived risk factors for surgical failure and people who were considered to have a high risk of surgical failure. We did not include people undergoing simultaneous bilateral surgery.

#### Types of interventions

We included trials in which trabeculectomy with beta radiation applied at any dose was compared to trabeculectomy alone, with placebo or with another anti-scarring agent. We excluded trials in which trabeculectomy was combined with cataract extraction.

#### Types of outcome measures

##### Primary outcomes

The primary outcome measures for this review were:

1. the proportion of failed trabeculectomies at 12 months or more after surgery (defined as repeat surgery or uncontrolled IOP more than 20 mmHg with or without additional topical or systemic medications);
2. mean IOP at 12 months or more. Intraocular pressure is used as a primary outcome measure for many clinical trials in glaucoma. The ideal outcome measure is visual function, most commonly expressed in terms of visual field (with allowance for visual acuity). However, the difficulty in determining visual field progression due to the inherently high noise in this psychophysical measure means that trials using this outcome measure need to be performed with large sample sizes and long follow-up time. For practical reasons, IOP was used as a surrogate measurement; the degree of IOP

reduction appears to be related to the rate of visual field progression (Gaasterland 2000).

##### Secondary outcomes

Secondary outcome measures included:

3. adverse effects including:
  - a. visual loss,
  - b. wound leaks: the presence of a positive Seidel test (visible aqueous flow with the tear film stained with fluorescein),
  - c. late endophthalmitis - an infection of the globe contents that even with prompt aggressive treatment often results in substantial loss of visual function. 'Late' here implies infection arising from organisms gaining access to the globe through thin walled drainage blebs or frank breaks in the conjunctival epithelium after the immediate postoperative period when infectious agents may have entered the eye during the surgical procedure,
  - d. hypotony: the IOP is below five millimetres mercury and/or associated with complications such as macular oedema and sight loss or choroidal detachments,
  - e. cataract: the subsequent development of visually significant cataract.
4. outcomes relating to quality of life or patient's perspective of care;
5. data relevant to economic evaluation.

### Search methods for identification of studies

#### Electronic searches

We searched the Cochrane Central Register of Controlled Trials (CENTRAL) 2012, Issue 3, part of *The Cochrane Library*. [www.thecochranelibrary.com](http://www.thecochranelibrary.com) (accessed 26 March 2012), MEDLINE (January 1950 to March 2012), EMBASE (January 1980 to March 2012), the *metaRegister* of Controlled Trials (*mRCT*) ([www.controlled-trials.com](http://www.controlled-trials.com)), ClinicalTrials.gov ([www.clinicaltrials.gov](http://www.clinicaltrials.gov)) and the WHO International Clinical Trials Registry Platform (ICTRP) ([www.who.int/ictrp/search/en](http://www.who.int/ictrp/search/en)). There were no language or date restrictions in the search for trials. The electronic databases were last searched on 26 March 2012. See: Appendices for details of search strategies for CENTRAL (Appendix 1), MEDLINE (Appendix 2), EMBASE (Appendix 3), *mRCT* (Appendix 4), ClinicalTrials.gov (Appendix 5) and the ICTRP (Appendix 6).

#### Searching other resources

We manually searched the Association for Research in Vision and Ophthalmology (ARVO) abstract books from 1990 to 2009 using a keyword index search for relevant studies. Keywords used were: radiation; beta radiation; strontium. All studies with these words in the permuted index were examined to see if they were relevant. We



contacted researchers who are active in the field and manufacturers of the emitter for information on further published or unpublished studies.

## Data collection and analysis

### Selection of studies

Two authors independently screened the titles and abstracts obtained by the searches and full copies of definitely or potentially relevant studies were obtained. Where the trial had not been fully reported, or was unreported, we attempted to contact the investigators and endeavour to obtain as much relevant data as possible. Details of the trial were not masked during the assessment.

### Data extraction and management

One review author entered the data into RevMan 5 (Review Manager 2011) and a second review author independently verified the data entry. Any disagreements between review authors were resolved by discussion.

### Assessment of risk of bias in included studies

Relevant details about study design were extracted independently by two authors. Risk of bias was assessed using The Cochrane Collaboration's tool for assessing risk of bias as described in Chapter 8 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We contacted study authors for further information on any item graded as unclear.

### Data synthesis

For description of dichotomous data such as failure rates, a summary odds ratio was calculated. For IOP, the weighted mean difference was reported. As there were three studies or less to be combined, we used a fixed-effect model. If further trials are included in future updates of this review we will use a random-effects model. There were not enough published studies to investigate potential publication bias nor to conduct sensitivity analyses as planned in the protocol (excluding poorer quality studies).

### Subgroup analysis and investigation of heterogeneity

No subgroup analyses were planned in the protocol. The included trials were conducted in different populations (black African, Caucasian and Chinese). There is some evidence that people of African origin have a higher risk of surgical failure (AGIS 2001). In the included trials there was some evidence that the control group risk of failure of surgery was different in different ethnic groups (see Table 1). We therefore did a (posthoc) subgroup analysis by ethnic group.

## RESULTS

### Description of studies

#### Results of the search

The electronic searches identified 1180 titles and abstracts. We screened the title and abstracts and obtained full-text copies of five reports to assess for potential inclusion in the review. Four studies were eligible for inclusion and one study was excluded, see the 'Characteristics of excluded studies' table for further information. An update search was done in March 2012 which yielded a further 93 references. We screened the search results but found no new studies which met the inclusion criteria.

#### Included studies

See Table 1 and 'Characteristics of included studies' for further details. Four trials were found (Barnes 2000; Lai 1994; Rehman 2002; Kirwan 2006). Two trials (Barnes 2000; Rehman 2002) were performed in Caucasian people, with primary open angle glaucoma with an expected high success rate. Lai 1994 was found in abstract form, which had been performed in Chinese people with open angle glaucoma. The success rate in this trial was also expected to be high, although not as high as for Caucasian people. One trial was performed in black African patients (Kirwan 2006). Two trials used a dosage of 750 cGy (Barnes 2000; Rehman 2002) and two trials used a dosage of 1000 cGy (Kirwan 2006; Lai 1994). A total of 551 people were randomised in these four trials. Follow-up ranged from 12 months (Rehman 2002) to four years (Kirwan 2006).

#### Excluded studies

See Characteristics of excluded studies for details.

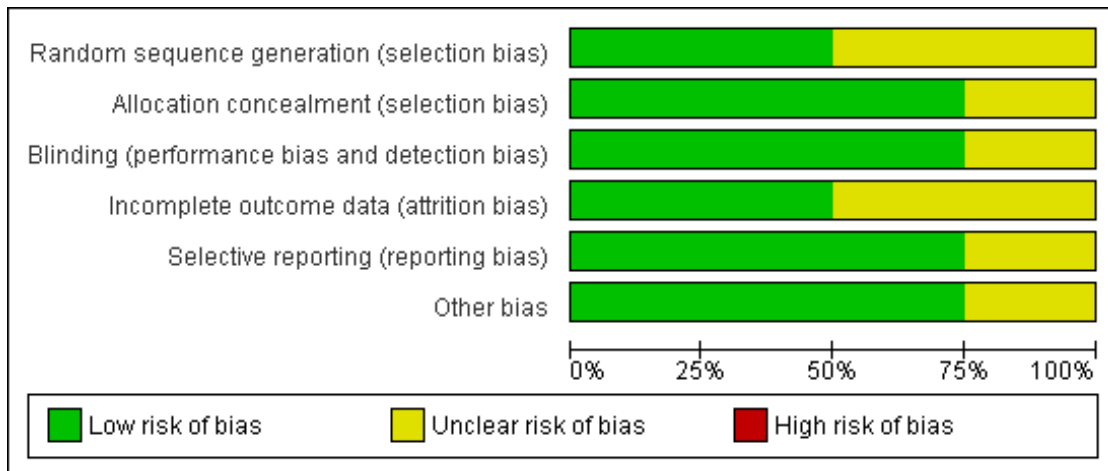
#### Risk of bias in included studies

See Figure 1 and Figure 2. The trials reported as research papers were of acceptable methodological quality (Barnes 2000; Kirwan 2006; Rehman 2002). In all three trials, sealed envelopes with a randomised number sequence were used for allocation concealment. Examiners determining outcome were unaware of trial allocation. The study published in abstract form has not been subsequently published but we were able to get some information from the trial investigator who reported adequate allocation concealment. Intention-to-treat analysis was not formally reported in any trial. In all trials, only one eye was entered into the trial.

Figure 1. Methodological quality summary: review authors' judgements about each methodological quality item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding (performance bias and detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Barnes 2000	?	?	+	?	+	+
Kirwan 2006	+	+	+	+	+	+
Lai 1994	?	+	?	?	?	?
Rehman 2002	+	+	+	+	+	+

**Figure 2. Methodological quality graph: review authors' judgements about each methodological quality item presented as percentages across all included studies.**



### Effects of interventions

See: [Summary of findings for the main comparison](#)

People who had trabeculectomy with beta irradiation had a lower risk of surgical failure one year after surgery compared to people who had trabeculectomy alone ([Analysis 1.1](#) pooled risk ratio 0.23 (95% confidence interval (CI) 0.14 to 0.40)). The treatment effect appeared to be similar in the different ethnic groups. However the three trials in non African people were smaller and the risk of surgical failure lower in Caucasian people. In the two trials of Caucasian people there were no cases of surgical failure in the treatment group. This analysis, therefore, does not have good power to detect true differences in treatment effect in the different ethnic groups. Only two trials reported mean IOP at the end of the study ([Analysis 2.1](#)). People receiving beta irradiation had on average 1 mmHg lower IOP 12 months after surgery. However, this result was not statistically significant and the result is compatible with both lower and higher IOP after surgery (pooled mean difference -0.97 95% CI -2.56 to 0.62).

The most recent study conducted on a South African population was the largest with the greatest power to detect a difference. In this study relative risk of failure was 0.21 (95% CI 0.11 to 0.40) at one year ([Kirwan 2006](#)).

The two trials reporting results in Caucasians both had very high success rates. The study from New Zealand, [Barnes 2000](#) showed a small trend for people having beta radiation to have a higher success rate, a lower IOP and a larger drop in IOP. None of these results were significant. The study from Leeds, [Rehman 2002](#) had success rates of 95% and 100% in each arm of the trial for the most generous definition of success and high success rates, even for more stringent definitions. There was no significant difference

in success rates between the two arms of the trial for any outcome measure. Neither study had sufficient power to compare adverse effect rates. The Hong Kong study, [Lai 1994](#) demonstrated a significant difference in success between the two arms of the study. The failure rate was 25% in the control arm and 9% in the treatment arm.

Adverse effects were reported in all studies but numbers were low. There was evidence of an increased risk of cataract requiring surgical intervention in the treatment group ([Analysis 3.1](#) pooled risk ratio 2.89 (95% CI 1.39 to 6.00)). Numbers of other complications were too low to comment on ([Analysis 3.2](#); [Analysis 3.3](#)), although there was a suggestion that beta radiation augmented trabeculectomy was associated with a small increase in the risk of ocular hypotony ([Analysis 3.2](#)).

### DISCUSSION

Several points are worthy of comment in considering the data. In the [Barnes 2000](#) study, four participants had only three months follow-up. We were able to obtain the raw data from the investigators. One of those four participants had failed surgery with an IOP 21mmHg or more at three months. We have counted this patient as a surgical failure at 12 months as well. For the other three patients who had an IOP less than 21mmHg at three months we cannot make any assumptions about their status at 12 months. The dose used for both the [Rehman 2002](#) and [Barnes 2000](#) studies was 750 cGy, this is less than the dose used in [Lai 1994](#) and cell culture and in vivo animal data suggests that the difference may be a biologically significant one. In the [Lai](#) study, the numbers

were reported as percentages only and no further information was obtained from the study author.

From the limited data available, it appears that beta radiation improves success rates in people with an intermediate surgical prognosis and possibly in those with an already good surgical prognosis. This may be at the cost of an increased rate of cataract development.

The degree of improvement in success is similar to that of other antimetabolites but there is no trial data comparing beta radiation with other metabolites such as Mitomycin C or 5-Fluorouracil.

In a developing world setting, with a high surgical volume, beta radiation has advantages in that there are no ongoing costs and it has a long service life (more than 20 years) although capital outlay is significant (approximately £5000 to purchase a Sr90 emitter in the UK). A further issue is that in the UK and many other countries, a government licence is required to use a radioactive device. Obtaining this may necessitate extra training; this and associated bureaucracy may place a barrier on the use of an emitter. In the UK, storage is in a lead lined repository, usually in the medical physics department. However, this may be excessive given the attenuation of Sr90 in air and storage in an area with a significant physical distance from those in the working environment is considered to be adequate in many settings.

## AUTHORS' CONCLUSIONS

### Implications for practice

Beta radiation-augmented trabeculectomy reduces the risk of surgical failure compared to standard trabeculectomy. It may be clinically useful but it is unknown how it compares in both effectiveness and safety to other, more widely used antimetabolites. Data on adverse effects are limited. A specific role for beta radiation is

unclear and this depends on there being more evidence available to guide practice.

### Implications for research

There are only four trials of beta radiation for increasing the probability of success in trabeculectomy. Much of the existing evidence for controlling wound healing after trabeculectomy concentrates on liquid antimetabolites. How beta radiation compares to liquid antimetabolites is unknown. Beta radiation may be particularly useful in a developing world setting but this has yet to be evaluated. Outcomes of particular interest are the degree of IOP lowering, success rates in the long term, and adverse effects - particularly cataract. Further trials comparing beta radiation with other antimetabolites would be useful in such a setting but may also be of relevance to richer countries.

## ACKNOWLEDGEMENTS

We are grateful to the peer reviewers including David Broadway, Catey Bunce and Suzanne Brodney-Folse for comments on this review and to investigators who shared their data with us. The Cochrane Eyes and Vision Group created and executed the search results. We thank Anupa Shah and Iris Gordon for their help with the review process.

Richard Wormald (Co-ordinating Editor for CEVG) acknowledges financial support for his CEVG research sessions from the Department of Health through the award made by the National Institute for Health Research to Moorfields Eye Hospital NHS Foundation Trust and UCL Institute of Ophthalmology for a Specialist Biomedical Research Centre for Ophthalmology. The views expressed in this publication are those of the authors and not necessarily those of the Department of Health.

## REFERENCES

### References to studies included in this review

#### Barnes 2000 *{published data only}*

\* Barnes RM, Mora JS, Best SJ. Beta radiation as an adjunct to low-risk trabeculectomy. *Clinical and Experimental Ophthalmology* 2000;**28**(4):259–62.

#### Kirwan 2006 *{published data only}*

\* Kirwan JF, Cousens S, Venter L, Cook C, Stulting A, Roux P, et al. Effect of beta radiation on success of glaucoma drainage surgery in South Africa: randomised controlled trial. *BMJ* 2006;**333**(7575):942.

#### Lai 1994 *{published data only}*

Lai JSM, Ho PCP. Trabeculectomy combined with beta irradiation in uncomplicated primary open angle glaucoma.

*Investigative Ophthalmology and Visual Science* 1994;**35**: Abstract number 825.

#### Rehman 2002 *{published data only}*

Rehman SU, Amoaku WM, Doran RM, Menage MJ, Morrell AJ. Randomized controlled clinical trial of beta irradiation as an adjunct to trabeculectomy in open-angle glaucoma. *Ophthalmology* 2002;**109**(2):302–6.

### References to studies excluded from this review

#### Tesha 2002 *{published data only}*

Tesha PE, Rehman SU, Amoaku WMK, Doran RML, Menage MJ, Morrell AJ. Investigation into the use of beta irradiation as an adjunct to trabeculectomy in

open angle glaucoma. Congress of the Royal College of Ophthalmologists. 2002.

## Additional references

### AGIS 2001

AGIS Investigators. The Advanced Glaucoma Intervention Study (AGIS): 9. Comparison of glaucoma outcomes in black and white patients within treatment groups. *American Journal of Ophthalmology* 2001;**132**(3):311–20.

### Broadway 1994

Broadway DC, Grierson I, O'Brien C, Hitchings RA. Adverse effects of topical antiglaucoma medication regimes: Part II. Effect on the outcome of filtration surgery. *Archives of Ophthalmology* 1994;**112**:1446–54.

### Buhrmann 2000

Buhrmann RR, Quigley HA, Barron Y, West SK, Oliva MS, Mmbaga BB. Prevalence of glaucoma in a rural East African population. *Investigative Ophthalmology and Visual Science* 2000;**41**(1):40–8.

### Burr 2004

Burr J, Azuara-Blanco A, Avenell A. Medical versus surgical interventions for open angle glaucoma. *Cochrane Database of Systematic Reviews* 2004, Issue 2. [DOI: 10.1002/14651858.CD004399.pub2]

### Chen 1983

Chen CW. Enhanced intraocular pressure controlling effectiveness of trabeculectomy by local application of mitomycin C. *Transactions of the Congress of the Asia-Pacific Academy of Ophthalmology* 1983:172.

### Constable 1999

Constable PH. *The effect of single doses of beta radiation on the wound healing functions of ocular fibroblasts [MD thesis]*. Cambridge: University of Cambridge, 1999.

### FFSG 1996

The Fluorouracil Filtering Surgery Study Group. Five-year follow-up of the Fluorouracil Filtering Surgery Study. *American Journal of Ophthalmology* 1996;**121**(4):349–66.

### Franks 1991

Franks WA, Hitchings RA. Complications of 5-fluorouracil after trabeculectomy. *Eye* 1991;**5**(Pt 4):385–9.

### Gaasterland 2000

Gaasterland DE, Ederer F, Beck A, Costarides A, Leef D, Closek J, et al. The AGIS Investigators. The advanced glaucoma intervention study (AGIS): 7. The relationship between control of intraocular pressure and visual field deterioration. *American Journal of Ophthalmology* 2000;**130**(4):429–40.

### Glanville 2006

Glanville JM, Lefebvre C, Miles JN, Camosso-Stefinovic J. How to identify randomized controlled trials in MEDLINE: ten years on. *Journal of the Medical Library Association* 2006;**94**(2):130–6.

### Heuer 1984

Heuer DK, Parrish RK 2d, Gressel MG, Hodapp E, Palmberg PF, Anderson DR. 5-fluorouracil and glaucoma

filtering surgery. II. A pilot study. *Ophthalmology* 1984;**91**(4):384–94.

### Higgins 2011

Higgins JPT, Altman DG, Sterne JAC (editors). Chapter 8: Assessing risk of bias in included studies. In: Higgins JPT, Green S (editors). *Cochrane Handbook for Systematic Reviews of Interventions* Version 5.1.0 (updated March 2011). The Cochrane Collaboration, 2011. Available from www.cochrane-handbook.org.

### Hitchings 1983

Hitchings RA, Grierson I. Clinico pathological correlation in eyes with failed fistulizing surgery. *Transactions of the Ophthalmological Societies of the United Kingdom* 1983;**103**(Pt 1):84–8.

### Hollows 1966

Hollows FC, Graham PA. Intra-ocular pressure, glaucoma, and glaucoma suspects in a defined population. *British Journal of Ophthalmology* 1966;**50**(10):570–86.

### Iliff 1944

Iliff CE. Surgical control of glaucoma in the negro. *American Journal of Ophthalmology* 1944;**27**:731–8.

### Khaw 1991

Khaw PT, Ward S, Grierson I, Rice NS. Effect of beta radiation on proliferating human Tenon's capsule fibroblasts. *British Journal of Ophthalmology* 1991;**75**(10):580–3.

### Lavin 1990

Lavin MJ, Wormald RP, Migdal CS, Hitchings RA. The influence of prior therapy on the success of trabeculectomy. *Archives of Ophthalmology* 1990;**108**(11):1543–8.

### Leske 1995

Leske MC, Connell AM, Wu SY, Hyman LG, Schachat AP. Risk factors for open-angle glaucoma. The Barbados Eye Study. *Archives of Ophthalmology* 1995;**113**(7):918–24.

### Lichter 2001

Lichter PR, Musch DC, Gillespie BW, Guire KE, Janz NK, Wren PA, et al. Interim clinical outcomes in the Collaborative Initial Glaucoma Treatment Study comparing initial treatment randomized to medications or surgery. *Ophthalmology* 2001;**108**(11):1943–53.

### Migdal 1986

Migdal C, Hitchings R. Control of chronic simple glaucoma with primary medical, surgical and laser treatment. *Transactions of the Ophthalmological Societies of the United Kingdom* 1986;**105**(Pt 6):653–6.

### Miller 1991

Miller MH, Rice NS. Trabeculectomy combined with beta irradiation for congenital glaucoma. *British Journal of Ophthalmology* 1991;**75**(10):584–90.

### Quigley 1996

Quigley HA. Number of people with glaucoma worldwide. *British Journal of Ophthalmology* 1996;**80**(5):389–93.

### Review Manager 2011 [Computer program]

The Nordic Cochrane Centre, The Cochrane Collaboration. Review Manager (RevMan). Version 5.1. Copenhagen:

The Nordic Cochrane Centre, The Cochrane Collaboration, 2011.

**Ritch 1996**

Ritch R, Shields MB, Krupin T. *The Glaucomas*. St Louis: Mosby, 1996.

**Rossetti 1993**

Rossetti L, Orzalesi N, Liberati A. The medical treatment of open angle glaucoma. *New England Journal of Medicine* 1993;**329**(10):735–6.

**Singh 2000**

Singh K, Mehta K, Shaikh NM, Tsai JC, Moster MR, Budenz DL, et al. Trabeculectomy with intraoperative mitomycin C versus 5- fluorouracil. Prospective randomized clinical trial. *Ophthalmology* 2000;**107**(12):2305–9.

**Vass 2007**

Vass C, Hirn C, Sycha T, Findl O, Sacu S, Bauer P, Schmetterer L. Medical interventions for primary open angle glaucoma and ocular hypertension. *Cochrane Database of Systematic Reviews* 2007, Issue 4. [DOI: 10.1002/14651858.CD003167.pub3]

**Wilkins 2005**

Wilkins M, Indar A, Wormald R. Intraoperative Mitomycin C for glaucoma surgery. *Cochrane Database of Systematic Reviews* 2005, Issue 4. [DOI: 10.1002/14651858.CD002897.pub2]

**Wolner 1991**

Wolner B, Liebmann JM, Sassani JW, Ritch R, Speaker M, Marmor M. Late bleb-related endophthalmitis after trabeculectomy with adjunctive 5-fluorouracil. *Ophthalmology* 1991;**98**(7):1053–60.

**Wormald 2001**

Wormald R, Wilkins M, Bunce C. Postoperative 5-Fluorouracil for glaucoma surgery. *Cochrane Database of Systematic Reviews* 2001, Issue 3. [DOI: 10.1002/14651858.CD001132]

**Yaldo 1993**

Yaldo MK, Stamper RL. Long-term effects of mitomycin on filtering blebs. Lack of fibrovascular proliferative response following severe inflammation. *Archives of Ophthalmology* 1993;**111**(6):824–6.

**Zacharia 1993**

Zacharia PT, Deppermann SR, Schuman JS. Ocular hypotony after trabeculectomy with mitomycin C. *American Journal of Ophthalmology* 1993;**116**(3):314–26.

**References to other published versions of this review**

**Kirwan 2009**

Kirwan JF, Rennie C, Evans JR. Beta radiation for glaucoma surgery. *Cochrane Database of Systematic Reviews* 2009, Issue 2. [DOI: 10.1002/14651858.CD003433.pub2]

\* Indicates the major publication for the study

## CHARACTERISTICS OF STUDIES

### Characteristics of included studies [ordered by study ID]

#### Barnes 2000

Methods	sealed envelopes, randomised number sequence, 3 lost to FU	
Participants	Caucasian > 45	
Interventions	trabeculectomy +/- beta radiation	
Outcomes	IOP < 21 = success (+/- medication) mean IOP	
Notes		
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Random sequence generation (selection bias)	Unclear risk	"The patients randomly received or did not receive a dose of beta radiation" Page 259, last paragraph
Allocation concealment (selection bias)	Unclear risk	-
Blinding (performance bias and detection bias) All outcomes	Low risk	"There was no record of beta radiation treatment in the patient notes, so at follow up the examining doctor did not know which arm of the study the patient belonged to" Bottom of page 259 top of page 260 We have made the assumption that the patient did not know either although this was not explicitly stated
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	"...follow-up periods ranged between 3 and 42 months. Three patients only had 3 months follow-up.." Page 261, first paragraph However, follow-up times for each treatment group were not given
Selective reporting (reporting bias)	Low risk	The main outcome reported in this study was intraocular pressure, as would be expected
Other bias	Low risk	-

## Kirwan 2006

Methods	opaque sealed envelopes, envelopes opened immediately before administration of treatment or placebo during surgery random number sequence with masking of allocation by outside agency, no exclusions, 30% lost to follow up
Participants	Africans > 40
Interventions	trabeculectomy + beta radiation or dummy application
Outcomes	IOP < 21 without medication
Notes	Risk of bias assessment made on <i>Kirwan et al. BMJ</i> , doi:10.1136/bmj.38971.395301.7C (published 5 October 2006)

### *Risk of bias*

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Randomisation was in blocks of 20.." <i>First sentence, page 2.</i>
Allocation concealment (selection bias)	Low risk	"... with the assigned groups distributed to each centre in opaque, sealed envelopes. Each participant was allocated a trial number on recruitment. The envelope with that number was opened during surgery to determine allocation" <i>Page 2, first paragraph.</i>
Blinding (performance bias and detection bias) All outcomes	Low risk	"Each centre had two applicators, one active and one placebo. The identity of the active applicator was only known to IM and SC, neither of whom participated in data collection". <i>Pages 1 &amp; 2, last sentence on page 1.</i> "Separate record sheets were used at follow-up to mask clinicians from the treatment allocation" <i>Page 2, first paragraph.</i>
Incomplete outcome data (attrition bias) All outcomes	Low risk	"Twenty participants (6%) dropped out of the study after surgery. Some evidence was found that the probability of not being seen was associated with sex (women 10% v men 4%: P = 0.03), blindness in the fellow eye (9% v 3%: P = 0.02), and centre (Bloemfontein 2%, Edendale 9%, Pretoria 3%: P = 0.09; table 1). No strong evidence was found for an association with other variables. A further 68 patients were lost to follow-up before 12 months (36 in the radiation arm). Similar mean numbers of fol-



**Kirwan 2006** (Continued)

		low-up visits were completed in each of the treatment groups (4.9 radiation group; 4.5 placebo group; P = 0.35). Participants who were followed up showed similar distributions for socio-demographic, ophthalmic, and surgical factors in the treatment arms (table 1).” <i>Page 2 first paragraph</i> <i>From figure 1:</i> 117/156 (75%) in control group followed at 12 months; 115/164 (70%) of radiation group followed at 12 months
Selective reporting (reporting bias)	Low risk	All outcomes specified in the protocol were reported in the paper. Jim Kirwan
Other bias	Low risk	-

**Lai 1994**

Methods	sealed envelopes, randomised number sequence, ? exclusions ? FU
Participants	Hong Kong Chinese > 30
Interventions	trabeculectomy +/- beta radiation
Outcomes	IOP < 21 = success (+/- medication) mean IOP
Notes	published in abstract form only

***Risk of bias***

<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Random sequence generation (selection bias)	Unclear risk	“Patients with primary open angle glaucoma whose pressure was over 21 mmHg on maximum anti-glaucoma medications were assigned randomly to either Group 1) : Trabeculectomy, or Group 2) Trabeculectomy + Intraoperative beta irradiation.”
Allocation concealment (selection bias)	Low risk	Adequate allocation concealment reported by investigator
Blinding (performance bias and detection bias) All outcomes	Unclear risk	-

Lai 1994 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	-
Selective reporting (reporting bias)	Unclear risk	-
Other bias	Unclear risk	-

Rehman 2002

Methods	random number table, no exclusions, no loss to FU
Participants	Caucasian > 40
Interventions	trabeculectomy +/- beta radiation
Outcomes	IOP < 21 = success (+/- medication) mean IOP
Notes	recruitment terminated early due to changes in clinical practice

*Risk of bias*

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"A random numbers table was used to assign each eye to either control or beta irradiation group," Page 303, materials and methods, first paragraph, last sentence.
Allocation concealment (selection bias)	Low risk	"..with ophthalmologist and patient masked to the assignment." Page 303, materials and methods, first paragraph, last sentence. Given this statement allocation concealment likely.
Blinding (performance bias and detection bias) All outcomes	Low risk	"Patients in the control group received an identical applicator but without an active source (an inactive plaque also designed by Amersham)." Page 303, materials and methods, middle of 3 <sup>rd</sup> paragraph. There is not much information on masking but given this statement it would be unlikely that patients and outcome assessors would know which had received beta-radiation.

**Rehman 2002** (Continued)

Incomplete outcome data (attrition bias) All outcomes	Low risk	No exclusions and no attrition reported. 61 patients were enrolled into the trial and data on 61 patients reported at 12 months for all outcomes (100% follow-up)
Selective reporting (reporting bias)	Low risk	The main outcome reported in this study was intraocular pressure. This is as would be expected and in any case there were no statistically significant differences between the groups with respect to this outcome.
Other bias	Low risk	Early termination of patient recruitment was reported because patient recruitment was slow. However, this early termination was not determined by the results of the study

FU: follow up

IOP: intraocular pressure

**Characteristics of excluded studies** [ordered by study ID]

Study	Reason for exclusion
Tesha 2002	Further report of data in Rehman's study in abstract form

## DATA AND ANALYSES

### Comparison 1. Surgical failure

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 IOP > 21 mmHg 12 months or more after surgery	4		RR (Fixed, 95% CI)	0.23 [0.14, 0.40]
1.1 Black African people	1		RR (Fixed, 95% CI)	0.21 [0.11, 0.40]
1.2 Caucasian people	2		RR (Fixed, 95% CI)	0.15 [0.02, 1.28]
1.3 Chinese people	1		RR (Fixed, 95% CI)	0.34 [0.12, 0.96]

### Comparison 2. Intraocular pressure (IOP)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Mean IOP 12 months or more after surgery	2	123	Mean Difference (IV, Fixed, 95% CI)	-0.97 [-2.56, 0.62]

### Comparison 3. Adverse effects

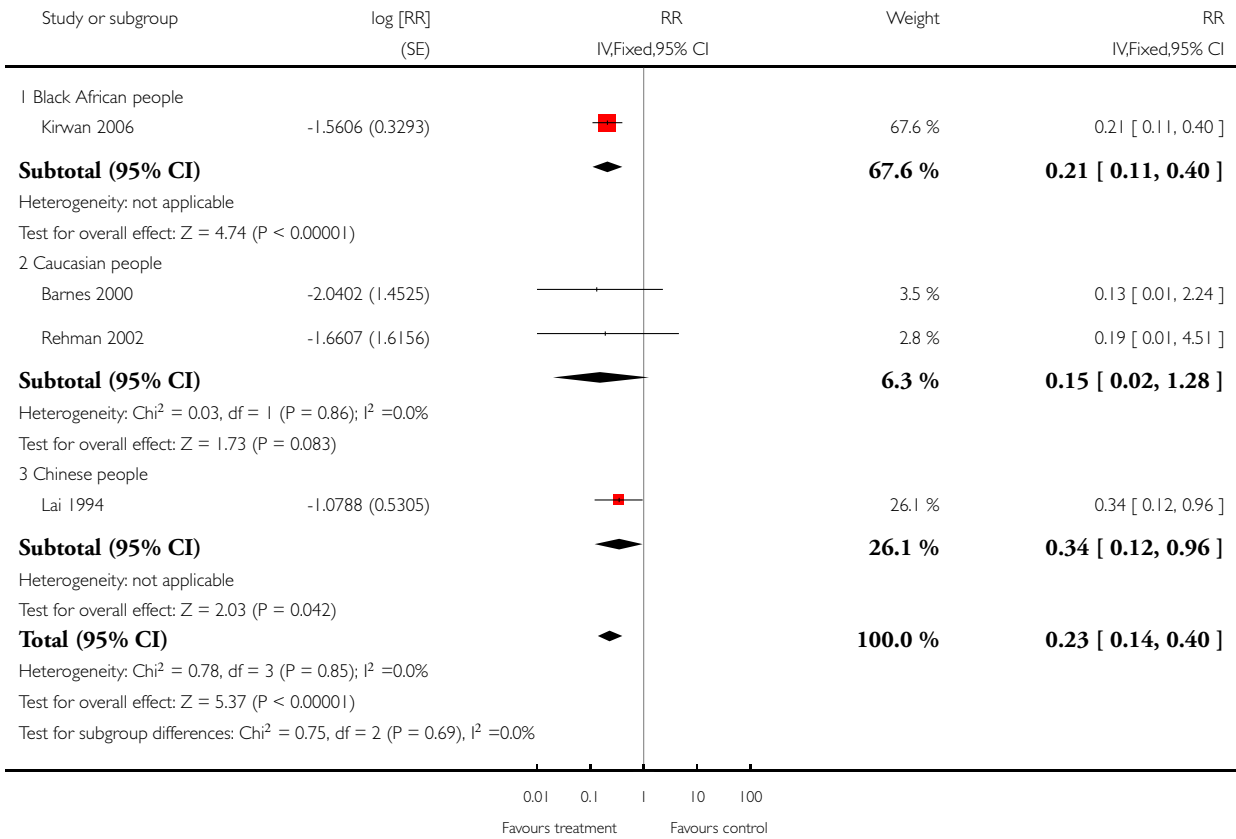
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Cataract	2	362	Risk Ratio (M-H, Fixed, 95% CI)	2.89 [1.39, 6.00]
2 Hypotony	3	423	Risk Ratio (M-H, Fixed, 95% CI)	1.79 [0.62, 5.14]
3 Bleb leak	2	123	Odds Ratio (M-H, Fixed, 95% CI)	0.53 [0.12, 2.38]

**Analysis 1.1. Comparison 1 Surgical failure, Outcome 1 IOP > 21 mmHg 12 months or more after surgery.**

Review: Beta radiation for glaucoma surgery

Comparison: 1 Surgical failure

Outcome: 1 IOP > 21 mmHg 12 months or more after surgery

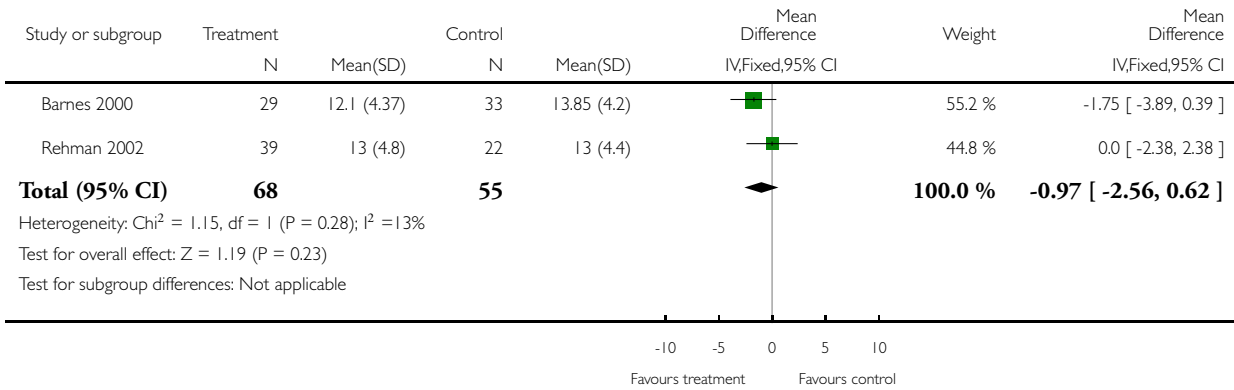


### Analysis 2.1. Comparison 2 Intraocular pressure (IOP), Outcome 1 Mean IOP 12 months or more after surgery.

Review: Beta radiation for glaucoma surgery

Comparison: 2 Intraocular pressure (IOP)

Outcome: 1 Mean IOP 12 months or more after surgery

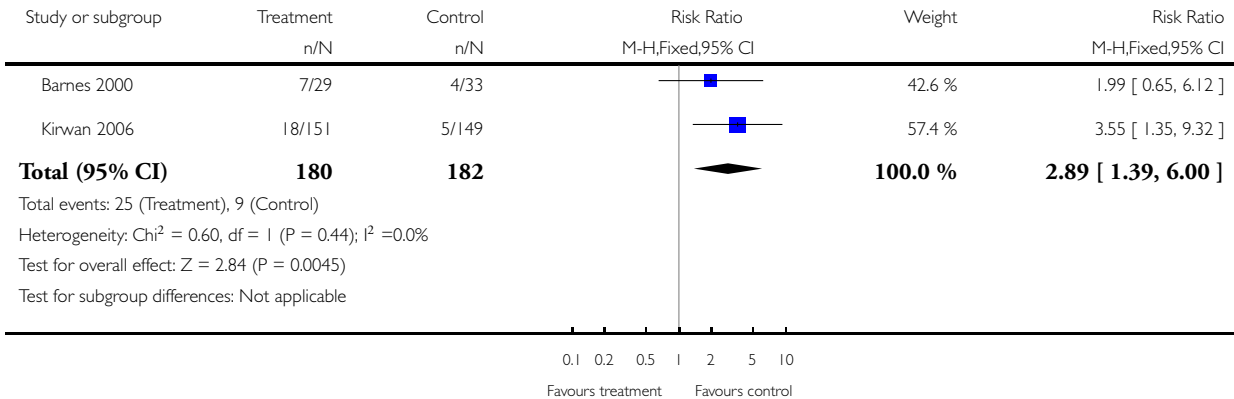


### Analysis 3.1. Comparison 3 Adverse effects, Outcome 1 Cataract.

Review: Beta radiation for glaucoma surgery

Comparison: 3 Adverse effects

Outcome: 1 Cataract

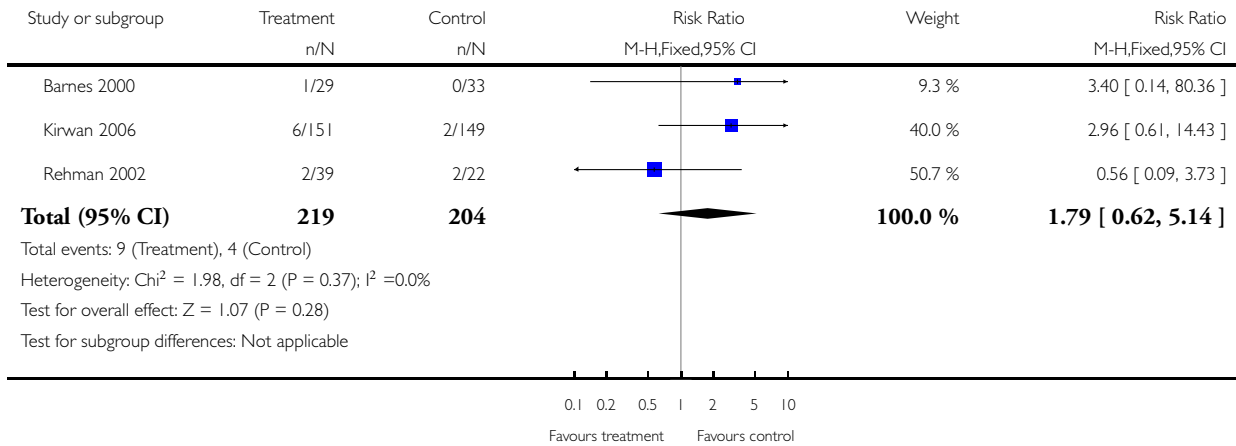


### Analysis 3.2. Comparison 3 Adverse effects, Outcome 2 Hypotony.

Review: Beta radiation for glaucoma surgery

Comparison: 3 Adverse effects

Outcome: 2 Hypotony

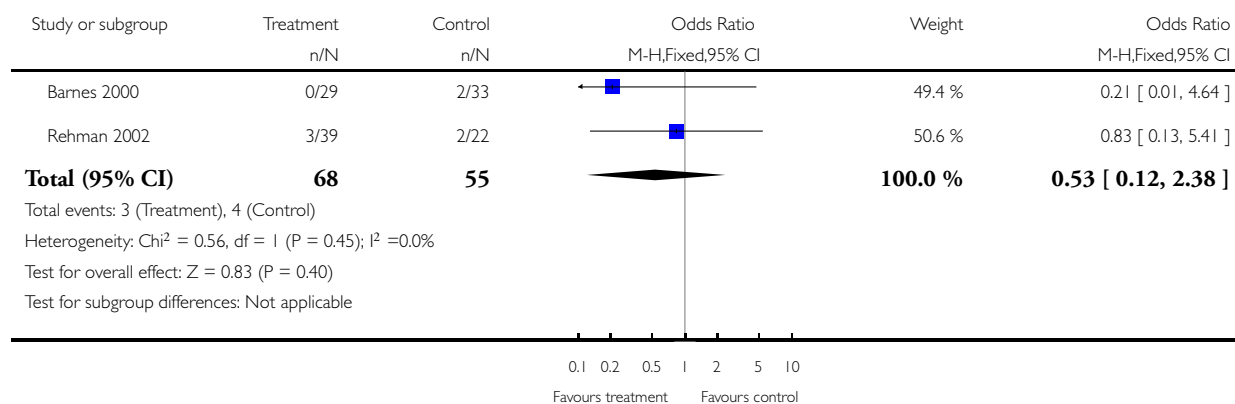


### Analysis 3.3. Comparison 3 Adverse effects, Outcome 3 Bleb leak.

Review: Beta radiation for glaucoma surgery

Comparison: 3 Adverse effects

Outcome: 3 Bleb leak



## ADDITIONAL TABLES

Table 1. Summary of trial characteristics

Study	Ethnic group	Dose	N treatment group	N control group	Surgical failure at 12 months in control group
Barnes 2000	Caucasian	750 cGy	31	34	12%
Kirwan 2006	Black African	1000 cGy	164	156	30%
Lai 1994	Chinese	1000 cGy	46	59	26%
Rehman 2002	Caucasian	750 cGy	39	22	14%



## APPENDICES

### Appendix 1. CENTRAL search strategy

- #1 MeSH descriptor Glaucoma
- #2 MeSH descriptor Filtering Surgery
- #3 (glaucoma\* or filter\* or filtrat\*) near (surg\*)
- #4 MeSH descriptor Trabeculectomy
- #5 trabeculectom\*
- #6 (#1 OR #2 OR #3 OR #4 OR #5)
- #7 MeSH descriptor Beta Rays
- #8 MeSH descriptor Radiation
- #9 MeSH descriptor Strontium
- #10 MeSH descriptor Brachytherapy
- #11 radiat\* or irradiat\*
- #12 strontium or SR 90
- #13 beta-rad\* or beta irradi\*
- #14 (#7 OR #8 OR #9 OR #10 OR #11 OR #12 OR #13)
- #15 (#6 AND #14)

### Appendix 2. MEDLINE (OvidSP) search strategy

1. randomized controlled trial.pt.
2. (randomized or randomised).ab,ti.
3. placebo.ab,ti.
4. dt.fs.
5. randomly.ab,ti.
6. trial.ab,ti.
7. groups.ab,ti.
8. or/1-7
9. exp animals/
10. exp humans/
11. 9 not (9 and 10)
12. 8 not 11
13. exp glaucoma/
14. exp filtering surgery/
15. ((glaucoma\$ or filter\$ or filtrat\$) adj3 surg\$).tw.
16. trabeculectom\$.tw.
17. or/13-16
18. beta particles/
19. exp radiation/
20. strontium/
21. exp brachytherapy/
22. (radiat\$ or irradiat\$).tw.
23. (strontium\$ or SR 90).tw.
24. (beta-radiat\$ or beta-irradiat\$).tw.
25. or/18-24
26. 17 and 25
27. 12 and 26

The search filter for trials at the beginning of the MEDLINE strategy is from the published paper by Glanville et al ([Glanville 2006](#)).

### Appendix 3. EMBASE (OvidSP) search strategy

1. exp randomized controlled trial/
2. exp randomization/
3. exp double blind procedure/
4. exp single blind procedure/
5. random\$.tw.
6. or/1-5
7. (animal or animal experiment).sh.
8. human.sh.
9. 7 and 8
10. 7 not 9
11. 6 not 10
12. exp clinical trial/
13. (clin\$ adj3 trial\$).tw.
14. ((singl\$ or doubl\$ or trebl\$ or tripl\$) adj3 (blind\$ or mask\$)).tw.
15. exp placebo/
16. placebo\$.tw.
17. random\$.tw.
18. exp experimental design/
19. exp crossover procedure/
20. exp control group/
21. exp latin square design/
22. or/12-21
23. 22 not 10
24. 23 not 11
25. exp comparative study/
26. exp evaluation/
27. exp prospective study/
28. (control\$ or prospectiv\$ or volunteer\$).tw.
29. or/25-28
30. 29 not 10
31. 30 not (11 or 23)
32. 11 or 24 or 31
33. exp glaucoma/
34. exp glaucoma surgery/
35. ((glaucoma\$ or filter\$ or filtrat\$) adj3 surg\$).tw.
36. trabeculectom\$.tw.
37. or/33-36
38. beta radiation/
39. exp irradiation/
40. strontium 90/
41. exp brachytherapy/
42. (radiat\$ or irradiat\$).tw.
43. (strontium\$ or SR 90).tw.
44. (beta-radiat\$ or beta-irradiat\$).tw.
45. or/38-44
46. 37 and 45
47. 32 and 46

#### **Appendix 4. metaRegister of Controlled Trials search strategy**

beta radiation and glaucoma

#### **Appendix 5. ClinicalTrials.gov search strategy**

Beta Radiation AND Glaucoma

#### **Appendix 6. ICTRP search strategy**

beta radiation AND glaucoma

### **WHAT'S NEW**

Last assessed as up-to-date: 26 March 2012.

<b>Date</b>	<b>Event</b>	<b>Description</b>
9 May 2012	New citation required but conclusions have not changed	Issue 6, 2012: New no trials were identified that met the inclusion criteria
9 May 2012	New search has been performed	Issue 6, 2012: Electronic searches were updated.

### **HISTORY**

Protocol first published: Issue 1, 2002

Review first published: Issue 2, 2009

<b>Date</b>	<b>Event</b>	<b>Description</b>
17 March 2008	Amended	Converted to new review format.

### **CONTRIBUTIONS OF AUTHORS**

JK wrote the protocol and assessed studies for inclusion, assessed studies for risk of bias, analysed the results and wrote the text of the review. CR assessed studies for inclusion and risk of bias and helped draft the review. JE assessed studies for risk of bias, extracted and entered data, did the summary of findings table and edited the text of the review.

## DECLARATIONS OF INTEREST

One of the review authors (JFK) has conducted an RCT into the effects of beta radiation on augmenting the success of glaucoma surgery in Africans in South Africa.

## SOURCES OF SUPPORT

### Internal sources

- Institute of Ophthalmology, UK.
- NIHR/Department of Health, UK.

JE was funded by NIHR during the updating of this review (Issue 6, 2012)

### External sources

- No sources of support supplied

## DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Analysis by ethnic group conducted in the review but not specified in the protocol.

## INDEX TERMS

### Medical Subject Headings (MeSH)

Beta Particles [adverse effects; \*therapeutic use]; Cataract [etiology]; Combined Modality Therapy [methods]; Glaucoma [\*surgery]; Randomized Controlled Trials as Topic; Trabeculectomy [\*methods]; Treatment Outcome; Wound Healing [\*radiation effects]

### MeSH check words

Humans