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Mitomycin C versus 5-Fluorouracil for wound healing in glaucoma surgery (Review)

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T A B L E   O F   C O N T E N T S

[Intervention Review]

Mitomycin C versus 5-Fluorouracil for wound healing in glaucoma surgery

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A B S T R A C T

Background

Raised intraocular pressure is a risk factor for glaucoma. One treatment option is glaucoma drainage surgery (trabeculectomy). Antimetabolites are used during surgery to reduce postoperative scarring during wound healing. Two agents in common use are mitomycin C (MMC) and 5-Fluorouracil (5-FU).

Objectives

To assess the effects of MMC compared to 5-FU as an antimetabolite adjunct in trabeculectomy surgery.

Search methods

We searched CENTRAL (which contains the Cochrane Eyes and Vision Group Trials Register) (2015 Issue 9), Ovid MEDLINE, Ovid MEDLINE In-Process and Other Non-Indexed Citations, Ovid MEDLINE Daily, Ovid OLDMEDLINE (January 1946 to October 2015), EMBASE (January 1980 to October 2015), Latin American and Caribbean Health Sciences Literature Database (LILACS) (January 1982 to October 2015), the ISRCTN registry (www.isrctn.com/editAdvancedSearch), ClinicalTrials.gov (www.clinicaltrials.gov) and the World Health Organization (WHO) International Clinical Trials Registry Platform (ICTRP) (www.who.int/ictrp/search/en). We did not use any date or language restrictions in the electronic searches for trials. We last searched the electronic databases on 2 October 2015.

Selection criteria

We included randomised controlled trials where wound healing had been modified with MMC compared to 5-FU.

Data collection and analysis

Two review authors independently selected trials and collected data. The primary outcome was failure of a functioning trabeculectomy one year after surgery. Secondary outcomes included mean intraocular pressure at one year. We considered three subgroups: high risk of trabeculectomy failure (people with previous glaucoma surgery, extracapsular cataract surgery, African origin and people with secondary glaucoma or congenital glaucoma); medium risk of trabeculectomy failure (people undergoing trabeculectomy with extracapsular cataract surgery) and low risk of trabeculectomy failure (people who have received no previous surgical eye intervention).

Main results

We identified 11 trials that enrolled 687 eyes of 679 participants. The studies were conducted in the United States, Europe, Asia and Africa. Five studies enrolled participants at low risk of trabeculectomy failure, five studies enrolled participants at high risk of failure, and one study enrolled people with both high and low risk of failure. None of the included trials enrolled participants with combined trabeculectomy/cataract surgery.

We considered one study to be at low risk of bias in all domains, six studies to be at high risk of bias in one or more domains, and the remaining four studies to be at an unclear risk of bias in all domains.

The risk of failure of trabeculectomy at one year after surgery was less in those participants who received MMC compared to those who received 5-FU, however the confidence intervals were wide and are compatible with no effect (risk ratio (RR) 0.54, 95% confidence interval (CI) 0.30 to 1.00; studies = 11; I^2 = 40%). There was no evidence for any difference between groups at high and low risk of failure (test for subgroup differences P = 0.69).

On average, people treated with MMC had lower intraocular pressure at one year (mean difference (MD) -3.05 mmHg, 95% CI -4.60 to -1.50), but the studies were inconsistent (I^2 = 52%). The size of the effect was greater in the high-risk group (MD -4.18 mmHg, 95% CI -6.73 to -1.64) compared to the low-risk group (MD -1.72 mmHg, 95% CI -3.28 to -0.16), but again the test for interaction was not statistically significant (P = 0.11).

Similar proportions of eyes treated with MMC lost 2 or more lines of visual acuity one year after surgery compared to 5-FU, but the confidence intervals were wide (RR 1.05, 95% CI 0.54 to 2.06).

Adverse events occurred relatively rarely, and estimates of effect were generally imprecise. There was some evidence for less epitheliopathy in the MMC group (RR 0.23, 95% CI 0.11 to 0.47) and less hyphaema in the MMC group (RR 0.62, 95% CI 0.42 to 0.91).

None of the studies reported quality of life.

Overall, we graded the quality of the evidence as low largely because of risk of bias in the included studies and imprecision in the estimate of effect.

Authors' conclusions

We found low-quality evidence that MMC may be more effective in achieving long-term lower intraocular pressure than 5-FU. Further comparative research on MMC and 5-FU is needed to enhance reliability and validity of the results shown in this review. Furthermore, the development of new agents that control postoperative scar tissue formation without side effects would be valuable and is justified by the results of this review.

P L A I N   L A N G U A G E   S U M M A R Y

Mitomycin C versus 5-Fluorouracil for wound healing in glaucoma surgery

Review question

Does mitomycin C (MMC) offer any advantage in comparison to 5-Fluorouracil (5-FU) as the antimetabolite used to augment glaucoma surgery (trabeculectomy)? Does MMC help to achieve lower rates of trabeculectomy failure than 5-FU at one year postoperatively?

Background

Raised intraocular pressure is a risk factor for glaucoma. One treatment option is glaucoma drainage surgery (trabeculectomy) to help lower intraocular pressure. Antimetabolites are medicines used during surgery to help reduce scarring after surgery during wound healing. If scarring occurs it can lead to treatment failure because the drainage channel no longer works. Two agents in common use are MMC and 5-FU.

Search date

The evidence is up to date to October 2015.

Study characteristics

We included 11 randomised controlled trials conducted in the United States, Europe, Asia and Africa in this review. In total, 687 eyes of 679 participants underwent routine trabeculectomy for glaucoma control. Some participants were at a higher risk of failure than others, for example if they had had previous glaucoma surgery, were of African origin, or if they had secondary glaucoma. Five studies enrolled participants at low risk of trabeculectomy failure, five studies enrolled participants at high risk of failure, and one study enrolled people with both high and low risk of failure. None of the included trials enrolled participants with combined trabeculectomy/cataract surgery.

Key results

Our review showed that the risk of failure of trabeculectomy at one year after surgery was slightly less in those participants treated with MMC compared to 5-FU. All of the included randomised controlled trials contributed to this result, with a mixed study population of high- and low-risk participants and varied methodology of antimetabolite application. We did not detect any significant differences between the subgroups of participants at low and high risk of failure, but the power of this analysis was low.

We identified no difference between the visual outcomes of the group that received MMC and the group that received 5-FU at one year postoperatively nor in the number of drops used postoperatively. However, we found evidence to suggest that MMC was more effective at lowering intraocular pressure than 5-FU in both high- and low-risk participants, achieving a lower mean intraocular pressure postoperatively than in those who were treated with 5-FU at one year. This effect seemed to be greater in the high-risk populations.

Evaluating the overall complications across all studies revealed a slight favour toward using MMC, particularly with the incidence of epitheliopathy and hyphaema. There was a trend towards bleb leaks, wound leaks, late hypotony and cataract formation in the MMC-treated group.

None of the studies reported quality of life.

Quality of the evidence

We graded the quality of the evidence as low, mostly due to the risk of bias in the included studies. One bias we commonly encountered came from the different techniques of antimetabolite administration, making it difficult to conceal which medicine was being used. Furthermore, most studies only had a few complications to report, which meant that there were low numbers overall to include in the analysis of complications.

SUMMARY OF FINDINGS FOR THE MAIN COMPARISON [Explanation]

MMC compared to 5-FU for wound healing in glaucoma surgery

Patient or population: wound healing in glaucoma surgery
Settings:
Intervention: MMC
Comparison: 5-FU

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants/eyes (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	**Corresponding risk**				
	5-FU	**MMC**				
Failure of functioning trabeculectomy at 1 year	**Study population**		Low-risk population RR 0.65 (95% CI 0.19 to 2. 20) High-risk population RR 0.49 (95% CI 0.22 to 1. 08)	634 (11 RCTs: 6 including low-risk population and 5 including high-risk population)	⊕⊕◯◯ LOW [1,2]	
	Low-risk population: 74 per 1000 High-risk population: 272 per 1000	Low-risk population: 50 per 1000 High-risk population: 137 per 1000				
Intraocular pressure at 1 year	The mean intraocular pressure at 1 year ranged across 5-FU groups Low-risk population: 10. 9 to 14.3 mmHg High-risk population: 14. 8 to 16.3 mmHg	The mean intraocular pressure at 1 year in the MMC groups had a range of values Low-risk population: 9.9 to 11.6 mmHg High-risk population: 8.6 to 13.7 mmHg	-	386 (7 RCTs: 3 including low-risk population and 4 including high-risk population)	⊕⊕◯◯ LOW [1,3]	
Loss of 2 or more lines of Snellen visual acuity at 1 year	**Study population**		Low-risk population RR 2.00 (95% CI 0.53 to 7. 59) High-risk population RR 0.81 (95% CI 0.36 to 1. 80)	328 (5 RCTs: 2 including low-risk population and 3 including high-risk population)	⊕⊕◯◯ LOW [2,4]	

	Low-risk population: 47 per 1000 High-risk population: 115 per 1000	Low-risk population: 94 per 1000 High-risk population: 96 per 1000		
Postoperative complications: late hypotony	**Study population** 37 per 1000	 59 per 1000	RR 1.37 (95% CI 0.41 to 4.63)	211 (4 RCTs)	⊕⊕◯◯ LOW [2,4]
Postoperative complications: choroidal detachment	**Study population** 68 per 1000	 70 per 1000	RR 0.86 (95% CI 0.45 to 1.63)	494 (8 RCTs)	⊕⊕◯◯ LOW [1,2]
Postoperative complications: endophthalmitis	**Study population** 0 per 1000	 19 per 1000	RR 3.89 (95% CI 0.44 to 34.57)	315 (4 RCTs)	⊕⊕◯◯ LOW [1,2]
Quality of life at 1 year				Not reported

*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).

5-FU: 5-Fluorouracil; **CI:** confidence interval; **MMC:** mitomycin C; **RCT:** randomised controlled trial; **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.

[1] Downgraded for risk of bias: only one study at low risk of bias in all domains
[2] Downgraded for imprecision: wide confidence intervals
[3] Downgraded for inconsistency: $I^2 = 60\%$
[4] Downgraded for risk of bias: no study at low risk of bias in all domains

BACKGROUND

Description of the condition

Glaucoma is a chronic, progressive optic neuropathy characterised by a progressive loss of ganglion cells that leads to a characteristic visual function loss. Intraocular pressure (IOP) is often considered to be a major risk factor for glaucoma, and it is the only factor that can be modified to try to change the course of the condition. The publication of a series of randomised controlled trials (RCTs) has established the evidence for treating glaucoma with IOP reduction (AGIS 1998; CNTGS 1998; Heijl 2002; Kass 2002; Maier 2005; Vass 2007).

Glaucoma drainage surgery remains an important treatment option for the control of IOP despite the addition of several new IOP-lowering drugs. Some evidence suggests that trabeculectomy is more effective than either medicine or laser treatment alternatives (Migdal 1994). However, a Cochrane systematic review from 2012 found that visual field deterioration up to five years is not significantly different whether treatment is initiated with medication or trabeculectomy (Burr 2012).

Optimum success rates are achieved when the eye has been exposed to no previous interventions, either surgical or medical, although this is not the usual situation in high-income countries. Risk factors for trabeculectomy failure are thought to be those that increase the scarring response and include previous exposure to topical medication, previous surgical manipulation of the conjunctiva or other injury, young age, African origin, a history of uveitis and neovascular glaucoma (EGS 2003).

Presentation and diagnosis

The diagnosis of glaucoma is made by the identification of a progressive optic neuropathy or a characteristic visual field defect. There are subgroups of glaucoma, primary open angle glaucoma being most common in European and African populations. A person with primary open angle glaucoma is often unaware of any symptoms until the late stages of the disease, making early diagnosis essential.

Description of the intervention

Treatment is usually initiated with topical treatment, and surgical options are considered if topical treatment fails to prevent progression of the disease. The trabeculectomy produces a guarded fistula between the anterior chamber and the subconjunctival space. There have been numerous modifications since its first description (Cairns 1968), including the use of antimetabolites to reduce fibroblast activity and postoperative scarring at the site of the scleral flap and the subconjunctival space.

How the intervention might work

Once trabeculectomy has been selected, the treatment decisions are whether to augment the surgery with antiscarring agents such as antimetabolites. Antimetabolites are applied to the surgical site to inhibit fibroblast activity and reduce postoperative scarring; the two agents commonly in use are mitomycin C (MMC) and 5-Fluorouracil (5-FU). Due to reported side effects such as increased risk of bleb leak, hypotony and endophthalmitis (DeBry 2002), there is concern that use of these agents should be restricted to high-risk cases only. A number of RCTs have reported the use of MMC (Andreanos 1997; Carlson 1997; Cohen 1996; Costa 1996; Martini 1997; Robin 1997; Shin 1995; Shin 1998; Wu 1996). A Cochrane systematic review concluded that compared to placebo, MMC reduces mean IOP at 12 months in all groups of participants (Wilkins 2010). Apart from increase in cataract formation, there was insufficient power to detect any increase in other serious side effects. Postoperative 5-FU injections to augment trabeculectomy have also been assessed with RCTs (FFSSG 1989; Goldenfeld 1994; Ophir 1992; Ruderman 1987), and also confer an improvement in IOP control at one year compared to placebo (Green 2014). Clinically, MMC and 5-FU can be applied intraoperatively on a sponge placed for one to five minutes between the conjunctiva and sclera at the start of the operation. Alternatively, 5-FU may be given as one or more postoperative subconjunctival injections. There is marked variation in the concentrations of both drugs used, the time of intraoperative application and the position and volume of postoperative injections.

Why it is important to do this review

The results of two Cochrane reviews comparing MMC, in Wilkins 2010, and 5-FU, in Green 2014, to placebo suggest a similar effect for the two agents in inhibiting scarring after trabeculectomy. However, there is no direct comparative evidence to influence which antimetabolite a surgeon should choose. The purpose of this review was to systematically summarise the RCTs in which MMC was compared to 5-FU in an attempt to clearly identify treatment benefits of one agent over the other.

OBJECTIVES

To assess the effects of MMC compared to 5-FU as an antimetabolite adjunct in trabeculectomy surgery.

METHODS

Criteria for considering studies for this review

Types of studies

We included RCTs where wound healing had been modified with one of the antimetabolites in one group of people undergoing trabeculectomy, compared to the other antimetabolite in the other group.

Types of participants

There were three separate subgroup populations:

- High risk of trabeculectomy failure: people with previous glaucoma or extracapsular cataract surgery, people of African origin and people with secondary glaucoma or congenital glaucoma.
- Medium risk of trabeculectomy failure: (combined surgery) people undergoing trabeculectomy with extracapsular cataract surgery.
- Low risk of trabeculectomy failure: (primary trabeculectomy): people who have received no previous surgical eye intervention. People who underwent previous laser procedures may be included in this group.

For the purpose of this review, there were no restrictions regarding age or gender.

Types of interventions

We included the following interventions:

1. Use of intraoperative MMC versus intraoperative 5-FU.
2. Use of intraoperative MMC versus postoperative 5-FU.
3. Use of intraoperative MMC versus intraoperative and postoperative 5-FU.
4. Use of intraoperative MMC and postoperative MMC versus intraoperative and postoperative 5-FU.

Types of outcome measures

Primary outcomes

The primary outcome was failure of a functioning trabeculectomy at one year from surgery (dichotomous).
We used the following definitions:

- Success: adequate pressure control (< 22 mmHg) without additional treatment.
- Failure: need for repeat filtration surgery or uncontrolled IOP (= or > 22 mmHg).

Secondary outcomes

- Survival analysis (time to event) for the previously given definition of failure
- Mean IOP for each group at one year from surgery
- Quality-of-life measures
- Economic data

Adverse outcomes

Adverse events in either group with reference to choroidal detachment, hypotony and late endophthalmitis were reported. Adverse events were reported at any time during the follow-up period. We used the following definitions:

- Bleb leakage: presence of a positive Seidel test (visible aqueous flow with the tear film stained with fluorescein).
- Hypotony: IOP below 5 mmHg and/or associated with complications such as macular oedema and sight loss or choroidal detachments.
- Endophthalmitis: an infection of the globe contents that even with prompt aggressive treatment results in substantial loss of visual function.

Search methods for identification of studies

Electronic searches

We searched CENTRAL (which contains the Cochrane Eyes and Vision Group Trials Register) (2015 Issue 9), Ovid MEDLINE, Ovid MEDLINE In-Process and Other Non-Indexed Citations, Ovid MEDLINE Daily, Ovid OLDMEDLINE (January 1946 to October 2015), EMBASE (January 1980 to October 2015), Latin American and Caribbean Health Sciences Literature Database (LILACS) (January 1982 to October 2015), the ISRCTN registry (www.isrctn.com/editAdvancedSearch), ClinicalTrials.gov (www.clinicaltrials.gov) and the World Health Organization (WHO) International Clinical Trials Registry Platform (ICTRP) (www.who.int/ictrp/search/en). We did not use any date or language restrictions in the electronic searches for trials. We last searched the electronic databases on 2 October 2015.
See: Appendices for details of search strategies for CENTRAL (Appendix 1), MEDLINE (Appendix 2), EMBASE (Appendix 3), LILACS (Appendix 4), ISRCTN (Appendix 5), ClinicalTrials.gov (Appendix 6) and the ICTRP (Appendix 7).

Searching other resources

We searched the reference lists of identified trial reports to find additional trials. We contacted investigators as necessary to identify additional published and unpublished studies.

Data collection and analysis

Selection of studies

Three review authors (JC/EC/JE) independently reviewed the titles and abstracts resulting from the searches. We obtained full copies of any report referring to possibly or definitely relevant trials and assessed them according to the definitions in the Criteria for

considering studies for this review section. We assessed only trials meeting these predefined criteria for methodological quality. We resolved any disagreements by discussion.

Data extraction and management

Three review authors (JC/EC/JE) independently extracted data with relation to the outcome measures outlined above. We resolved discrepancies by discussion. One review author entered the data into Review Manager (RevMan 2014), and the other review authors checked the data entry.

Assessment of risk of bias in included studies

Three review authors (JC/EC/JE) independently assessed risk of bias according to methods set out in Chapter 8 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We considered six domains: random sequence generation, allocation concealment, masking, incomplete outcome data, selective reporting and any other identified bias. We graded each domain as low risk of bias, high risk of bias, or unclear risk of bias. For example, in allocation concealment the grading was low risk if there was central randomisation of subjects, high risk if there was simple alternating methods used to allocate subjects and unclear if there was no real qualifying statement. We resolved disagreements by discussion. Review authors were not masked to trial details during the assessment. We excluded trials scoring 'high risk' on allocation concealment. In cases where missing or confusing data did not permit a clear grading of the trial, we contacted the study authors in order to obtain further information.

Measures of treatment effect

We measured the effect of dichotomous data by risk ratio; continuous data by difference in means; and time to event data by hazard ratio.

Unit of analysis issues

All studies were parallel-group RCTs. In the majority of studies, one eye per person was enrolled, and therefore there were no unit of analysis issues. In Lamping 1995, WuDunn 2002 and Xinyu 2001, both eyes of some participants were enrolled, but in most cases this was less than 10%, and overall less than 5% of the data would be affected by this. None of the trials took into account the potential correlation between eyes, and we have analysed the data from the trials as reported.

Dealing with missing data

We did an available case analysis. This assumes that data are missing at random. We assessed whether this assumption was reasonable by collecting data from each included trial on the number of participants excluded or lost to follow-up and reasons for loss to follow-up by treatment group, if reported. We collected this information as part of the assessment of attrition bias.

Assessment of heterogeneity

We examined the overall characteristics of the studies, in particular the types of participants and interventions, in order to assess the extent to which the studies were similar enough to make pooling study results sensible.

We looked at the forest plot of study results to see how consistent the studies were, in particular looking at the size and direction of effects.

We calculated I^2, which is the percentage of the variability in effect estimates that is due to heterogeneity rather than sampling error (chance) (Higgins 2002). We considered I^2 values over 50% to indicate substantial inconsistency or heterogeneity. We also considered Chi^2 P values; when the number of studies was few we used P less than 0.1 to indicate statistical significance of the Chi^2 test.

Assessment of reporting biases

We planned to do a 'funnel plot' to investigate reporting (publication) bias, but there were not enough included trials (fewer than 10 in each meta-analysis) to make this possible.

Data synthesis

If there was inconsistency between individual study results such that a pooled result may not have been a good summary of the individual trial results, for example the effects were in different directions, or I^2 was greater than 50% and P less than 0.1, we did not pool the data but did describe the pattern of the individual study results.

If I^2 was greater than 50%, but all the effect estimates were in the same direction such that a pooled estimate would seem to have provided a good summary of the individual trial results, we did pool the data.

If there was inconsistency between individual study results such that a pooled result may not have been a good summary of the individual trial results, for example the effects were in different directions, or I^2 was greater than 50% and P less than 0.1, we did not pool the data but did describe the pattern of the individual study results.

If I^2 was greater than 50%, but all the effect estimates were in the same direction such that a pooled estimate would have provided a good summary of the individual trial results, we did pool the data.

Subgroup analysis and investigation of heterogeneity

We compared the effect of intervention in a pre-planned analysis comparing effects in groups at high and low risk of failure.

Sensitivity analysis

We conducted sensitivity analyses to determine the impact of risk of bias on effect size. We repeated the analyses excluding trials at high risk of bias in one or more domains.

R E S U L T S

Description of studies

Results of the search

The electronic searches yielded a total of 446 references (Figure 1). The Trials Search Co-ordinator scanned the search results, removed 58 duplicates and then removed 298 references that were not relevant to the scope of the review. We screened the remaining 90 reports and discarded 69 reports as not relevant. After assessing the reports, we identified a further two studies studies for potential inclusion in the review (Oh 1994; Uva 1996). In total, we obtained 23 full-text reports for potential inclusion in the review. After consideration of each report, we included a total of 13 reports of 11 studies in the final review; see Characteristics of included studies and excluded seven studies; see Characteristics of excluded studies for reasons. We have categorised three studies as awaiting assessment, two of which we are unable to source copies of the reports and one is awaiting a response from the authors regarding information on methods of randomisation (Liu 2015).

Figure 1. Study flow diagram


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┌─────────────────────────────┐
│ 446 records identified through │
│ electronic database searching │
└─────────────────────────────┘
              │
              ▼
┌─────────────────────────────┐
│ 388 records after duplicates │
│ removed                      │
└─────────────────────────────┘
              │
              ▼
┌─────────────────────────────┐        ┌─────────────────────────────┐
│ 388 records screened by the │───────▶│ 298 records excluded by the │
│ Trials Search Co-ordinator  │        │ TSC after initial screening │
│ (TSC)                       │        └─────────────────────────────┘
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┌─────────────────────────────┐        ┌─────────────────────────────┐
│ 90 records screened by the  │───────▶│ 69 records excluded by      │
│ review authors              │        │ review authors as not       │
│                             │        │ relevant                    │
└─────────────────────────────┘        └─────────────────────────────┘
              │
              ▼
┌─────────────────────────────┐
│ Review authors identified 2 │
│ additional full-text reports for │
│ potential inclusion         │
└─────────────────────────────┘
              │
              ▼
┌─────────────────────────────┐        ┌─────────────────────────────┐
│ 23 full-text articles assessed │────▶│ 7 full-text articles excluded, │
│ for eligibility             │        │ with reasons                 │
└─────────────────────────────┘        └─────────────────────────────┘
              │
              ▼
┌─────────────────────────────┐
│ 3 studies awaiting assessment, │
│ unable to source 2 reports and │
│ one report awaiting response │
│ from authors regarding       │
│ request for information on   │
│ methods of randomisation.    │
└─────────────────────────────┘
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              ▼
┌─────────────────────────────┐
│ 13 reports of 11 studies    │
│ included in qualitative     │
│ synthesis                   │
└─────────────────────────────┘
              │
              ▼
┌─────────────────────────────┐
│ 13 reports of 11 studies    │
│ included in quantitative    │
│ synthesis (meta-analysis)   │
└─────────────────────────────┘
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Included studies

Design

We included a total of 11 studies in this Cochrane Review and summarised them in the Characteristics of included studies. All 11 studies were designed as a prospective RCTs. One study in this review was a multicentre study (Singh 2000); the rest were single-centre.

Setting

Four studies were based in the United States (Katz 1995; Lamping 1995; Singh 2000; WuDunn 2002), two in Italy (Sisto 2007; Uva 1996) and the remainder in Ghana (Singh 1997), Japan (Kitazawa 1991), China (Xinyu 2001), Israel (Zadok 1995) and Iran (Mostafaei 2011). All research was carried out in clinical ophthalmic institutes.

Participants and sample sizes

In total, 687 eyes of 679 participants underwent routine trabeculectomy for glaucoma control. The smallest study was of 20 eyes of 20 participants (Zadok 1995), and the largest study included 115 eyes of 103 people (WuDunn 2002). Five studies included high-risk cases only (Katz 1995; Kitazawa 1991; Lamping 1995; Singh 1997; Sisto 2007), one study enrolled both high- and low-risk cases (Xinyu 2001), and the participants in the remaining five studies were low risk. Participants across the studies were a mixture of male and female; the percentage female ranged from 19% to 67%. The average age in the studies ranged from 47 years to 71 years, with a median average age of 62 years. One study had a significant age difference (P = 0.01), with a mean age of 41.2 in the MMC group and 54.2 in the 5-FU group (Kitazawa 1991).

Interventions

We have summarised the interventions in Table 1.

The majority of trials applied MMC using an intraoperative sponge; the exception was Mostafaei 2011, where 0.02mg MMC was applied by intraoperative subconjunctival injection. Subconjunctival application of MMC is not consistent with current practice (Dhingra 2009). The MMC dose given by intraoperative sponge varied between studies:

- Two studies used 0.5 mg/ml applied for 5 minutes, in Katz 1995, or 3.5 minutes, in Singh 1997.
- Three studies used 0.4 mg/ml applied for 5 minutes (Kitazawa 1991), 2.5 minutes (Lamping 1995), or 2 minutes (Singh 2000).

- Five studies used 0.2 mg/ml applied for 5 minutes, in Xinyu 2001 and Zadok 1995, or 2 minutes, in Sisto 2007, Uva 1996 and WuDunn 2002.

The method of administration of the 5-FU varied between studies: four studies used an intraoperative sponge technique similar to that of MMC application (Singh 1997; Singh 2000; Uva 1996; WuDunn 2002), six trials used a series of postoperative subconjunctival injections (Katz 1995; Kitazawa 1991; Lamping 1995; Sisto 2007; Xinyu 2001; Zadok 1995), and one study used intraoperative subconjunctival 5-FU 5 mg (Mostafaei 2011). All four studies with a group receiving intraoperative sponge-applied 5-FU used 50 mg/ml for 5 minutes, which is consistent with current practice (Dhingra 2009).

Different dosing regimens were used for the postoperative injections.

- Four studies used 10 postoperative injections
  - daily for 1 week, 3 times the following week (Katz 1995);
  - each day for 1 week, every other day for the following week (Kitazawa 1991);
  - first 10 days (Lamping 1995);
  - starting on day 7, 2 injections per week for 2 weeks and then 1 injection per week for 6 weeks (Sisto 2007).
- Two studies used approximately 7 postoperative injections
  - once daily up to 7 times in the first week after surgery (Zadok 1995);
  - 6 to 8 (alternate days, starting on day 3) (Xinyu 2001)

Outcomes

All of the 11 included studies stated an optimal postoperative IOP to achieve in order to accept success: five studies used a level of below 21 mmHg as desirable (Kitazawa 1991; Singh 1997; Singh 2000; Sisto 2007; Zadok 1995), two studies used equal to or less than 21 mmHg (Lamping 1995; WuDunn 2002), two used equal to or less than 12 mmHg (Katz 1995; Uva 1996), one used less than 21.06 mmHg (Xinyu 2001), and one study used 6 to 22 mmHg (Mostafaei 2011). Each study group reported their findings either as a percentage success or mean IOP.

Excluded studies

We excluded seven studies from the review: Ashworth 2003; Dreyer 1995; Li 2001; Membrey 2000; Membrey 2001; Rodriguez-Bermejo 1993; Oh 1994. For further details please see Characteristics of excluded studies.

Risk of bias in included studies
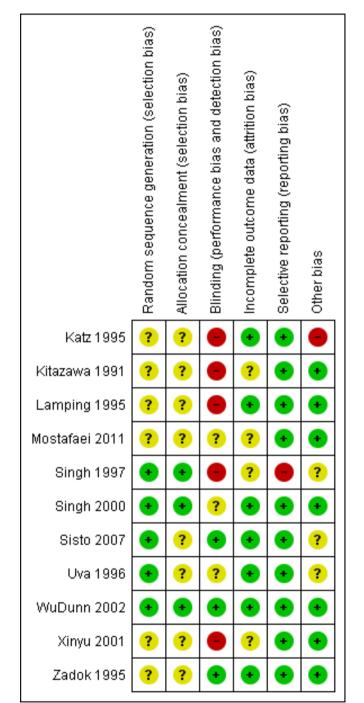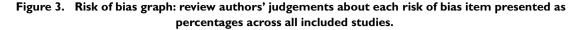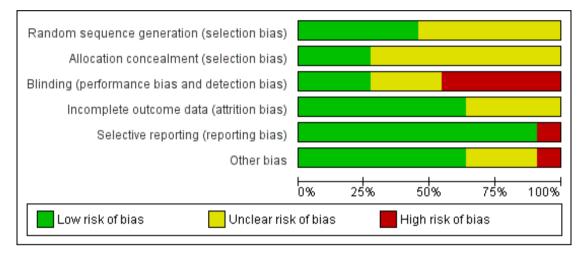
See Figure 2; Figure 3

Figure 2.   Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

Figure 3.   Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.

The review authors individually assessed the risk of bias. If the relative point was addressed in a study's manuscript, then a true assessment of 'high' or 'low' risk was carried out. If we deemed the risk as unclear, then this indicated we could make no true assessment because the required information was not given either in the published manuscript or after making contact with the lead author.

Allocation

Five studies reported adequate methods to generate a random allocation sequence: Singh 1997 tossed a coin in the operating theatre to allocate participants; Uva 1996 used a table of random numbers; and the remaining three studies used computer-generated allocation sequences (Singh 2000; Sisto 2007; WuDunn 2002).

It was unclear how the allocation schedule was generated in the remaining six studies.

Five studies reported adequate methods of allocation concealment (Kitazawa 1991; Singh 1997; Singh 2000; Uva 1996; WuDunn 2002)

Blinding

In four of the included studies, 5-FU was administered using a different technique to that of MMC, and no report was given about whether or not the follow-up information was gathered from masked assessors. We classified all these studies as high risk of performance and detection bias (Katz 1995; Kitazawa 1991; Lamping 1995; Xinyu 2001).

In one study, the method of 5-FU administration was the same as for MMC, but information gathered in the follow-up period was not from masked assessors. We therefore classified this study as high risk (Singh 1997).

Two studies used different techniques for antimetabolite administration but assessors were masked during the follow-up period. We graded these two studies as low risk of performance and detection bias for the primary outcome of this review (Sisto 2007; Zadok 1995).

Only one study used a placebo to mask allocation, which we graded as at low risk of perfomance and detection bias (WuDunn 2002). We graded the other three studies as unclear because the surgical administration of the antimetabolites was the same, but there was no mention of masking during follow-up (Mostafaei 2011; Singh 2000; Uva 1996).

Incomplete outcome data

Four studies did not comment on the exclusion or inclusion of participants in their analysis (Kitazawa 1991; Mostafaei 2011; Singh 1997; Xinyu 2001); we classified these as unclear risk. We classified the other seven studies as low risk as participants were clearly identified as included or not. No studies raised any concern over their intention to include or exclude participants.

Selective reporting

Singh 1997 did not specify in the methods of the paper what outcomes they considered, thus we cannot be certain that all the intended outcomes were addressed; we highlighted this as high risk. All other studies commented on all stated outcomes.

Other potential sources of bias

The only other sources of bias identified were that of postoperative care with regard to what other care or medications participants received and the varied amount of 5-FU a participant would receive with an incomplete postoperative regimen. This was highlighted in the study by Katz 1995. Other studies had no other clear identifiable bias.

Effects of interventions

See: **Summary of findings for the main comparison** MMC compared to 5-FU for wound healing in glaucoma surgery

Failure of a functioning trabeculectomy at one year from surgery (primary outcome)

All 11 studies reported failure of a functioning trabeculectomy at approximately one year, which was defined as IOP above (approximately) 22 mmHg or more (Analysis 1.1).

The risk of failure of trabeculectomy at one year after surgery was lower in those treated with MMC compared to 5-FU (risk ratio (RR) 0.54, 95% confidence interval (CI) 0.30 to 1.00; studies = 11; I^2 = 40%). However, the confidence intervals of the studies were wide, and we cannot exclude important differences.

There was no evidence for any difference between groups at high and low risk of failure (test for subgroup differences P = 0.69), but with only a few trials in each group, the power of the analysis to detect any differences was low.

The dose of MMC varied across the studies included in the review, and consequently we performed a dose-response analysis. We identified a trend showing that studies increasingly favoured the use of MMC rather than 5-FU as the intraoperative exposure to MMC increased (Analysis 1.2). Overall exposure was calculated by multiplying the concentration of MMC by the duration of exposure for each study. We then listed the studies in descending order of MMC exposure to view the overall effect. We excluded one study that administered the MMC by subconjunctival injection from this analysis.

When considering the method of 5-FU administration as in Analysis 1.3, there was no significant effect on the overall outcome whether the 5-FU was administered by postoperative subconjunctival injections or by the more current method of intraoperative sponge application (subgroup difference P = 0.93)

Time to failure of functioning trabeculectomy

No trial reported this outcome.

Mean IOP one year from surgery

Seven studies reported mean IOP at 12 months (range 6 to 18 months). On average, people treated with MMC had lower IOP at one year (mean difference (MD) -3.05 mmHg, 95% CI -4.60

to -1.50) Analysis 1.4. There was inconsistency between trials (I^2 = 52%), the MD showing a large range in the studies.

The size of the effect was greater in the high-risk group (MD -4.18 mmHg, 95% CI -6.73 to -1.64) compared to the low-risk group (MD -1.72 mmHg, 95% CI -3.28 to -0.16), but the test for interaction was not statistically significant (P = 0.11).

Postoperative use of antiglaucoma medications

Seven studies reported on the frequency of postoperative use of antiglaucoma medications. Similar proportions of people treated with MMC and 5-FU required postoperative medication to control pressure (RR 1.03, 95% CI 0.57 to 1.85) (Analysis 1.5). There was no evidence for any difference in effect between high-risk and low-risk groups (P = 0.88). The low-risk group trials were consistent (I^2 = 0%), but we saw different results in the three higher-risk group trials (I^2 = 74%).

Four studies reported the mean number of antiglaucoma medications used. On average, people receiving MMC used fewer antiglaucoma medications (MD -0.33, 95% CI -0.70 to 0.05), but the effect was uncertain (CIs include 0.00), and the studies were inconsistent (I^2 = 71%) (Analysis 1.6). The inconsistency in the trials came from those with a higher risk of failure (I^2 = 62%). However, there was a difference between the trials including participants at high risk of failure and those including participants at low risk of failure with a greater relative effect of MMC in the higher-risk groups (test for interaction P = 0.06). The main caveat was that there were only two trials in each group of the analysis.

Reduction in visual acuity

Five studies reported postoperative visual acuity. The proportion of eyes treated with MMC that lost 2 or more lines of visual acuity one year after surgery was similar to that of 5-FU, but the CIs were wide (RR 1.05, 95% CI 0.54 to 2.06) Analysis 1.7.

Quality of life

No trial reported this outcome.

Economic data

No trial reported this outcome.

Adverse outcomes

Bleb leak

Two studies reported bleb leak as a complication encountered following trabeculectomy. Participants receiving MMC were more likely to have a postoperative bleb leak, although the CI was wide,

and only two studies reported this outcome (RR 1.22, 95% CI 0.32 to 4.68; I^2 = 0%).

Six studies used the term 'wound leak' rather than 'bleb leak' in their assessment of postoperative complications. These studies also showed, with similar statistics, that participants receiving MMC were more likely to have a postoperative wound leak than those receiving 5-FU, although the CI was wide (RR 1.17, 95% CI 0.51 to 2.71; I^2 = 0%).

Late hypotony

Five studies reported hypotony post-trabeculectomy. Participants receiving MMC were more likely to have postoperative hypotony compared to those participants who received 5-FU, however the effect was uncertain with wide CIs compatible with no effect or increased hypotony in the 5-FU group (RR 1.37, 95% CI 0.41 to 4.63; I^2 = 0%).

Maculopathy

Four studies reported maculopathy following trabeculectomy. Participants receiving MMC were more likely to encounter maculopathy postoperatively than those receiving 5-FU, but the effect was uncertain and CI compatible with no effect or increased maculopathy in the 5-FU group (RR 1.71, 95% CI 0.35 to 8.33; I^2 = 0%).

Cataract

Four studies reported the incidence of postoperative cataract development. Participants receiving MMC were more likely to develop cataract than those receiving 5-FU, but again the CIs include 1 (null effect) (RR 1.73, 95% CI 0.65 to 4.61; I^2 = 24%).

Shallow anterior chamber

Five studies noted postoperative shallowing of the anterior chamber. Those participants receiving MMC were more likely to present with a shallow anterior chamber than those who received 5-FU. The statistical analysis showed a wide CI (RR 1.22, 95% CI 0.67 to 2.21; I^2 = 0%).

Choroidal detachment

Nine studies (549 eyes) reported a choroidal detachment as a postoperative complication following trabeculectomy. Three studies (303 eyes) reported the same event as a 'suprachoroidal haemorrhage'. The former group of studies found no difference in the

rate of events between those participants who received MMC and those who received 5-FU (RR 0.86, 95% CI 0.45 to 1.63; I^2 = 0%). The latter group of studies favoured those participants who received MMC, although the CI was wide (RR 0.73, 95% CI 0.09 to 5.66; I^2 = 18%).

Epitheliopathy

Nine studies (474 eyes) reported this complication following trabeculectomy. Those participants who received MMC were less likely to have an epitheliopathy following surgery than those who received 5-FU, which is most likely a result of the differences in the technique of antimetabolite application (RR 0.23, 95% CI 0.11 to 0.47; I^2 = 0%).

Tenon's cyst

Four studies (232 eyes) reported Tenon's cysts in their postoperative complication analysis. Those participants who received MMC were less likely to have a Tenon's cyst following surgery, although the CI was wide (RR 0.94, 95% CI 0.20 to 4.38; I^2 = 34%).

Hyphaema

Four studies (250 eyes) documented postoperative hyphaema during their follow-up of participants. Participants who received MMC were less likely to have a postoperative hyphaema than those who received 5-FU, which may be a consequence of antimetabolite application differences (RR 0.62, 95% CI 0.42 to 0.91; I^2 = 0%).

Endophthalmitis

Four studies (315 eyes) published rates of postoperative endophthalmitis. Participants receiving MMC were more likely to have endophthalmitis following trabeculectomy than those who received 5-FU. The CI was wide (RR 3.89, 95% CI 0.44 to 34.57; I^2 = 0%).

Sensitivity analyses (excluding studies at high risk of bias)

An interesting feature of these analyses was that the trials at high risk of bias were also the trials recruiting participants at high risk of failure. In general, excluding these studies improved the consistency (reduced I^2). Although the estimate of effect changed in these analyses, in general the conclusions (of uncertainty in most cases) did not.

Outcome	Name	All trials	Excluding trials at high risk of bias in 1 or more domains
Analysis 1.1*	Failure of functioning trabeculectomy at 1 year	RR 0.54, 95% CI 0.30 to 1.00	RR 1.02, 95% CI 0.50 to 2.04
Analysis 1.4*	Mean intraocular pressure at 1 year	MD -3.05 mmHg, 95% CI -4.60 to -1.50	MD -1.72 mmHg, 95% CI -3.28 to -0.16
Analysis 1.5**	Use of postoperative medication at 1 year	RR 1.03, 95% CI 0.57 to 1.85	RR 1.39, 95% CI 0.75 to 2.57
Analysis 1.6*	Mean number of postoperative medications at 1 year	MD -0.33, 95% CI -0.70 to 0.05	MD -0.08, 95% CI -0.27 to 0.11
Analysis 1.7**	Loss of 2 or more lines of visual acuity at 1 year	RR 1.05, 95% CI 0.54 to 2.06	RR 2.00, 95% CI 0.53 to 7.59

CI: confidence interval; MD: mean difference; RR: risk ratio

* The trials at high risk of bias were also the trials of the subgroup at high risk of failure.

** Two trials with a high risk of failure and one trial with a low risk of failure were excluded.

DISCUSSION

Summary of main results

We have summarised the results in the Summary of findings for the main comparison.

We identified 11 trials conducted in the United States, Europe, Asia and Africa. Five studies enrolled participants at low risk of trabeculectomy failure, five studies enrolled participants at high risk of failure, and one study enrolled people with both high and low risk of failure. None of the included trials enrolled participants with combined trabeculectomy/cataract surgery.

We considered one study to be at low risk of bias in all domains, six studies at high risk of bias in one or more domains, and the remaining four studies at an unclear risk of bias.

Our review showed that the risk of failure of trabeculectomy at one year after surgery was lower in those participants treated with MMC compared to those treated with 5-FU. However, the estimate of effect was imprecise, and we cannot exclude important differences. All 11 RCTs contributed to this finding with an overall mixed study population and varied methodology of antimetabolite application. Although MMC appeared to have a greater suc-

cess and more of an IOP-lowering effect in the higher-risk populations, we detected no significant difference between the subgroups of participants at low and high risk of failure in these analyses. We identified no difference between the visual outcomes of the people receiving MMC or 5-FU at one year postoperatively nor in the number of drops used postoperatively.

Evaluation of postoperative complications showed that there was a higher incidence of epitheliopathy and hyphaema when using 5-FU compared to MMC. However, we found those participants who received MMC to have more reported bleb leaks, wound leaks, late hypotony and cataract formation than those who received 5-FU. The quality of the evidence was low given that in general adverse outcomes were rare, and hence estimates of effect were imprecise. Although there were trends, any real significance cannot be determined from this review alone.

None of the studies reported quality of life.

Overall completeness and applicability of evidence

This review is limited owing to the small numbers of and large variability between studies, for example in participant demographics, methodology, masking of participants and varied follow-up. Some of the included studies had only 20 or 30 participants in their study population, which contrasts with the largest study, which had 115 participants.

After many years of widespread use of antimetabolite agents, uncertainty remains about the relative benefits and harms of their use in trabeculectomy surgery. Newer agents and techniques may

be developed and evaluated to then eventually take over the role of antimetabolites.

The majority of studies were carried out in the United States, although we included studies from European, Asian, African and Middle Eastern countries. Two of the included papers, one in Chinese and one in Italian, were translated. The analysis has taken into account the risk of failure of each study population and reported the risk as high or low. This is important when interpreting the results in a clinical setting in order to reflect the practice population. However, results showed a similar trend between high- and low-risk participants, which perhaps may be due to the inclusion of poor-quality evidence as discussed.

Quality of the evidence

Overall, we graded the quality of the evidence as low, in most cases because of risk of bias in the included studies and imprecision in the estimate of effect. One commonly encountered bias came from the difficulty in masking participants and surgeons owing to the different techniques of antimetabolite administration. All studies included in the review were RCTs, but the variability in outcome reporting reduced the quality of the evidence for some outcomes. Each study group reported few complications, which subsequently led to small numbers being incorporated into the analysis of complications.

Potential biases in the review process

We identified no obvious bias from the review process.

Agreements and disagreements with other studies or reviews

Fendi 2013 completed a meta-analysis that showed significantly higher success rates with the use of MMC when compared with 5-FU. This analysis included only five studies with participants who had recieved previous surgical treatment. Lin 2012 found in an analysis of eight studies that MMC achieved a significantly lower postoperative IOP than 5-FU, but MMC and 5-FU were comparable in achieving success. Likewise, Abdu 2010 found similar results to both Lin 2012 and this review with little difference between the two antimetabolites at achieving success. Abdu 2010 also also found that there was no difference in the mean postoperative IOP between participants who received MMC and partici-

pants who received 5-FU and suggest that further research in this area would enhance results to determine any true superiority of either MMC or 5-FU.

AUTHORS' CONCLUSIONS

Implications for practice

This review provided low-quality evidence that to achieve lower IOP following trabeculectomy MMC may be a more effective antimetabolite than 5-FU across both high- and low-risk populations. The risk associated with using either MMC or 5-FU as an antimetabolite in a routine trabeculectomy was low given the infrequent reporting of adverse outcomes.

Implications for research

Antimetabolites are a widely used adjunct in trabeculectomy surgery to help achieve lower postoperative IOP. However, the use of these medications may be associated with an increased risk of sight-threatening complications, predominantly due to the toxic effects on the conjunctiva and Tenon's capsule.

Further comparative research on MMC and 5-FU would be required to enhance reliability and validity of the results shown in this review. However, the development of newer, safer agents to control wound healing in glaucoma surgery may be of more benefit to patients in the longer term. These future agents would require full evaluation with well-designed trials to become integrated into clinical practice, particularly through the inclusion of trials with higher power to detect minimally important clinical differences and to consider cost and patient-orientated outcomes.

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* *Indicates the major publication for the study*

CHARACTERISTICS OF STUDIES

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

Katz 1995

Methods	Parallel-group randomised controlled trial, 1 eye per person
Participants	Country: USA Number of participants (eyes): 39 (39) % women: 67% Average age: 63 years (range not reported) Risk of trabeculectomy failure: high Inclusion criteria: • requiring trabeculectomy • history of prior cataract surgery, uveitic glaucoma, neovascular glaucoma, or previously failed filtering surgery Exclusion criteria: • younger than 18 years • corneal decompensation

Interventions	• MMC (20 eyes)
	  ○ Application: 1 intraoperative application
	  ○ Dose: 0.5 mg/ml for 5 minutes
	  ○ Location: between the conjunctiva and the episclera (n = 20 eyes)
	• 5-FU (19 eyes)
	  ○ Application: 10 postoperative injections (daily for 1 week, 3 times in following week)
	  ○ Dose: 5 mg
	  ○ Location: subconjunctival
	All surgeries involved a limbus-based conjunctival flap. Scleral flap was closed by 10-0 nylon sutures. Postoperative topical steroids were used in all participants and tapered over several weeks. 4 surgeons were involved in the study

Outcomes	Postoperative IOP Number of glaucoma medications used Change in visual acuity Follow-up: 1 and 2 years
Notes	Date study conducted: May 1990 to March 1991 Conflict of interest: None declared Funding source: Not reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Stated "randomised" but no elaboration of methods used

Katz 1995 *(Continued)*

Allocation concealment (selection bias)	Unclear risk	No mention of patient concealment of allocation
Blinding (performance bias and detection bias) All outcomes	High risk	Surgical method varied between both the groups, so masking for the surgeon and participant was impossible. No mention as to masking during the follow-up period
Incomplete outcome data (attrition bias) All outcomes	Low risk	Complete follow-up recorded for all participants with recognition of participants lost to follow-up
Selective reporting (reporting bias)	Low risk	All intended outcomes were identified and discussed
Other bias	High risk	Participants may have received different postoperative treatment: "The use of antibiotics, cycloplegics, digital massage and laser suture lysis were left to the discretion of the surgeon" Participants received different doses of 5-FU (average 46.0 mg, +/- 4.9 mg)

Kitazawa 1991

Methods	Parallel-group randomised controlled trial, 1 eye per person
Participants	Country: Japan Number of participants (eyes): 32 (32) % women: 38% Average age: 47 years (range 22 to 81) Risk of trabeculectomy failure: high Inclusion criteria: • failure of medical treatment to control IOP • 2 or more failed trabeculectomies, neovascular glaucoma, inflammatory glaucoma, congenital glaucoma, or aphakia Exclusion criteria: not reported
Interventions	• MMC (17 eyes) ○ Application: 1 intraoperative sponge application ○ Dose: 0.4 mg/ml for 5 minutes ○ Location: underneath the conjunctival flap and beneath the scleral flap • 5-FU (15 eyes) ○ Application: 10 postoperative injections (each day for 1 week and every other day for the following week) ○ Dose: 5 mg ○ Location: subconjunctival, 90 to 180 degrees away from the surgical site Following the trabeculectomy, 10-0 monofilament nylon suture was used for the scleral

Kitazawa 1991 (*Continued*)

flap, and 10-0 nylon shoelace suture was used for the conjunctival wound closure. Post-operatively 1.2 mg of dexamethasone was injected subconjunctivally. Topical atropine and antibiotics were given at the time of surgery. 0.1% betamethasone, 1% atropine sulfate and 0.3% ofloxacine were used as a standard for all participants postop

Outcomes	Mean IOP at 12 months Category 1 success: IOP controlled without antiglaucoma medication Category 2 success: IOP controlled with or without topical eye drops Category 3 success: IOP controlled without any medication or with oral carbonic anhydrase inhibitors in addition to topical medication Success was defined as IOP equal to or less than 20 mmHg without any medication Follow-up: 7 to 12 months
Notes	Date study conducted: December 1989 to November 1990 Conflict of interest: None declared Funding source: Research grant for Aging and Health from the Ministry of Health and Welfare, Japan

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	No clear methods described. Significant difference between the ages of each group (P = 0.01)
Allocation concealment (selection bias)	Unclear risk	Randomly allocated to intervention groups, but no elaboration on method used
Blinding (performance bias and detection bias) All outcomes	High risk	No masking due to nature of 2 techniques of administration for the interventions, and no mention of follow-up masking
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No statements about attrition or exclusion made
Selective reporting (reporting bias)	Low risk	Stated outcome measures were reported
Other bias	Low risk	The study appears to be free from other bias

Lamping 1995

Methods	Parallel-group randomised controlled trial, 1 or both eyes included
Participants	Country: USA Number of participants (eyes): 74 (80) % women: 41%

Lamping 1995 *(Continued)*

	Average age: 71 years (range not reported) Risk of trabeculectomy failure: high Inclusion criteria: • medically uncontrolled glaucoma and posterior lens implants, requiring glaucoma filtration surgery Exclusion criteria: • vitreous in the anterior chamber
Interventions	• MMC (40 eyes) ○ Application: 1 intraoperative sponge application ○ Dose: 0.4 mg/ml for 2.5 minutes ○ Location: between the conjunctival and scleral flap • 5-FU (40 eyes) ○ Application: 10 postoperative injections (once daily for first 10 days) ○ Dose: 5 mg ○ Location: subconjunctival, 180 degrees away from the surgical site No steroid or antibiotic was used at the time of surgery, but topical prednisolone, tobramycin and dexamethasone and atropine were applied postoperatively Single surgeon
Outcomes	Postoperative IOP Follow-up: week 1, week 2 and months 1, 2, 3, 6, 9 and 12
Notes	Date study conducted: Not reported Conflict of interest: None declared Funding source: Not reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Consecutive eyes were selected, no random sequence generation mentioned
Allocation concealment (selection bias)	Unclear risk	Randomised allocation to each intervention group, but no elaboration of methods used
Blinding (performance bias and detection bias) All outcomes	High risk	No masking due to different methods of application of the 2 interventions, and no mention of follow-up masking
Incomplete outcome data (attrition bias) All outcomes	Low risk	Recognition of 1 postop complication that stopped the use of antimetabolite therapy in this participant. This participant was not excluded from the study
Selective reporting (reporting bias)	Low risk	Stated outcome measures were reported

Other bias	Low risk	Participants in the 5-FU group received varied amounts of antimetabolite due to withholding of treatments if indicated by complications. 8 participants did not receive the full dose. This was taken into account in the data analysis

Mostafaei 2011

Methods	Parallel-group randomised controlled trial, 1 eye per person
Participants	Country: Iran Number of participants (eyes): 40 (40) % women: 19% Average age: 68 years (range 48 to 83) Risk of trabeculectomy failure: low Inclusion criteria: • open-angle glaucoma and uncontrolled IOP with evidence of optic nerve damage and visual field restriction Exclusion criteria: none reported

Interventions	• MMC (18 eyes)
	  ○ Application: intraoperative
	  ○ Dose: 0.02 mg
	  ○ Location: subconjunctival, 180 degrees away from operating site
	• 5-FU (22 eyes)
	  ○ Application: intraoperative
	  ○ Dose: 5 mg
	  ○ Location: subconjunctival

Outcomes	Primary outcome of successful surgery defined as an IOP of 6 to 22 mmHg at 6 months postoperatively Secondary outcome: complications identified at the 6-month follow-up IOP using Goldmann applanation Complications Follow-up: baseline, 2 weeks postoperatively, 1, 3 and 6 months
Notes	Date study conducted: Not reported Conflict of interest: None declared Funding source: Not reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Parallel trial design, but the details not described

Mostafaei 2011   *(Continued)*

Allocation concealment (selection bias)	Unclear risk	No report on concealment
Blinding (performance bias and detection bias) All outcomes	Unclear risk	No report on masking
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No report on loss to follow-up. One reported complication of surgery was observed but it was not made clear to which study group this participant belonged to
Selective reporting (reporting bias)	Low risk	All intended outcomes were identified and reported
Other bias	Low risk	The study appeared to be free from other bias

Singh 1997

Methods	Parallel-group randomised controlled trial, 1 eye per person
Participants	Country: Ghana Number of participants (eyes): 81 (81) % women: 40% Average age: 54 years (range not reported) Risk of trabeculectomy failure: high Inclusion criteria: • diagnosis of primary open-angle glaucoma Exclusion criteria: • No discussion about previous hypotensive drops. May have been primary trabeculectomies

Interventions	• MMC (44 eyes)
	  ○ Application: intraoperative sponge
	  ○ Dose: 0.5 mg/ml for 3.5 minutes
	  ○ Location: between scleral flap and conjunctiva
	• 5-FU (37 eyes)
	  ○ Application: intraoperative sponge
	  ○ Dose: 50 mg/ml for 5 minutes
	  ○ Location: between scleral flap and conjunctiva
	Limbal-based conjunctival flaps. Antimetabolite delivered with a sponge and thoroughly irrigated after required time. 5 surgeons with small variation on technique. Day 1 postop is when topical gentamycin, prednisolone acetate and atropine therapy started

Outcomes	IOP outcomes: < 21 mmHg, < 18 mmHg and < 15 mmHg Visual acuity Postoperative complications Follow-up: Post-operative days 1, 3, 7 and 14 and then average longer term follow up of

Singh 1997 (*Continued*)

	10 months (+/- 4.41)	
Notes	Date study conducted: Reported completed in 1995 Conflict of interest: Not reported Funding source: Not reported	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Treated decided by the flick of a coin in the operating theatre
Allocation concealment (selection bias)	Low risk	Surgeons masked from allocation up until time of surgery (minimal influence)
Blinding (performance bias and detection bias) All outcomes	High risk	Surgeons were masked up until time of surgery but not thereafter. Method of administration of treatment similar between groups. Follow-up team were not masked as to which antimetabolite the participant had received
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Attempts made to contact missing participants. No description about the number of participants lost to follow-up. 81/85 participants had at least 3 months' follow-up
Selective reporting (reporting bias)	High risk	No stated outcomes in the methods
Other bias	Unclear risk	Insufficient information

Singh 2000

Methods	Parallel-group randomised controlled trial, 1 eye per person Multicentre

Participants	Country: USA
	Number of participants (eyes): 108 (108)
	% women: not reported
	Average age: 66 years (range not reported)
	Risk of trabeculectomy failure: low
	Inclusion criteria:
	    • primary open-angle glaucoma, pseudoexfoliative glaucoma, or pigmentary glaucoma
	    • poorly controlled IOP despite maximal topical treatment
	Exclusion criteria:
	    • previous conjunctival or intraocular surgery

Singh 2000 (*Continued*)

Interventions	- MMC (54 eyes) - Application: intraoperative - Dose: 0.4 mg/ml for 2 minutes - Location: not stated 5-FU (54 eyes) - Application: intraoperative - Dose: 50 mg/ml for 5 minutes - Location: not stated Limbal-based conjunctival flaps, closed with 8-0 or 9-0 polyglactin 910 (Vicryl) suture. Antimetabolite delivered with a sponge and thoroughly irrigated after required time. 18 surgeons were involved in the study. 8 centres
Outcomes	This is the preliminary report with plans to extend follow-up time Main outcome measures were IOP and proportion of participants achieving successful outcomes, with varying IOP criteria for success (< 21 mmHg, < 18 mmHg, < 15 mmHg and < 12 mmHg). Post-operative visual acuity, complications and use of IOP-lowing medications were also included in the follow up data
Notes	Date study conducted: December 1996 Conflict of interest: None declared Funding source: Not reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation performed using a modified Moses-Oakford algorithm, and the randomisation envelope mailed to the study co-ordinators at the respective sites
Allocation concealment (selection bias)	Low risk	Participating surgeons were masked with regard to antimetabolite use until after participant enrolment into the study and written informed consent
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Surgeons were masked with regard to antimetabolite use until after participant enrolment into the study. Participants and follow-up period were not masked
Incomplete outcome data (attrition bias) All outcomes	Low risk	5 participants not included in analysis due to lack of pre- or intraoperative information
Selective reporting (reporting bias)	Low risk	Stated outcome measures in the methods were reported in the results

Singh 2000 (*Continued*)

Other bias	Low risk	The study appears free of other sources of bias

Sisto 2007

Methods	Parallel-group randomised controlled trial, 1 eye per person
Participants	Country: Italy Number of participants (eyes): 40 (40) % women: 35% Average age: 61 years (range 36 to 75) Risk of trabeculectomy failure: high Inclusion criteria: • neovascular glaucoma • IOP < 21 mmHg and resistant to medical therapy • no previous ocular surgery • best corrected visual acuity > -1.5 logMAR units Exclusion criteria: not reported

Interventions	• MMC (22 eyes)
	  ○ Application: intraoperative sponge application
	  ○ Dose: 0.2 mg/ml for 2 minutes
	  ○ Location: between the sclera and the Tenon's capsule
	• 5-FU (18 eyes)
	  ○ Application: postoperative injections, commencing on day 7, 2 injections per week for 2 weeks and then 1 injection per week for 6 weeks
	  ○ Dose: 0.1 ml of 50 mg/ml
	  ○ Location: subconjunctival injections near the bleb
	Fornix-based conjunctival flaps with single surgeon. No releasable sutures or suture lysis employed

Outcomes	Success defined as IOP < 21 mmHg at final postoperative visit Qualified success defined if IOP < 21 mmHg with addition of topical treatment. Failure is uncontrolled IOP equal or above 21 mmHg or vision dropped to no perception of light Follow-up: every 3 months in the first year, every 6 months thereafter to maximum 60 months (5 years)
Notes	Date study conducted: January 1993 to November 2000 Conflict of interest: Not reported Funding source: Not reported

Risk of bias

Bias	Authors' judgement	Support for judgement

Sisto 2007 (*Continued*)

Random sequence generation (selection bias)	Low risk	40 consecutive people with neovascular glaucoma selected. "All eyes had been assigned with a computer generated randomization code."
Allocation concealment (selection bias)	Unclear risk	No statement made
Blinding (performance bias and detection bias) All outcomes	Low risk	Surgeon not masked due to technique, but the follow-up staff were masked on collecting postoperative data
Incomplete outcome data (attrition bias) All outcomes	Low risk	All participant data included in analysis. However, length of follow-up was variable and no statement was made regarding the participants lost to follow-up
Selective reporting (reporting bias)	Low risk	Success criteria defined in the methods
Other bias	Unclear risk	No attempted power calculations

Uva 1996

Methods	Parallel-group randomised controlled trial, 1 eye per person
Participants	Country: Italy Number of participants (eyes): 30 (30) % women: 47% Average age: 54.1 years (range 45 to 60) Risk of trabeculectomy failure: low Inclusion criteria: - primary open-angle glaucoma uncontrolled with medication or laser therapy Exclusion criteria: - previous ocular surgery - aged 60 years or more - had been on antiglaucoma medication for less than 3 years
Interventions	- MMC (15 eyes) - Application: intraoperative sponge application - Dose: 0.2 mg/ml for 2 minutes - Location: between the sclera and the Tenon's capsule 5-FU (15 eyes) - Application: intraoperative sponge application - Dose: 50 mg/ml was applied for 5 minutes - Location: between the sclera and the Tenon's capsule Limbal flap was used that was closed with 10-0 nylon suture. 1% atropine, antibiotic and steroid was applied at the time of surgery. Conjunctiva was closed with 8-0 polyglactin synthetic suture

Uva 1996 (*Continued*)

Outcomes	Postoperative IOP Visual acuity Postoperative complications Follow-up: Mean follow-up 292 days +/- 46.1 days
Notes	Date study conducted: Not reported Conflict of interest: Not reported Funding source: Not reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomised with a "table of numbers" technique
Allocation concealment (selection bias)	Unclear risk	No statement made
Blinding (performance bias and detection bias) All outcomes	Unclear risk	Participant masking carried out given same surgical procedure for both antimetabolite interventions. Surgeons are presumed to not be masked given different duration of antimetabolite application. No mention of follow-up masking
Incomplete outcome data (attrition bias) All outcomes	Low risk	Commented on all intended outcomes. Short period of follow-up with all participants recorded within similar follow-up period
Selective reporting (reporting bias)	Low risk	Stated outcome measures in the methods were reported in the results
Other bias	Unclear risk	No obvious further bias

WuDunn 2002

Methods	Parallel-group randomised controlled trial, 1 or both eyes per person
Participants	Country: USA Number of participants (eyes): 103 (115) % women: 44% Average age: 65 years Risk of trabeculectomy failure: low Inclusion criteria: • poorly controlled IOP despite maximal tolerated medical treatment • primary open-angle glaucoma, pigmentary glaucoma, chronic angle-closure glaucoma, secondary glaucoma and pseudoexfoliation

	Exclusion criteria: • previous intraocular surgery

Interventions	• MMC (58 eyes)
	   ○ Application: intraoperative sponge application
	   ○ Dose: 0.2 mg/ml for 2 minutes
	   ○ Location: not stated
	• 5-FU (57 eyes)
	   ○ Application: intraoperative sponge application
	   ○ Dose: 50 mg/ml for 5 minutes
	   ○ Location: not stated
	Limbal-based conjunctival flaps, closed with 8-0 polyglactin 910 (Vicryl) suture. Antimetabolite was delivered with a cellulose sponge and thoroughly irrigated after required time. The application was divided into 2 phases to allow surgeon masking through same time of antimetabolitie/sham application. Corticosteroid, antibiotic ointment and atropine were instilled at the time of surgery. Postoperatively, all eyes received 1% prednisolone acetate, 1% atropine and an antibiotic

Outcomes	• Target IOP outcomes < 22, 19, 16, 13 with or without additional topical treatment were stated, and the IOP reduction had to be greater than 20%. • Visual acuity • Glaucoma medication needs • Complications • Fail defined as those participants whose preoperative IOP was less than 21 and did not have a 20% reduction; whose postoperative IOP was above the target level; or if further surgery to control IOP was required. • Follow-up: 6 and 12 months
Notes	Date study conducted: 1997 to 2001 Conflict of interest: Not reported Funding source: Research to Prevent Blindness, Inc, New York, New York NCT00346489

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	The assignment schedule was generated in blocks of 50 (25 per group) by a study co-ordinator who was not involved in the surgical procedure or clinical care. If the second eye of the participant was also enrolled, it was assigned to the opposite group of the first eye
Allocation concealment (selection bias)	Low risk	Group assignment was made randomly on the day of the surgery by the study co-ordinator and relayed directly to the operating room circulating nurse who prepared the

WuDunn 2002   *(Continued)*

		antimetabolite solutions
Blinding (performance bias and detection bias) All outcomes	Low risk	Only the study co-ordinator and the circulating nurse knew the allocation. The assignment code was kept in a locked drawer in the office of the study coordinator. The code was broken at 6 months postoperative to allow data analysis Surgeons were kept masked by using the same colour solution and the same duration of sponge application for both intervention groups (the MMC group had 3-minute application of balanced salt solution to equal the 5-minute 5-FU group in total)
Incomplete outcome data (attrition bias) All outcomes	Low risk	Clearly identified participants: 1 participant in each group failed to reach 6-month follow-up. By 1 year, the lost to follow-up were 9 in the 5-FU group and 4 in the MMC group
Selective reporting (reporting bias)	Low risk	Stated outcome measures in the methods were reported in the results
Other bias	Low risk	The study appears free of other sources of bias

Xinyu 2001

Methods	Parallel-group randomised controlled trial, 1 or both eyes included
Participants	Country: China Number of participants (eyes): 98 (108) % women: 57% Average age: 54 years (range 16 to 76) Risk of trabeculectomy failure: Mixed population: mostly low-risk population, 4 participants with previous surgery were termed "high risk" Inclusion criteria: Of those in the 5-FU group, 33 eyes were angle-closure glaucoma, 3 open-angle glaucoma, 2 eyes had glaucoma recurrent following previous control and 2 with previous glaucoma surgery. Of those in the MMC group, 25 had angle-closure glaucoma, 2 open-angle, 1 recurrent and 2 with previous glaucoma surgery. In the control group, 33 had angle-closure glaucoma, 3 open-angle and 2 with recurrence Exclusion criteria: Not mentioned

Interventions	• MMC (30 eyes)
	    ◦ Application: intraoperative sponge application
	    ◦ Dose: 0.2 mg/ml for 5 minutes
	    ◦ Location: not stated

	• 5-FU (40 eyes)
	  ○ Application: 6 to 8 postoperative injections on alternate days starting on day 3
	  ○ Dose: 5 mg/ml for 5 minutes
	  ○ Location: subconjunctival injections, 180 degrees away from the site of scleral flap
	All surgeons used same standard surgical technique.
	Control (untreated) group also included. (n = 38)

Outcomes	Measure of corneal scarring and corneal staining, postoperative IOP and occurrence of complications (conjunctivitis, vitreous detachment, hyphaema, corneal epithelial defect, hypotony and corneal ulcer) Successful reduction in IOP defined as < 21.06 mmHg. Follow-up: twice a week for 2 weeks, once a week for the subsequent 4 weeks, and then 1 or 2 times a month thereafter
Notes	Date study conducted: May 1995 to October 1999 Conflict of interest: Unable to ascertain with manuscript translation Funding source: Unable to ascertain with manuscript translation

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Random division into groups, but no detail given
Allocation concealment (selection bias)	Unclear risk	No information of allocation methods
Blinding (performance bias and detection bias) All outcomes	High risk	Application methods different for all 3 groups, and therefore difficult to mask surgeons and participants Follow-up of participants varied between 3 and 34 months with no clear statement about minimum length of follow-up
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No mention of attrition/exclusion
Selective reporting (reporting bias)	Low risk	All intended outcomes were addressed
Other bias	Low risk	No other obvious bias identified

Zadok 1995

Methods	Parallel-group randomised controlled trial, 1 eye per person
Participants	Country: Israel Number of participants (eyes): 20 (20) % women: 45% Average age: 69 years (range not reported) Risk of trabeculectomy failure: low Inclusion criteria: • uncontrolled primary open-angle glaucoma Exclusion criteria: Not mentioned

Interventions	• MMC (10 eyes)
	  ○ Application: intraoperative sponge application
	  ○ Dose: 0.2 mg/ml for 5 minutes
	  ○ Location: between the conjunctiva and episclera
	• 5-FU (10 eyes)
	  ○ Application: up to 7 postoperative injections, once daily in the first week after surgery
	  ○ Dose: 5 mg
	  ○ Location: subconjunctival injections, 180 degrees away from the site of surgery
	Closure of scleral flap by 10-0 nylon sutures. Conjunctiva closed by running suture. All participants received 1% atropine sulphate twice daily for 4 weeks and dexamethasone 4 times daily, tapered over several weeks

Outcomes	IOP < 21 mmHg as a primary outcome with or without antiglaucoma medication Follow-up: Participants reviewed at 1 week postoperatively, 2 weeks, 1, 2, 3, 6 and 12 months
Notes	Date study conducted: Not reported Conflict of interest: Not reported Funding source: Not reported

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	No mention of method of selection. Randomised
Allocation concealment (selection bias)	Unclear risk	No mention of how participants were concealed from their respective allocation
Blinding (performance bias and detection bias) All outcomes	Low risk	Surgeons not masked given different administration techniques. Follow-up completed by masked professionals
Incomplete outcome data (attrition bias) All outcomes	Low risk	Evidence from Table 3 (IOP distributions at 6 and 12 months) that all participants in

Zadok 1995 *(Continued)*

		study reached full follow-up period
Selective reporting (reporting bias)	Low risk	All outcomes reported in manuscript
Other bias	Low risk	No other bias evident

5-FU: 5-Fluorouracil
IOP: intraocular pressure
MMC: mitomycin C

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

Study	Reason for exclusion
Ashworth 2003	Prospective, non-randomised trial
Dreyer 1995	Not a randomised controlled study. No data on intraocular pressure as an outcome, therefore does not match inclusion criteria
Li 2001	Prospective, non-randomised trial
Membrey 2000	Retrospective study
Membrey 2001	Case-control; not a randomised controlled study
Oh 1994	Random allocation not mentioned; no reply from authors to request for clarification
Rodriguez-Bermejo 1993	Manuscript not available for review

Characteristics of studies awaiting assessment *[ordered by study ID]*

Liu 2015

Methods	
Participants	
Interventions	
Outcomes	
Notes	Awaiting response from authors regarding our query on methods of randomisation

Susanna 1995

Methods	
Participants	
Interventions	
Outcomes	
Notes	Unable to locate copy of the report

Yamamoto 1997

Methods	
Participants	
Interventions	
Outcomes	
Notes	Unable to locate copy of the report

DATA AND ANALYSES

Comparison 1. MMC versus 5-FU

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Failure of functioning trabeculectomy at one year	11	634	Risk Ratio (M-H, Random, 95% CI)	0.54 [0.30, 1.00]
1.1 Low risk of failure	6	370	Risk Ratio (M-H, Random, 95% CI)	0.65 [0.19, 2.20]
1.2 High risk of failure	5	264	Risk Ratio (M-H, Random, 95% CI)	0.49 [0.22, 1.08]
2 Failure of functioning trabeculectomy at one year in descending order of MMC exposure (dose x duration)	10	594	Risk Ratio (M-H, Random, 95% CI)	0.54 [0.30, 1.00]
3 Failure of functioning trabeculectomy at one year depending on 5-FU administration technique	10	594	Risk Ratio (M-H, Random, 95% CI)	0.54 [0.30, 1.00]
3.1 5-FU by postoperative injections	6	273	Risk Ratio (M-H, Random, 95% CI)	0.55 [0.27, 1.15]
3.2 5-FU by intraoperative sponge application	4	321	Risk Ratio (M-H, Random, 95% CI)	0.52 [0.13, 2.08]
4 Intraocular pressure at one year	7	386	Mean Difference (IV, Random, 95% CI)	-3.05 [-4.60, -1.50]
4.1 Low risk of failure	3	162	Mean Difference (IV, Random, 95% CI)	-1.72 [-3.28, -0.16]
4.2 High risk of failure	4	224	Mean Difference (IV, Random, 95% CI)	-4.18 [-6.73, -1.64]
5 Use of postoperative anti-glaucoma medications at final follow up	7	426	Risk Ratio (M-H, Random, 95% CI)	1.03 [0.57, 1.85]
5.1 Low risk of failure	4	273	Risk Ratio (M-H, Random, 95% CI)	1.11 [0.60, 2.07]
5.2 High risk of failure	3	153	Risk Ratio (M-H, Random, 95% CI)	0.99 [0.26, 3.76]
6 Mean number of postoperative anti-glaucoma medications	4	342	Mean Difference (IV, Random, 95% CI)	-0.33 [-0.70, 0.05]
6.1 Low risk of failure	2	223	Mean Difference (IV, Random, 95% CI)	-0.08 [-0.27, 0.11]
6.2 High risk of failure	2	119	Mean Difference (IV, Random, 95% CI)	-0.71 [-1.34, -0.09]
7 Loss of 2 or more lines of Snellen visual acuity postoperatively	5	328	Risk Ratio (M-H, Fixed, 95% CI)	1.05 [0.54, 2.06]
7.1 Low risk of failure	2	128	Risk Ratio (M-H, Fixed, 95% CI)	2.0 [0.53, 7.59]
7.2 High risk of failure	3	200	Risk Ratio (M-H, Fixed, 95% CI)	0.81 [0.36, 1.80]
8 Postoperative Complications	11		Risk Ratio (M-H, Random, 95% CI)	Subtotals only
8.1 Bleb leak	2	154	Risk Ratio (M-H, Random, 95% CI)	1.22 [0.32, 4.68]
8.2 Wound leak	6	391	Risk Ratio (M-H, Random, 95% CI)	1.17 [0.51, 2.71]
8.3 Late hypotony	4	211	Risk Ratio (M-H, Random, 95% CI)	1.37 [0.41, 4.63]
8.4 Maculopathy	4	342	Risk Ratio (M-H, Random, 95% CI)	1.71 [0.35, 8.33]
8.5 Cataract	4	275	Risk Ratio (M-H, Random, 95% CI)	1.73 [0.65, 4.61]
8.6 Shallow anterior chamber	5	311	Risk Ratio (M-H, Random, 95% CI)	1.22 [0.67, 2.21]
8.7 Choroidal detachment	8	494	Risk Ratio (M-H, Random, 95% CI)	0.86 [0.45, 1.63]
8.8 Epitheliopathy	8	419	Risk Ratio (M-H, Random, 95% CI)	0.23 [0.11, 0.47]
8.9 Tenon cyst	3	177	Risk Ratio (M-H, Random, 95% CI)	0.94 [0.20, 4.38]
8.10 Hyphaema	4	250	Risk Ratio (M-H, Random, 95% CI)	0.62 [0.42, 0.91]

8.11 Suprachoroidal haemorrhage		3	303	Risk Ratio (M-H, Random, 95% CI)				0.73 [0.09, 5.66]
8.12 Endophthalmitis		4	315	Risk Ratio (M-H, Random, 95% CI)				3.89 [0.44, 34.57]

ADDITIONAL TABLES

Table 1. Interventions

Study	MMC*				5-FU			
	Dose	Duration (minutes)	Location	Intraoperative or postoperative	Dose	Number of injections	Duration	Location
Katz 1995	0.5 mg/ml	5	Between the conjunctiva and the episclera	Postoperative	5 mg	10 (daily for 1 week, 3 times following week)	NA (injection)	Subconjunctival injection
Kitazawa 1991	0.4 mg/ml	5	Between the conjunctival and scleral flap	Postoperative	5 mg	10 (each day for 1 week and every other day for the following week)	NA (injection)	Subconjunctival injections, 90 to 180 degrees away from the surgical site
Lamping 1995	0.4 mg/ml	2.5	Between the conjunctival and scleral flap	Postoperative	5 mg	10 (first 10 days)	NA (injection)	Subconjunctival injection, 180 degrees from operating site
Mostafaei 2011	0.02 mg	not stated	Subconjunctival injection, 180 degrees away from operating site	Intraoperative	5 mg	NA	Not stated	Subconjunctival injection
Singh 1997	0.5 mg/ml	3.5	Between scleral flap and conjunctiva	Intraoperative	50 mg/ml	NA	5	Between scleral flap and conjunctiva
Singh 2000	0.4 mg/ml	2	Not stated	Intraoperative	50 mg/ml	NA	5	Not stated

Table 1. Interventions   (*Continued*)

Sisto 2007	0.2 mg/ml	2	Between the sclera and the Tenon's capsule	Postoperative	0.1 ml of 50 mg/ml	10 (starting on day 7, 2 injections per week for 2 weeks and then 1 injection per week for 6 weeks	NA (injection)	Subconjunctival injections near the bleb
Uva 1996	0.2 mg/ml	2	Between the sclera and the Tenon's capsule	Intraoperative	50 mg/ml	NA	5	Between the sclera and the Tenon's capsule
WuDunn 2002	0.2 mg/ml	2	Not stated	Intraoperative	50 mg/ml	NA	5	Not stated
Xinyu 2001	0.2 mg/ml	5	Not stated	Postoperative	5 mg	6 to 8 (alternate days, starting on day 3)	NA (injection)	Subconjunctival, 180 degrees away from the site of scleral flap
Zadok 1995	0.2 mg/ml	5	Between the conjunctiva and episclera	Postoperative	5 mg (0.5 ml of 10 mg/ml solution)	7 (once daily up to 7 times in the first week after surgery)	NA (injection)	Subconjunctival, 180 degrees from site of surgery

NA: not applicable
* All MMC only one intraoperative application

H I S T O R Y

Protocol first published: Issue 4, 2006

Review first published: Issue 11, 2015

Date	Event	Description
10 September 2008	Amended	Converted to new review format.

CONTRIBUTIONS OF AUTHORS

Conceiving the review: JCKC, PGS

Designing the review: JCKC, PGS

Coordinating the review: JCKC, EC

Undertaking manual searches: JCKC, PGS

Screening search results: JCKC, PGS

Organising retrieval of papers: JCKC, PGS, EC

Screening retrieved papers against inclusion criteria: JCKC, PGS, EC, JE

Appraising quality of papers: JCKC, PGS, EC

Abstracting data from papers: JCKC, PGS, EC, JE

Writing to authors of papers for additional information: JCKC, PGS, EC

Obtaining and screening data on unpublished studies: JCKC, PGS

Data management for the review: JCKC, PGS, EC, JE

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Interpretation of data: JCKC, PGS, EC, JE

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Securing funding for the review: JCKC, PGS

DECLARATIONS OF INTEREST

Emily Cabourne: None known.

Jonathan CK Clarke: None known.

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DIFFERENCES BETWEEN PROTOCOL AND REVIEW

We did not include outomes 1.4, 1.5 and 1.6 in the protocol. Use of postoperative glaucoma medication is a surrogate for partial trabeculectomy failure and was measured as an outcome in several of the included studies. Visual acuity was another commonly reported outcome in the included studies, in particular a loss of 2 lines of Snellen visual acuity. We therefore considered it appropriate to include these two outcomes in our review given their use in the assessment of trabeculectomy outcomes.

No data were available on time to failure as no studies were found to use Kaplan-Meier survival analysis as an outcome measure. Additionally, only one study commented clearly on non-attendance rates. No data were available on quality-of-life measures. We therefore did not report outcomes for time to failure, quality of life and non-attendance rate.

Late hypotony, endophthalmitis and choroidal detachment are of clinical interest and significance when concerning trabeculectomy. We therefore chose these adverse outcomes as priority in the Summary of findings for the main comparison.

INDEX TERMS

Medical Subject Headings (MeSH)

*Trabeculectomy; Antimetabolites [*therapeutic use]; Chemotherapy, Adjuvant; Cicatrix [prevention & control]; Fluorouracil [*therapeutic use]; Glaucoma [drug therapy; *surgery]; Intraocular Pressure [drug effects]; Mitomycin [*therapeutic use]; Randomized Controlled Trials as Topic; Risk; Treatment Failure; Wound Healing [*drug effects]

MeSH check words

Humans