

## Retinal detachment

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### ABSTRACT

**INTRODUCTION:** Rhegmatogenous retinal detachment (RRD) is the most common form of retinal detachment, where a retinal 'break' allows the ingress of fluid from the vitreous cavity to the subretinal space, resulting in retinal separation. It occurs in about 1 in 10,000 people a year. **METHODS AND OUTCOMES:** We conducted a systematic review and aimed to answer the following clinical questions: What are the effects of different surgical interventions in people with rhegmatogenous retinal detachment? What are the effects of interventions to treat proliferative vitreoretinopathy occurring as a complication of retinal detachment or previous treatment for retinal detachment? We searched: Medline, Embase, The Cochrane Library, and other important databases up to September 2013 (Clinical Evidence reviews are updated periodically; please check our website for the most up-to-date version of this review). We included harms alerts from relevant organisations such as the US Food and Drug Administration (FDA) and the UK Medicines and Healthcare products Regulatory Agency (MHRA). **RESULTS:** We found 14 studies that met our inclusion criteria. We performed a GRADE evaluation of the quality of evidence for interventions. **CONCLUSIONS:** In this systematic review, we present information relating to the effectiveness and safety of the following interventions: corticosteroids, daunorubicin, fluorouracil plus low molecular weight heparin, pneumatic retinopexy, scleral buckling, short-acting or long-acting gas tamponade, silicone oil tamponade, and vitrectomy.

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### INTERVENTIONS

#### DIFFERENT SURGICAL INTERVENTIONS FOR RHEGMATOGENOUS RETINAL DETACHMENT

##### Unknown effectiveness

Scleral buckling versus pneumatic retinopexy (there is consensus that both surgical techniques are effective: insufficient evidence to compare effects of scleral buckling versus pneumatic retinopexy) . . . . . 4

Scleral buckling versus primary vitrectomy (there is consensus that both surgical techniques are effective, but effects of scleral buckling compared with vitrectomy are unclear: in pseudophakic or aphakic rhegmatogenous retinal detachment [RRD], rate of retinal re-attachment after one operation may be lower post-scleral buckling compared with post-vitrectomy, but scleral buckling is associated with a lower rate of development or progression of cataract in phakic RRD) . . . . . 7

#### TREATING RRD ASSOCIATED WITH PROLIFERATIVE VITREORETINOPATHY

##### Likely to be beneficial

Silicone oil or long-acting gas tamponade (silicone oil and long-acting gas are equally effective in people receiving vitrectomy for RRD with severe proliferative vitreoretinopathy (PVR); silicone oil is more effective than short-acting gas at increasing re-attachment rates) . . . . . 1

##### Unknown effectiveness

Corticosteroid injection during vitrectomy surgery . . . . . 15  
 Daunorubicin infusion during vitrectomy surgery . . . . . 22  
 Fluorouracil plus low molecular weight heparin added to infusion solution during vitrectomy surgery . . . . . 25

### Key points

- Rhegmatogenous retinal detachment (RRD) is the most common form of retinal detachment, where a retinal 'break' allows the ingress of fluid from the vitreous cavity to the subretinal space, resulting in retinal separation. It occurs in about 1 in 10,000 people a year.  
 This review considers only acute progressive RRD.
- There is consensus that scleral buckling, pneumatic retinopexy, and vitrectomy are all effective for treating RRD.  
 We found insufficient evidence to assess effects of scleral buckling compared with [pneumatic retinopexy](#).  
 The effects of scleral buckling compared with [primary vitrectomy](#) are unclear. There is limited evidence that, in phakic RRD, scleral buckling improves visual acuity and is associated with a reduced risk of development or progression of cataract. However, we don't know whether scleral buckling is more effective than primary vitrectomy at increasing re-attachment rates in people with pseudophakic and aphakic RRD.
- In people undergoing vitrectomy for RRD with severe proliferative vitreoretinopathy (occurring as a complication of retinal detachment or previous treatment for retinal detachment), [silicone oil and long-acting gas](#) are equally effective for increasing re-attachment rates and improving visual acuity; silicone oil is better than short-acting gas.
- We found insufficient evidence assessing the effects of [fluorouracil plus heparin](#), [corticosteroids](#), or [daunorubicin](#) given during vitrectomy surgery for proliferative vitreoretinopathy.

**DEFINITION**

Retinal detachment can be defined as the separation of the neurosensory retina from the underlying retinal pigment epithelium (RPE). Direct apposition of the retina to the RPE is essential for normal retinal function, and retinal detachment involving the foveal centre leads to profound loss of vision in the affected eye.<sup>[1] [2]</sup> **Rhegmatogenous retinal detachment (RRD)** is the most common form of retinal detachment, where a retinal 'break' allows the ingress of fluid from the vitreous cavity to the subretinal space, resulting in retinal separation. Retinal break refers to a full-thickness defect in the neurosensory retina. Retinal breaks that develop from a tear in the retina at the time of posterior vitreous detachment (PVD) are usually referred to as retinal tears. Lattice degeneration can lead to the formation of circular retinal holes, which are typically referred to as atrophic holes. Retinal breaks can also develop as a result of trauma to and inflammation of the eye; examples include retinal dialysis, which is typically secondary to blunt trauma, and tears associated with retinal necrosis, resulting from trauma or inflammation. Rarer causes of retinal detachment include tractional retinal detachment, secondary to fibrous tissue on the surface of the retina; exudative retinal detachment, as a result of choroidal tumours that produce increased fluid flow through the subretinal space;<sup>[3]</sup> and ocular inflammatory conditions. Retinal detachments can also be a mixture of two or more of the above types. Asymptomatic and non-progressive chronic retinal detachment can also occur. This review considers only acute progressive RRD. **Diagnosis:** RRD is often, but not universally, associated with symptoms of flashes of light (retinal photopsia), visual floaters, and peripheral and usually progressive visual field loss. It is diagnosed by ophthalmoscopy. Acute RRD is seen as an oedematous folded retina with loss of the normal retinal transparency. The detachment can assume a bullous configuration that moves when the eye moves. There can be associated signs of PVD, as well as vitreous haemorrhage or RPE cells circulating in the vitreous cavity after retinal break formation. The presence of pigment cells in the anterior vitreous — visible on slit-lamp biomicroscopy (termed 'Shafer's sign') — is a sensitive indicator of the presence of a retinal break in a person presenting with an acute PVD.<sup>[4]</sup> Chronic retinal detachments can be associated with retinal cyst formation and 'tidemarks' demarcating the extent of the detachment, as well as subretinal fibrosis.<sup>[1] [2]</sup>

**INCIDENCE/  
PREVALENCE**

RRD can occur at any age, but reaches peak prevalence in people aged 60 to 70 years.<sup>[5] [6] [7]</sup> It affects men more than women, and white people more than black people. Observational studies from the US, Europe, and New Zealand found that non-traumatic, phakic (lens intact) RRD occurred in about 6 to 18 in 100,000 people a year (i.e., about 1/10,000).<sup>[5] [6] [7] [8] [9] [10] [11]</sup>

**AETIOLOGY/  
RISK FACTORS**

The occurrence of retinal detachment is related to the interplay between predisposing retinal lesions and vitreoretinal traction, and occurs when fluid moves from the vitreous cavity through a retinal break into the subretinal space.<sup>[12]</sup> Most (80%–90%) retinal detachments are associated with retinal-break formation at the time of PVD.<sup>[13] [14]</sup> PVD is a naturally occurring phenomenon, with a rapidly increasing prevalence in the 60- to 70-year-old age group. Most (70%) retinal breaks, formed at the time of PVD, are seen as tears in the retina or as holes with a free-floating retinal operculum. Retinal breaks can occur in areas of previously abnormal retina; for example, lattice degeneration.<sup>[14] [15]</sup> Symptoms and signs of acute PVD are known to be associated with a higher risk of immediate progression to RRD in people with predisposing retinal lesions. However, people with established (chronic) PVD and predisposing retinal lesions who have not immediately progressed to RRD are at lower risk than those without a PVD. Symptomatic retinal tears with persistent vitreoretinal traction (not a complete PVD) have a high rate of progression to retinal detachment (>50% if left untreated).<sup>[16]</sup> The risk of retinal detachment is increased to a variable extent in people with a symptomatic pre-existing retinal disease or lesions, especially retinal-flap tears, operculated retinal holes after separation of a retinal flap, atrophic retinal holes, lattice degeneration (areas of retinal thinning with abnormal vitreoretinal adhesion), and retinal dialyses. Autopsy studies have shown that about 6% to 11% of people aged over 20 years have retinal breaks in one form or another. However, the chances of an RRD occurring in an asymptomatic eye with a retinal break and with no history of fellow-eye RRD is 0.5% over a follow-up period of 11 years.<sup>[17] [18]</sup> Similarly, 7% to 8% of adults have areas of lattice degeneration, but only a small proportion of these lesions progress to RRD.<sup>[17] [19] [20] [21]</sup> Asymptomatic retinal dialysis is thought to have a high risk of progression to retinal detachment, especially after trauma.<sup>[22] [23]</sup> Increased risk of RRD is associated with several factors. There is a higher prevalence of RRD in short-sighted (myopic) people,<sup>[24]</sup> with around a 10-fold increased incidence in people with over three dioptres of myopia.<sup>[25]</sup> Approximately 50% of phakic RRD cases are myopic.<sup>[26]</sup> The fellow eye in people with an RRD is at a higher risk, with 2% to 10% of RRDs being bilateral.<sup>[27] [28] [29] [30] [31]</sup> Although some RRD occurring in a fellow eye will develop from pre-existing retinal lesions, most subsequent RRD (at least 50%, and possibly as high as 80%–90%) in the fellow eye will occur from ophthalmoscopically normal areas of retina,<sup>[14]</sup> and so prophylaxis to visible abnormal areas may not completely reduce the incidence of fellow-eye RRD. There is also a higher incidence of RRD in people with a family history of retinal detachment, especially in conditions such as Stickler's syndrome. People who have had previous cataract surgery also have a higher incidence of RRD, and about 1 in 5 patients presenting with RRD in the UK will have had cataract surgery.<sup>[26]</sup> About 0.5% to 0.6% of

people experience RRD after phacoemulsification surgery for cataracts, with the risk being increased by 15 to 20 times with rupture of the posterior capsule.<sup>[32]</sup> <sup>[33]</sup> About 10% of RRDs are associated with trauma. There are other conditions which, more rarely, increase the risk of RRD, including uveitis — especially CMV retinitis — and other degenerative retinal conditions, such as retinoschisis. Idiopathic macular holes may cause RRD in highly myopic eyes, but rarely in emmetropic or hypermetropic eyes.

<b>PROGNOSIS</b>	On presentation, retinal detachment is usually divided into 'macula on', when the fovea is still attached, and 'macula off', where the retina is detached centrally. <sup>[34]</sup> People with macula-on retinal detachments typically have good initial visual acuity and a better prognosis with successful surgery. Rapidly progressive cases are therefore treated as a matter of urgency. Macula-off retinal detachments have worse initial visual acuity, and have a worse prognosis even with successful re-attachment of the retina. Overall, about 95% of people have anatomically successful repair of RRD, with 70% to 90% achieving this in one operation. In 90% of successfully repaired macula-on retinal detachments, vision is 6/12 or better. However, in those with macula-off retinal detachments, only 50% of eyes achieve a visual acuity of 6/15, and, if the macula has been detached for 1 week or more, this level of visual acuity is rarely achieved. <sup>[34]</sup> Reasons for anatomical failure of surgery include new or missed retinal breaks and proliferative vitreoretinopathy (PVR). PVR is classified based on extent, position, and type of PVR: the American Retina Society proposed the first classification of PVR in 1983, <sup>[35]</sup> and, although updated in 1991 following the Silicone Oil Study, <sup>[36]</sup> this classification system continues to be widely used. Causes of poor visual acuity after successful repair include macular epiretinal membranes (fibrosis), cystoid macular oedema, and foveal photoreceptor degeneration in macula-off retinal detachments. <sup>[37]</sup>
<b>AIMS OF INTERVENTION</b>	To achieve retinal re-attachment in people with RRD; to achieve retinal re-attachment in people with PVR occurring as a complication of RRD or previous treatment for RRD; to achieve these aims with minimal re-operation rates and adverse effects of treatment.
<b>OUTCOMES</b>	<i>Treatment:</i> <b>Anatomical re-attachment rate</b> (after one operation and final rate), <b>re-operation rate, visual acuity</b> . <i>Treatment of eyes with proliferative vitreoretinopathy:</i> <b>rate of retinal re-attachment</b> (after one operation and final rate), <b>re-operation rate, visual acuity. Adverse effects.</b>
<b>METHODS</b>	<i>Clinical Evidence</i> search and appraisal September 2013. The following databases were used to identify studies for this systematic review: Medline 1966 to September 2013, Embase 1980 to September 2013, and The Cochrane Database of Systematic Reviews 2013, issue 9 (1966 to date of issue). Additional searches were carried out in the Database of Abstracts of Reviews of Effects (DARE) and Health Technology Assessment (HTA) Database. We also searched for retractions of studies included in the review. Titles and abstracts identified by the initial search, run by an information specialist, were first assessed against predefined criteria by an evidence scanner. Full texts for potentially relevant studies were then assessed against predefined criteria by an evidence analyst. Studies selected for inclusion were discussed with an expert contributor. All data relevant to the review were then extracted by an evidence analyst. Study design criteria for inclusion in this review were: published systematic reviews and RCTs, at least single-blinded, and containing more than 20 individuals of whom more than 80% were followed up. There was no minimum length of follow-up. We excluded all studies described as 'open', 'open label', or not blinded unless blinding was impossible. We included RCTs and systematic reviews of RCTs where harms of an included intervention were assessed, applying the same study design criteria for inclusion as we did for benefits. In addition, we use a regular surveillance protocol to capture harms alerts from organisations such as the FDA and the MHRA, which are added to the reviews as required. To aid readability of the numerical data in our reviews, we round many percentages to the nearest whole number. Readers should be aware of this when relating percentages to summary statistics such as relative risks (RRs) and odds ratios (ORs). We have performed a GRADE evaluation of the quality of evidence for interventions included in this review (see table, p 31 ). The categorisation of the quality of the evidence (high, moderate, low, or very low) reflects the quality of evidence available for our chosen outcomes in our defined populations of interest. These categorisations are not necessarily a reflection of the overall methodological quality of any individual study, because the Clinical Evidence population and outcome of choice may represent only a small subset of the total outcomes reported, and population included, in any individual trial. For further details of how we perform the GRADE evaluation and the scoring system we use, please see our website ( <a href="http://www.clinicalevidence.com">www.clinicalevidence.com</a> ).

**QUESTION** What are the effects of different surgical interventions in people with rhegmatogenous retinal detachment?

**OPTION** SCLERAL BUCKLING VERSUS PNEUMATIC RETINOPEXY

- For GRADE evaluation of interventions for Retinal detachment, see table, p 31 .
- There is consensus that scleral buckling and pneumatic retinopexy are both effective for treating rhegmatogenous retinal detachment (RRD). However, we found insufficient evidence to assess the effects of scleral buckling compared with pneumatic retinopexy.
- Scleral buckling has been associated with higher rates of refractive change (usually a myopic shift in refraction), diplopia with extraocular muscle dysfunction, and subretinal haemorrhage compared with pneumatic retinopexy.

**Benefits and harms**

**Scleral buckling versus pneumatic retinopexy:**

We found two RCTs. <sup>[38]</sup> <sup>[39]</sup>

**Re-attachment rate**

*Scleral buckling compared with pneumatic retinopexy* Scleral buckling and pneumatic retinopexy seem equally effective at increasing re-attachment rates (after one operation and final rate) in people with phakic, pseudophakic, or aphakic rhegmatogenous retinal detachment (RRD) and superior retinal breaks (*moderate-quality evidence*).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rates</b>					
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) rhegmatogenous retinal detachment (RRD), involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe proliferative vitreoretinopathy (PVR)	<b>Re-attachment rates after one operation , 6 months</b> 78/95 (82%) eyes with scleral buckling 75/103 (73%) eyes with pneumatic retinopexy	Reported as not significant P value not reported	↔	Not significant
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) RRD, involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe PVR	<b>Final re-attachment rates</b> 93/95 (98%) eyes with scleral buckling 102/103 (99%) eyes with pneumatic retinopexy	Reported as not significant P value not reported	↔	Not significant
[39] RCT	20 people with RRD and single retinal break or small group of breaks in phakic or pseudophakic eyes without severe PVR	<b>Re-attachment rate</b> 8/10 (80%) with scleral buckling 7/10 (70%) with pneumatic retinopexy	Significance not assessed		

**Visual acuity**

*Scleral buckling compared with pneumatic retinopexy* Scleral buckling and pneumatic retinopexy seem equally effective at improving visual acuity in people with phakic, pseudophakic, or aphakic rhegmatogenous retinal detachment and

superior retinal breaks. However, in eyes with preoperative detachment of the macula for up to 14 days, scleral buckling seems less effective at improving visual acuity (moderate-quality evidence).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) rhegmatogenous retinal detachment (RRD), involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe proliferative vitreoretinopathy (PVR)	<b>Proportion of eyes with visual acuity of 20/50 or better on the Snellen scale , 6 months</b> 64/95 (68%) eyes with scleral buckling 90/103 (87%) eyes with pneumatic retinopexy See further information on studies for details on final visual outcome after successful repair compared with failed surgery	Significance not assessed		
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) RRD, involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe PVR	<b>Proportion of eyes with visual acuity of 20/25</b> 12/95 (13%) eyes with scleral buckling 25/103 (24%) eyes with pneumatic retinopexy See further information on studies for details on final visual outcome after successful repair compared with failed surgery	Reported as not significant P = 0.05	↔	Not significant
[38] RCT	109 eyes with preoperative detachment of the macula for up to 14 days Subgroup analysis There were insufficient data to assess eyes with macula detachment for more than 14 days	<b>Proportion of eyes with visual acuity of 20/50 or better on the Snellen scale</b> 27/48 (56%) eyes with scleral buckling 49/61 (80%) eyes with pneumatic retinopexy	P = 0.01	○○○	pneumatic retinopexy

No data from the following reference on this outcome. [39]

**Re-operation rate**

No data from the following reference on this outcome. [38] [39]

**Adverse effects**

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Adverse effects</b>					
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic	<b>Proportion of eyes with at least 1 dioptre of myopia</b> 65/95 (68%) eyes with scleral buckling	P = 0.0001	○○○	pneumatic retinopexy

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
	(20 eyes) rhegmatogenous retinal detachment (RRD), involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe proliferative vitreoretinopathy (PVR)	3/103 (3%) eyes with pneumatic retinopexy			
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) RRD, involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe PVR	<b>Persistent diplopia</b> 3/95 (3%) eyes with scleral buckling 0/103 (0%) eyes with pneumatic retinopexy	Significance not assessed		
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) RRD, involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe PVR	<b>Proliferative vitreoretinopathy (PVR)</b> 5/95 (5%) eyes with scleral buckling 3/103 (3%) eyes with pneumatic retinopexy	Significance not assessed		
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) RRD, involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe PVR	<b>Macular pucker requiring surgery</b> 2/95 (2%) eyes with scleral buckling 0/103 (0%) eyes with pneumatic retinopexy	Significance not assessed		
[38] RCT	198 eyes with phakic (108 eyes), pseudophakic (70 eyes), or aphakic (20 eyes) RRD, involving retinal breaks within 1 disc diameter of each other located in the superior retina, without severe PVR	<b>Macular subretinal haemorrhage</b> 2/95 (2%) eyes with scleral buckling 0/103 (0%) eyes with pneumatic retinopexy	Significance not assessed		
[38] RCT	108 phakic eyes Subgroup analysis	<b>Progressive lens opacities , 24 months</b> 21/44 (47%) eyes with scleral buckling 10/53 (19%) eyes with pneumatic retinopexy	Significance not assessed		



Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
[38] RCT	108 phakic eyes Subgroup analysis	<b>Proportion of people requiring cataract surgery</b> 8/44 (18%) eyes with scleral buckling 2/53 (4%) eyes with pneumatic retinopexy	Significance not assessed		
[39] RCT	20 people with RRD and single retinal break or small group of breaks in phakic or pseudophakic eyes without severe PVR	<b>Proliferative vitreoretinopathy (PVR) with RRD</b> 0/10 (0%) eyes with scleral buckling 2/10 (20%) eyes with pneumatic retinopexy One person in each group had re-detachment because of new retinal holes	Significance not assessed		

#### Further information on studies

[38] The RCT found no significant difference in final visual outcome between eyes with successful RRD repair with initial scleral buckling surgery and eyes that failed with initial pneumatic retinopexy and required further surgery (P >0.05; absolute data for final vision outcome in each group not reported).

#### Comment:

##### Clinical guide:

Rhegmatogenous retinal detachment (RRD) is repaired using techniques to close retinal breaks and relieve vitreoretinal traction. Although some RRDs could potentially be repaired by all three surgical techniques (scleral buckling, [vitrectomy](#), or pneumatic retinopexy), this is not universally the case, and choice of surgery will depend on various factors, including the number, location, and size of retinal breaks present; the ability of the patient to posture to position tamponade agents in the correct place; lens status; and surgeon experience (including access to equipment). Vitrectomy techniques require specialist training, and equipment is expensive; access is thus limited in resource-poor areas. In clinical practice in the UK, Europe, and North America, people with phakic eyes and localised RRD with small anterior holes or [retinal dialysis](#) are usually treated with scleral buckling, especially if there is no associated [posterior vitreous detachment](#). Eyes in which a scleral buckle cannot be placed (e.g., thin sclera) and people with vitreous opacity obstructing the retinal view, giant retinal breaks, or very posterior retinal breaks are usually treated with vitrectomy. Pneumatic retinopexy is usually reserved for people with a single or localised group of breaks in the superior retina. People with pseudophakic RRD represent about 20% of all RRD that presents in clinical practice in the UK. Retinal breaks in these cases are often small and difficult to see because of the intra-ocular lens and capsule remnants restricting the fundal view. There is an increasing trend to treat these people with vitrectomy rather than scleral buckling or pneumatic retinopexy, which allows accurate break localisation with the technique of internal searching. Furthermore, with the eye already being pseudophakic, a common adverse effect of vitrectomy surgery — cataract formation — is avoided, as well as the occasional refractive and extra-ocular muscle imbalance changes associated with scleral buckling. In some clinical situations, surgeons may choose vitrectomy surgery combined with scleral buckling; for example, in cases with superior and inferior retinal breaks, especially if there is pre-existing proliferative vitreoretinopathy.

#### OPTION SCLERAL BUCKLING VERSUS PRIMARY VITRECTOMY

- For GRADE evaluation of interventions for Retinal detachment, [see table, p 31](#).
- There is consensus that [scleral buckling](#) and primary [vitrectomy](#) are both effective for treating rhegmatogenous retinal detachment (RRD).
- The effects of scleral buckling compared with primary vitrectomy are unclear. There is limited evidence that, in phakic RRD, scleral buckling improves visual acuity and is associated with a reduced risk of development or

progression of cataract. However, we don't know whether scleral buckling is more effective than primary vitrectomy at increasing re-attachment rates in people with pseudophakic and aphakic RRD.

- Vitrectomy has been associated with higher rates of cataract formation in phakic eyes compared with scleral buckling.

## Benefits and harms

### Scleral buckling versus primary vitrectomy in pseudophakic or aphakic rhegmatogenous retinal detachment (RRD):

We found two systematic reviews (search date 2010<sup>[40]</sup> and search date 2011<sup>[41]</sup>) comparing scleral buckling with primary vitrectomy in people with pseudophakic or aphakic RRD. Both reviews included the same six RCTs, but they reported some different outcomes, so both are reported here.

#### Re-attachment rate

*Scleral buckling compared with primary vitrectomy in people with pseudophakic or aphakic rhegmatogenous retinal detachment* We don't know whether scleral buckling is more effective than primary vitrectomy at increasing re-attachment rates after one operation at 6 months to 1 year, but it may be less effective than vitrectomy at increasing secondary re-attachment rates (*low-quality evidence*).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rates</b>					
[40] Systematic review	780 pseudophakic/aphakic subjects 5 RCTs in this analysis	<b>Primary re-attachment after the first intervention , 6 months or 1 year</b> 274/407 (67%) with scleral buckling 279/373 (75%) with pars plana vitrectomy	OR 1.46 95% CI 0.79 to 2.71 P = 0.23 Heterogeneity I <sup>2</sup> = 65% P = 0.02	↔	Not significant
[40] Systematic review	690 pseudophakic/aphakic subjects 4 RCTs in this analysis	<b>Secondary re-attachment , at end of study</b> 327/359 (91%) with scleral buckling 317/331 (96%) with pars plana vitrectomy	OR 2.08 95% CI 1.08 to 4.03 P = 0.03	●●○	vitrectomy

#### Visual acuity

*Scleral buckling compared with primary vitrectomy* We don't know how scleral buckling and primary vitrectomy compare at improving visual acuity at 6 months or longer in people with pseudophakic or aphakic rhegmatogenous retinal detachment (*very low-quality evidence*).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[40] Systematic review	743 pseudophakic/aphakic subjects 5 RCTs in this analysis	<b>Mean BCVA (logMAR units) , 6 months or less</b> with scleral buckling with pars plana vitrectomy Absolute results not reported	Mean difference -0.03 95% CI -0.01 to +0.04 P = 0.36 Authors note that some people had secondary procedures; and outcome of this size may not be clinically relevant	↔	Not significant
[41] Systematic review	465 pseudophakic/aphakic subjects 3 RCTs in this analysis	<b>Final visual success (preservation or improvement in visual acuity)</b> 202/233 (87%) with scleral buckling 210/232 (91%) with pars plana vitrectomy	OR 1.49 95% CI 0.82 to 2.68 P = 0.19	↔	Not significant



**Re-operation rate**

*Scleral buckling compared with primary vitrectomy in people with pseudophakic or aphakic rhegmatogenous retinal detachment* Scleral buckling and primary vitrectomy seem equally effective at reducing re-operation rate (*moderate-quality evidence*).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-operation rate</b>					
[42] RCT	150 people, 150 eyes, with pseudophakic rhegmatogenous retinal detachment In review [40] [41]	<b>Re-operation rate</b> 13/75 (17%) with scleral buckling 4/75 (5%) with pars plana vitrectomy with infusion of short-acting gas	P = 0.38	↔	Not significant

No data from the following reference on this outcome. [43] [44] [45]

**Adverse effects**

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Diplopia</b>					
[40] Systematic review	225 pseudophakic/aphakic subjects Subgroup analysis 1 RCT in this analysis [44]	<b>Diplopia/EOM dysfunction</b> 0/99 (0%) with pars plana vitrectomy 5/126 (4%) with scleral buckling	OR 0.11 95% CI 0.01 to 2.03	↔	Not significant
[40] Systematic review	150 pseudophakic/aphakic subjects Subgroup analysis 1 RCT in this analysis [42]	<b>Diplopia/EOM dysfunction</b> 0/75 (0%) with pars plana vitrectomy 3/75 (4%) with scleral buckling	OR 0.14 95% CI 0.01 to 2.70	↔	Not significant
[40] Systematic review	50 pseudophakic/aphakic subjects Subgroup analysis 1 RCT in this analysis [43]	<b>Diplopia/EOM dysfunction</b> 0/25 (0%) with pars plana vitrectomy 1/25 (4%) with scleral buckling	OR 0.32 95% CI 0.01 to 8.25	↔	Not significant
<b>Proliferative vitreoretinopathy</b>					
[41] Systematic review	465 pseudophakic/aphakic subjects 3 RCTs in this analysis	<b>PVR</b> 76/359 (21%) with scleral buckling 59/331 (18%) with pars plana vitrectomy	OR 0.85 95% CI 0.58 to 1.26 P = 0.42	↔	Not significant

**Scleral buckling versus primary vitrectomy in phakic rhegmatogenous retinal detachment (RRD):**

We found two systematic reviews (search date 2010 [40] and search date 2011 [41]) comparing scleral buckling with primary vitrectomy in people with phakic rhegmatogenous retinal detachment (RRD). Both reviews included the same six RCTs, but they reported some different outcomes, so both are reported here.

**Re-attachment rate**

*Scleral buckling compared with primary vitrectomy in people with phakic rhegmatogenous retinal detachment* We don't know how scleral buckling and primary vitrectomy compare at increasing re-attachment rates after one operation at 6 months to 1 year ([low-quality evidence](#)).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rate</b>					
[40] Systematic review	523 phakic subjects 3 RCTs in this analysis	<b>Primary re-attachment after the first intervention , at 6 months or 1 year</b> 179/263 (68%) with scleral buckling 177/260 (68%) with pars plana vitrectomy	OR 1.00 95% CI 0.69 to 1.46 P = 0.99	↔	Not significant
[40] Systematic review	523 phakic subjects 3 RCTs in this analysis	<b>Secondary re-attachment , at end of study</b> 256/263 (97%) with scleral buckling 253/260 (97%) with pars plana vitrectomy	OR 0.99 95% CI 0.34 to 2.87 P = 0.99	↔	Not significant

**Visual acuity**




*Scleral buckling compared with primary vitrectomy* We don't know how scleral buckling and primary vitrectomy compare at improving visual acuity at 6 months or longer in people with phakic rhegmatogenous retinal detachment as clinical relevance is unclear ([very low-quality evidence](#)).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[40] Systematic review	477 phakic subjects 2 RCTs in this analysis	<b>Mean BCVA (logMAR units) , 6 months or less</b> with scleral buckling with pars plana vitrectomy Absolute results not reported	Mean difference 0.14 95% CI 0.06 to 0.21 P = 0.0004 Authors note that some people had secondary procedures; and outcome of this size may not be clinically relevant		scleral buckling
[41] Systematic review	440 phakic subjects 3 RCTs in this analysis	<b>Final visual success (preservation or improvement in visual acuity)</b> 233/263 (89%) with scleral buckling 207/260 (80%) with pars plana vitrectomy	OR 0.50 95% CI 0.31 to 0.82 P = 0.005	●●○	scleral buckling

**Re-operation rate**

No data from the following reference on this outcome. [\[40\]](#) [\[41\]](#)

**Adverse effects**

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Cataracts</b>					
[41] Systematic review	477 phakic subjects 2 RCTs in this analysis	<b>Cataract</b> 96/240 (40%) with scleral buckling 165/237 (70%) with pars plana vitrectomy	OR 4.18 95% CI 2.75 to 6.35 P <0.00001		scleral buckling
<b>Proliferative vitreoretinopathy</b>					
[40] Systematic review	523 phakic subjects 3 RCTs in this analysis	<b>PVR</b> 27/263 (10%) with scleral buckling 39/260 (15%) with pars plana vitrectomy	OR 1.54 95% CI 0.91 to 2.59 P = 0.11		Not significant
<b>Other adverse effects</b>					
[40] Systematic review	46 phakic subjects Subgroup analysis 1 RCT in this analysis [46]	<b>Diplopia/EOM dysfunction</b> 2/23 (7%) with pars plana vitrectomy 3/23 (13%) with scleral buckling	OR 0.63 95% CI 0.10 to 4.21		Not significant

**Comment:** Cataract formation is more common after vitrectomy surgery than after scleral buckling surgery. The power of the included studies to detect differences in less frequent adverse outcomes, including diplopia and orbital/scleral buckle infection is limited. These can occur in scleral buckling surgery but are very rare in vitrectomy surgery. Similarly, endophthalmitis occurs with an incidence of approximately 1 in 2000 in vitrectomy surgery, but would not occur in completely extra-ocular scleral buckling surgery.

**Clinical guide:**

See scleral buckling versus pneumatic retinopexy, p 4 .

**QUESTION** What are the effects of interventions to treat rhegmatogenous retinal detachment associated with proliferative vitreoretinopathy?

**OPTION** DIFFERENT SUBSTANCES FOR TAMPONADE

- For GRADE evaluation of interventions for Retinal detachment, see table, p 31 .
- In people undergoing vitrectomy for rhegmatogenous retinal detachment (RRD) with severe proliferative vitreoretinopathy (occurring as a complication of retinal detachment or previous treatment for retinal detachment), silicone oil and long-acting gas are equally effective for increasing re-attachment rates and improving visual acuity; silicone oil is better than short-acting gas.
- Silicone oil may cause less hypotony compared with long-acting gas, especially in people with severe anterior proliferative vitreoretinopathy.

**Benefits and harms**

**Silicone oil tamponade versus long-acting gas tamponade:**

We found one systematic review (search date 2009), [47] which identified one RCT [48] comparing silicone oil tamponade versus long-acting gas (C<sub>3</sub>F<sub>8</sub>, perfluoropropane) in people with severe proliferative vitreoretinopathy (PVR) receiving vitrectomy. The RCT compared treatments in two distinct groups: 131 eyes undergoing initial vitrectomy, and 134 eyes undergoing a second vitrectomy after previous failed vitrectomy surgery. [48] We also identified two further reports of the RCT identified by the review, which reported on only adverse effects. [49] [50]

**Re-attachment rate**

*Silicone oil tamponade compared with long-acting gas tamponade* Silicone oil tamponade and long-acting gas tamponade seem equally effective at increasing re-attachment rates in people with severe proliferative vitreoretinopathy undergoing initial or repeat vitrectomy for rhegmatogenous retinal detachment (**moderate-quality evidence**).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rate</b>					
[48] RCT	131 eyes undergoing initial vitrectomy for rhegmatogenous retinal detachment (RRD) with severe proliferative vitreoretinopathy (PVR) In review [47]	<b>Re-attachment , last follow-up examination (up to 36 months)</b> 38/59 (64%) with silicone oil 45/62 (73%) with C <sub>3</sub> F <sub>8</sub> Last observation carried forward (LOCF) analysis See Further information on studies for details of follow-up at different time frames	P = 0.33	↔	Not significant
[48] RCT	134 eyes undergoing a second vitrectomy after previous failed vitrectomy surgery In review [47]	<b>Re-attachment , last follow-up examination (up to 36 months)</b> 37/61 (61%) with silicone oil 50/68 (74%) with C <sub>3</sub> F <sub>8</sub> LOCF analysis See Further information on studies for details of follow-up at different time frames	Significance not assessed		

**Visual acuity**

*Silicone oil tamponade compared with long-acting gas tamponade* Silicone oil tamponade and long-acting gas tamponade seem equally effective at improving visual acuity in people with severe proliferative vitreoretinopathy undergoing initial or repeat vitrectomy for rhegmatogenous retinal detachment (**high-quality evidence**).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[48] RCT	131 eyes undergoing initial vitrectomy for rhegmatogenous retinal detachment (RRD) with severe proliferative vitreoretinopathy (PVR) In review [47]	<b>Proportion who achieved a visual acuity of 5/200 or better , last follow-up examination (up to 36 months)</b> 29/64 (45%) with silicone oil 29/67 (43%) with C <sub>3</sub> F <sub>8</sub> Last observation carried forward (LOCF) analysis See Further information on studies for details of follow-up at different time frames	P = 0.82	↔	Not significant
[48] RCT	134 eyes undergoing a second vitrectomy after previous failed vitrectomy surgery In review [47]	<b>Proportion who achieved a visual acuity of 5/200 or better , last follow-up examination (up to 36 months)</b> 21/63 (33%) with silicone oil 27/71 (38%) with C <sub>3</sub> F <sub>8</sub> LOCF analysis See Further information on studies for details of follow-up at different time frames	P = 0.57	↔	Not significant

**Re-operation rate**

No data from the following reference on this outcome. <sup>[48]</sup>

**Adverse effects**


Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Adverse effects</b>					
<sup>[48]</sup> RCT	265 eyes undergoing initial or repeat vitrectomy for rhegmatogenous retinal detachment (RRD) with severe proliferative vitreoretinopathy (PVR) In review <sup>[47]</sup>	<b>Keratopathy , last follow-up examination (up to 36 months)</b> 30% with silicone oil 33% with C <sub>3</sub> F <sub>8</sub> Absolute numbers not reported Last observation carried forward (LOCF) analysis See Further information on studies for details of follow-up at different time frames	P = 0.70	↔	Not significant
<sup>[49]</sup> RCT	245 eyes undergoing initial or repeat vitrectomy for RRD with severe PVR Further report of reference <sup>[48]</sup>	<b>Elevated intraocular pressure (raised intraocular pressure to &gt;25 mmHg on 2 or more consecutive visits) , 6 months</b> 9/120 (8%) eyes with silicone oil 2/121 (2%) eyes with C <sub>3</sub> F <sub>8</sub>	P <0.05	○○○	C <sub>3</sub> F <sub>8</sub>
<sup>[49]</sup> RCT	245 eyes undergoing initial or repeat vitrectomy for RRD with severe PVR Further report of reference <sup>[48]</sup>	<b>Chronic hypotony (intra-ocular pressure &lt;5 mmHg on 2 or more consecutive or 3 visits) , 6 months</b> 21/120 (18%) eyes with silicone oil 37/121 (31%) eyes with C <sub>3</sub> F <sub>8</sub>	P <0.05	○○○	silicone oil
<sup>[50]</sup> RCT	245 eyes undergoing initial or repeat vitrectomy for RRD with severe PVR Further report of reference <sup>[48]</sup>	<b>Macular pucker , 6 months</b> 12% with silicone oil 19% with C <sub>3</sub> F <sub>8</sub> Absolute numbers not reported	P = 0.15	↔	Not significant

**Silicone oil tamponade versus short-acting gas tamponade:**

We found one systematic review (search date 2009), <sup>[47]</sup> which identified one RCT comparing **silicone oil tamponade** versus short-acting gas (SF<sub>6</sub>, sulphur hexafluoride) in people with severe **proliferative vitreoretinopathy (PVR)** without previous vitrectomy receiving **vitrectomy**. <sup>[51]</sup>


**Re-attachment rate**

*Silicone oil tamponade compared with short-acting gas tamponade* Silicone oil tamponade seems more effective at increasing re-attachment rates at 6 months in people with severe proliferative vitreoretinopathy undergoing initial vitrectomy for rhegmatogenous retinal detachment (**moderate-quality evidence**).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rate</b>					
[51] RCT	101 eyes undergoing initial <b>vitrectomy</b> for rhegmatogenous retinal detachment (RRD) with severe <b>proliferative vitreoretinopathy (PVR)</b> In review [47]	<b>Re-attachment , 6 months</b> 31/51 (61%) with silicone oil 23/46 (50%) with SF <sub>6</sub> See Further information on studies for rate of eyes with subtotal retinal attachment but subsequent successful macular attachment	P <0.05		silicone oil

**Visual acuity**



*Silicone oil tamponade compared with short-acting gas tamponade* Silicone oil tamponade seems more effective at improving visual acuity at 6 months in people with severe proliferative vitreoretinopathy undergoing vitrectomy for rhegmatogenous retinal detachment (**moderate-quality evidence**).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[51] RCT	101 eyes undergoing initial <b>vitrectomy</b> for rhegmatogenous retinal detachment (RRD) with severe <b>proliferative vitreoretinopathy (PVR)</b> In review [47]	<b>Proportion of eyes with 5/200 or better on the Snellen scale , 6 months</b> 31/51 (61%) with silicone oil 15/46 (33%) with SF <sub>6</sub>	P <0.05		silicone oil

**Re-operation rate**

No data from the following reference on this outcome. [51]

**Adverse effects**

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Adverse effects</b>					
[51] RCT	101 eyes undergoing initial <b>vitrectomy</b> for rhegmatogenous retinal detachment (RRD) with severe <b>proliferative vitreoretinopathy (PVR)</b> In review [47]	<b>Keratopathy</b> 10/47 (21%) with silicone oil 19/40 (48%) with SF <sub>6</sub> See Further information on studies for rate of keratopathy in eyes with detached macula	P = 0.01		silicone oil
[51] RCT	101 eyes undergoing initial vitrectomy for RRD with severe PVR In review [47]	<b>Hypotony , 24 months</b> 5/47 (11%) with silicone oil 7/40 (18%) with SF <sub>6</sub> See Further information on studies for rate of hypotony in eyes with detached macula	P = 0.35		Not significant



**Further information on studies**

- [48] The RCT aimed to follow people up for 36 months. Data were available for 100% of eyes at 3 months, 91% to 95% of eyes at 12 months, 64% to 73% of eyes at 24 months, and 49% to 53% of eyes at 36 months.
- [51] The RCT found that silicone oil significantly increased the proportion of eyes with subtotal retinal attachment but successful macular attachment at 6 months post-operatively compared with SF<sub>6</sub> (10/51 [20%] with silicone oil v 5/46 [11%] with SF<sub>6</sub>; P <0.05). Rates of both hypotony and keratopathy were higher in eyes with detached macula, although differences between groups were not significant (reported as not significant; P value not reported).

**Comment:**

**Clinical guide:**

In people with rhegmatogenous retinal detachment and advanced proliferative vitreoretinopathy (PVR), the PVR-associated membranes can sometimes prevent closure of retinal breaks when using either [scleral buckling surgery](#) or [pneumatic retinopexy](#). In this situation, vitrectomy surgery may be indicated to allow the surgical removal of these membranes, and hence allow retinal re-attachment. Tamponade of retinal breaks postoperatively can be achieved with long-acting gas or silicone oil. There does not seem to be any major difference in clinical outcome between the two agents, and the choice of tamponade agent can be individualised for each patient. The advantages of silicone oil include its transparency, which allows some vision when walking immediately after surgery. Silicone oil also facilitates postoperative [laser photocoagulation](#), which is more difficult through a gas bubble. Being non-dissolvable, silicone oil also provides long-term tamponade over a large area of the retina. Disadvantages include the need to remove the oil at a second operation to avoid complications. Oil can be left *in situ* to provide continuous retinal tamponade and avoid retinal detachment, but this carries the risk of long-term complications, including glaucoma and keratopathy related to silicone oil emulsification. Newer oil formulations and high viscosity oils carry less risk of this. Leaving oil *in situ* may be necessary in conditions such as CMV-associated retinal detachment with multiple atrophic breaks in areas previously affected by retinitis; cases with persistent unrelieved retinal traction; or cases at high risk of hypotony after oil removal. In these cases, higher viscosity oils are often used to reduce the risk of emulsification and complications.

**OPTION CORTICOSTEROIDS DURING VITRECTOMY SURGERY FOR PROLIFERATIVE VITREO-RETINOPATHY**

- For GRADE evaluation of interventions for Retinal detachment, [see table, p 31](#) .
- We found insufficient evidence assessing the effects of corticosteroids given during vitrectomy surgery for proliferative vitreoretinopathy.

**Benefits and harms**

**Corticosteroids versus no corticosteroids/placebo/standard care:**

We found two RCTs comparing corticosteroid with placebo or standard care. [52] [53]

**Re-attachment rate**

*Corticosteroids compared with placebo or standard treatment* We don't know whether using corticosteroids during vitrectomy surgery improves re-attachment rates in people with proliferative retinopathy ([low-quality evidence](#)).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rate</b>					
[52] RCT	75 people; 75 eyes with rhegmatogenous retinal detachment (RRD) and <a href="#">grade C proliferative vitreoretinopathy (PVR)</a>	<b>Retinal re-attachment rate after 1 operation , 6 months</b> 32/38 (84%) with triamcinolone acetonide 29/37 (78%) with no triamcinolone acetonide	P = 0.5	↔	Not significant

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
		Triamcinolone acetonide was injected into the silicone-filled vitreous cavity on completion of surgery			
[52] RCT	75 people; 75 eyes with RRD and grade C PVR	<b>Final retinal re-attachment rate (with or without re-operation) , 6 months</b> 35/38 (92.1%) with triamcinolone acetonide 34/37 (91.9%) with no triamcinolone acetonide Triamcinolone acetonide was injected into the silicone-filled vitreous cavity on completion of surgery	P = 0.97	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Retinal re-attachment , at end of first operation</b> 81/110 (74%) with oral prednisone tapering for 15 days 72/110 (65%) with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Retinal re-attachment after 1 operation , 24 hours</b> 84/110 (76%) with oral prednisone tapering for 15 days 82/110 (74%) with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Retinal re-attachment after 1 operation , 2 to 4 days</b> 91/110 (82%) with oral prednisone tapering for 15 days 86/110 (78%) with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of	<b>Retinal re-attachment after 1 operation , 4 to 8 days</b> 91/110 (82%) with oral prednisone tapering for 15 days 88/110 (80%) with placebo	Reported as not significant	↔	Not significant

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
	eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic				
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Retinal re-attachment after 1 operation , &gt;8 days</b> 95/110 (86%) with oral prednisone tapering for 15 days 94/110 (85%) with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Re-attachment , at 6 months</b> 97/102 (95%) with oral prednisone tapering for 15 days 84/94 (89%) with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Macular re-attachment , at 6 months</b> 100/102 (98%) with oral prednisone tapering for 15 days 92/94 (98%) with placebo	Reported as not significant	↔	Not significant

**Visual acuity**

*Corticosteroids compared with placebo or standard treatment* We don't know whether using corticosteroids during vitrectomy surgery improves visual acuity in people with proliferative retinopathy ([low-quality evidence](#)).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[52] RCT	75 people; 75 eyes with rhegmatogenous retinal detachment (RRD) and <a href="#">grade C proliferative</a>	<b>Mean best corrected visual acuity (logMAR units)</b> 1.2 with triamcinolone acetate 1.4 with no triamcinolone acetate	P = 0.21	↔	Not significant

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
	tive vitreoretinopathy (PVR)	Triamcinolone acetonide was injected into the silicone-filled vitreous cavity on completion of surgery			
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Visual acuity , at 1 month</b> 0.25 logMAR with oral prednisone tapering for 15 days 0.27 logMAR with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Visual acuity , at 3 months</b> 0.21 logMAR with oral prednisone tapering for 15 days 0.24 logMAR with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Visual acuity , at 6 months</b> 0.20 logMAR with oral prednisone tapering for 15 days 0.21 logMAR with placebo	Reported as not significant	↔	Not significant

### Re-operation rate

*Corticosteroids compared with placebo or standard treatment* We don't know whether using corticosteroids during vitrectomy surgery improves re-operation rates in people with proliferative vitreoretinopathy (*low-quality evidence*).






Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-operation rate</b>					
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of	<b>Re-operations</b> 29 operations in 14/110 eyes (13%) with oral prednisone tapering for 15 days 39 operations in 22/110 eyes (20%) with placebo	Reported as not significant	↔	Not significant

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
	eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic				

No data from the following reference on this outcome. <sup>[52]</sup>

### Adverse effects

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Adverse effects</b>					
<sup>[52]</sup> RCT	75 people; 75 eyes with rhegmatogenous retinal detachment (RRD) and grade C proliferative vitreoretinopathy (PVR)	<b>Intraocular pressure , 6 months</b> 14.7 mmHg with triamcinolone acetonide 16.4 mmHg with no triamcinolone acetonide Triamcinolone acetonide was injected into the silicone-filled vitreous cavity on completion of surgery	P = 0.25	↔	Not significant
<sup>[52]</sup> RCT	75 people; 75 eyes with RRD and grade C PVR	<b>Recurrence of PVR , 6 months</b> 11/38 (29%) with triamcinolone acetonide 11/37 (37%) with no triamcinolone acetonide Triamcinolone acetonide was injected into the silicone-filled vitreous cavity on completion of surgery	P = 0.94	↔	Not significant
<sup>[52]</sup> RCT	75 people; 75 eyes with RRD and grade C PVR	<b>Macular pucker , 6 months</b> 8/38 (21%) with triamcinolone acetonide 13/37 (35%) with no triamcinolone acetonide Triamcinolone acetonide was injected into the silicone-filled vitreous cavity on completion of surgery	P = 0.2	↔	Not significant
<sup>[53]</sup> RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage A , at 1 month</b> 5/106 (5%) with oral prednisone tapering for 15 days 7/99 (7%) with placebo	Reported as not significant	↔	Not significant

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage B , at 1 month</b> 28/106 (26%) with oral prednisone tapering for 15 days 40/99 (40%) with placebo	P <0.05		prednisone
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage C , at 1 month</b> 5/106 (5%) with oral prednisone tapering for 15 days 3/99 (3%) with placebo	Reported as not significant		Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage A , at 3 months</b> 6/107 (6%) with oral prednisone tapering for 15 days 4/99 (4%) with placebo	Reported as not significant		Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage B , at 3 months</b> 27/107 (25%) with oral prednisone tapering for 15 days 45/99 (46%) with placebo	P <0.005		prednisone
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of	<b>Post-operative PVR stage C , at 3 months</b> 3/107 (3%) with oral prednisone tapering for 15 days 2/99 (2%) with placebo	Reported as not significant		Not significant



Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
	eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic				
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage A , at 6 months</b> 1/102 (1%) with oral prednisone tapering for 15 days 4/94 (4%) with placebo	Reported as not significant	↔	Not significant
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage B , at 6 months</b> 23/102 (23%) with oral prednisone tapering for 15 days 43/94 (46%) with placebo	P <0.0005	○○○	prednisone
[53] RCT	220 people with RRD (220 consecutive eyes) having sclera buckling procedure plus air or SF <sub>6</sub> gas in around 50% of eyes plus primary vitrectomy in around 14% of eyes; posterior Nd:Yag capsulotomy in around 9% of eyes; 79.5% phakic	<b>Post-operative PVR stage C , at 6 months</b> 2/102 (2%) with oral prednisone tapering for 15 days 3/94 (3%) with placebo	Reported as not significant	↔	Not significant

#### Further information on studies

#### Comment:

#### Clinical guide:

The antiproliferative and anti-inflammatory properties of corticosteroids are thought to promote repair in rhegmatogenous retinal detachment (RRD). Topical corticosteroids are routinely given by most surgeons post-surgery to correct RRD, and some surgeons give periocular and systemic corticosteroids in cases of RRD with proliferative vitreoretinopathy (PVR). Triamcinolone acetate is a slow-release corticosteroid preparation, and injection into the vitreous cavity provides a locally higher corticosteroid concentration than can be achieved by systemic or topical administration. Although

the RCTs reported here found no benefit associated with using corticosteroids in eyes with RRD and established PVR, [52] corticosteroids may still have a role in other conditions, such as in eyes with inflammation, or after trauma or previous surgery.

**OPTION DAUNORUBICIN DURING VITRECTOMY SURGERY FOR PROLIFERATIVE VITREORETINOPATHY**

- For GRADE evaluation of interventions for Retinal detachment, see table, p 31 .
- We found insufficient evidence assessing the effects of daunorubicin given during vitrectomy surgery for proliferative vitreoretinopathy.

**Benefits and harms**

**Daunorubicin versus no daunorubicin/placebo/standard care:**

We found two RCTs comparing the use of daunorubicin intraoperatively during vitrectomy surgery for rhegmatogenous retinal detachment (RRD) and proliferative vitreoretinopathy (PVR) versus no adjunctive treatment. [54] [55]

**Re-attachment rate**

*Daunorubicin compared with no daunorubicin* Infusing daunorubicin intravitreally for 10 minutes during vitrectomy surgery seems no more effective at improving retinal re-attachment rate (after 1 operation) at 3 to 6 months in patients with rhegmatogenous retinal detachment and grade C and D proliferative vitreoretinopathy (moderate-quality evidence).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rate</b>					
[54] RCT	286 people; 286 eyes with rhegmatogenous retinal detachment (RRD) and proliferative vitreoretinopathy (PVR) of grade C2 or more; multicentre RCT; 24 surgeons	<b>Retinal re-attachment rate (after 1 operation) , 6 months</b> 89/142 (63%) with daunorubicin 73/135 (54%) with no daunorubicin See Further information on studies for details on daunorubicin injections	OR 1.43 95% CI 0.88 to 2.30 P = 0.07	↔	Not significant
[54] RCT	286 people; 286 eyes with RRD and PVR of grade C2 or more; multicentre RCT; 24 surgeons	<b>Overall re-attachment rate (with or without re-operation) , 1 year</b> 105/131 (80%) with daunorubicin 103/126 (82%) with no daunorubicin See Further information on studies for details on daunorubicin injections	Reported as not significant P value not reported	↔	Not significant
[55] RCT	30 people; 30 eyes with RRD and grade D1 PVR or more	<b>Retinal re-attachment , 3 months</b> 13/15 (87%) with daunorubicin 10/15 (67%) with no daunorubicin See Further information on studies for details on daunorubicin injections	Reported as not significant P value not reported	↔	Not significant


**Visual acuity**

*Daunorubicin compared with no daunorubicin* Infusing daunorubicin intravitreally for 10 minutes during vitrectomy surgery seems no more effective at improving visual acuity at 3 to 6 months in people with RRD and grade C and D proliferative vitreoretinopathy (moderate-quality evidence).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[54] RCT	286 people; 286 eyes with rhegmatogenous retinal detachment (RRD) and proliferative vitreoretinopathy (PVR) of grade C2 or more; multicentre RCT; 24 surgeons	<b>Proportion of people with visual acuity rated as improved , 6 months</b> 113/136 (83%) with daunorubicin 98/128 (77%) with no daunorubicin Improved visual acuity defined as positive difference between preoperative logMAR score and score at 6 months See Further information on studies for details on daunorubicin injections	P = 0.17 for between-group comparison (combined analysis of improved, unchanged, and deteriorated)	↔	Not significant
[54] RCT	286 people; 286 eyes with RRD and PVR of grade C2 or more; multicentre RCT; 24 surgeons	<b>Proportion of people with visual acuity rated as unchanged , 6 months</b> 17/136 (13%) with daunorubicin 25/128 (20%) with no daunorubicin Unchanged visual acuity defined as no difference between preoperative logMAR score and score at 6 months See Further information on studies for details on daunorubicin injections	P = 0.17 for between-group comparison (combined analysis of improved, unchanged, and deteriorated)	↔	Not significant
[54] RCT	286 people; 286 eyes with RRD and PVR of grade C2 or more; multicentre RCT; 24 surgeons	<b>Proportion of people with visual acuity rated as deteriorated , 6 months</b> 6/136 (4.4%) with daunorubicin 5/128 (3.9%) with no daunorubicin Deteriorated visual acuity defined as negative difference between preoperative logMAR score and score at 6 months See Further information on studies for details on daunorubicin injections	P = 0.17 for between-group comparison (combined analysis of improved, unchanged, and deteriorated)	↔	Not significant
[55] RCT	30 people; 30 eyes with RRD and grade D1 PVR or more	<b>Proportion of people with improvement in visual acuity , 3 months</b> 14/15 (93%) with daunorubicin 12/15 (80%) with no daunorubicin See Further information on studies for details on daunorubicin injections	Reported as not significant P value not reported	↔	Not significant

**Re-operation rate**

*Daunorubicin compared with no daunorubicin* Infusing daunorubicin intravitreally for 10 minutes during vitrectomy surgery is more effective at reducing the requirement for further vitreoretinal surgery at 1 year in people with rhegmatogenous retinal detachment (RRD) and grade C2 proliferative vitreoretinopathy ([high-quality evidence](#)).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-operation rate</b>					
[54] RCT	286 people; 286 eyes with rhegmatogenous retinal detachment (RRD) and proliferative vitreoretinopathy (PVR) of grade C2 or more; multicentre RCT; 24 surgeons	<b>Proportion of people requiring further vitreoretinal surgery after initial surgery , 1 year</b> 50/145 (34%) with daunorubicin 65/141 (46%) with no daunorubicin  See Further information on studies for details on daunorubicin injections	P = 0.005		daunorubicin

No data from the following reference on this outcome. [55]

### Adverse effects

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Adverse effects</b>					
[54] RCT	286 people; 286 eyes with rhegmatogenous retinal detachment (RRD) and proliferative vitreoretinopathy (PVR) of grade C2 or more; multicentre RCT; 24 surgeons	<b>Adverse effects</b> with daunorubicin with no daunorubicin  The RCT gave no information on adverse effects associated with daunorubicin  The authors reported no treatment-related adverse effects with daunorubicin, and reported no data on other possible adverse effects associated with its use  See Further information on studies for details on daunorubicin injections			
[55] RCT	30 people; 30 eyes with RRD and grade D1 PVR or more	<b>Adverse effects</b> with daunorubicin with no daunorubicin  The RCT gave no information on adverse effects associated with daunorubicin  The authors reported no treatment-related adverse effects with daunorubicin, and reported no data on other possible adverse effects associated with its use  See Further information on studies for details on daunorubicin injections			

### Further information on studies

[54] If the surgeon determined that the retina could be successfully re-attached, the vitreous cavity was perfused with daunorubicin (7.5 micrograms/mL in balanced saline solution) for 10 minutes. Before silicone oil exchange, daunorubicin was exchanged with balanced saline, perfluorocarbon liquid, or air.

[55] Daunorubicin (5 micrograms in 0.1 mL balanced saline solution) was injected into the vitreous cavity and left for 10 minutes, after which time it was flushed out of the vitreous cavity and silicone oil exchange was carried out.

**Comment:** **Clinical guide:** Daunorubicin acts by inhibiting both cell proliferation and cell migration. It can be infused into the vitreous cavity for short periods during vitrectomy surgery without apparent adverse effects. However, it is unclear whether use of daunorubicin as an adjunctive treatment confers benefits in eyes with rhegmatogenous retinal detachment and established proliferative vitreoretinopathy. Further study is required.

**OPTION FLUOROURACIL PLUS HEPARIN DURING VITRECTOMY SURGERY FOR PROLIFERATIVE VITREORETINOPATHY**

- For GRADE evaluation of interventions for Retinal detachment, see table, p 31 .
- We found insufficient evidence assessing the effects of fluorouracil plus heparin given during vitrectomy surgery for proliferative vitreoretinopathy.

**Benefits and harms**

**Fluorouracil plus heparin versus placebo:**

We found one RCT comparing adding perioperative fluorouracil plus low molecular weight heparin to the intraocular infusion versus adding placebo to the intraocular infusion in people having vitrectomy with silicone oil tamponade. [56]

**Re-attachment rate**

*Fluorouracil plus low molecular weight heparin compared with placebo* Adding fluorouracil plus low molecular weight heparin to the intraocular infusion may be no more effective at increasing surgery success rates (re-attaching with removal of silicone oil without further operations and final re-attachment rate) in people with grade C anterior or posterior proliferative vitreoretinopathy (low-quality evidence).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Re-attachment rate</b>					
[56] RCT	157 people with grade C anterior or posterior proliferative vitreoretinopathy (PVR)	<b>Proportion of people with successful surgery , 6 months</b> 39/70 (56%) with fluorouracil plus heparin 40/78 (51%) with placebo Successful surgery defined as re-attachment with removal of silicone oil without further operations	P = 0.589	↔	Not significant
[56] RCT	157 people with grade C anterior or posterior PVR	<b>Overall complete retinal re-attachment rate with or without re-operation , 6 months</b> 56/67 (83.5%) with fluorouracil plus heparin 65/77 (84.4%) with placebo	Reported as not significant P value not reported	↔	Not significant

**Visual acuity**

*Fluorouracil plus low molecular weight heparin compared with placebo* Adding fluorouracil plus low molecular weight heparin to the intraocular infusion seems no more effective at improving visual acuity in people with grade C anterior or posterior proliferative vitreoretinopathy (moderate-quality evidence).

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Visual acuity</b>					
[56] RCT	157 people with grade C anterior or posterior proliferative vitreoretinopathy (PVR)	<b>Mean visual acuity on logMAR scale</b> 1.8 with fluorouracil plus heparin 1.4 with placebo	P = 0.126	↔	Not significant

### Re-operation rate

No data from the following reference on this outcome. [56]

### Adverse effects

Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
<b>Adverse effects</b>					
[56] RCT	157 people with grade C anterior or posterior proliferative vitreoretinopathy (PVR)	<b>Macular pucker , 6 months</b> 4/66 (6%) with fluorouracil plus heparin 13/77 (17%) with placebo	P = 0.068	↔	Not significant
[56] RCT	157 people with grade C anterior or posterior PVR	<b>Glaucoma , 12 months</b> 0 with fluorouracil plus heparin 3 with placebo Adverse effects data at 12 months available for 98/157 (62%) participants Unclear whether figures represent percentages or absolute number of people with adverse effect in each group	Reported as not significant P value not reported	↔	Not significant
[56] RCT	157 people with grade C anterior or posterior PVR	<b>Hypotony , 12 months</b> 9 with fluorouracil plus heparin 7 with placebo Adverse effects data at 12 months available for 98/157 (62%) participants Unclear whether figures represent percentages or absolute number of people with adverse effect in each group	Reported as not significant P value not reported	↔	Not significant
[56] RCT	157 people with grade C anterior or posterior PVR	<b>Keratopathy , 12 months</b> 5 with fluorouracil plus heparin 2 with placebo Adverse effects data at 12 months available for 98/157 (62%) participants Unclear whether figures represent percentages or absolute number of people with adverse effect in each group	Reported as not significant P value not reported	↔	Not significant



Ref (type)	Population	Outcome, Interventions	Results and statistical analysis	Effect size	Favours
[56] RCT	157 people with grade C anterior or posterior PVR	<b>Need for cataract extraction , 12 months</b> 21 with fluorouracil plus heparin 29 with placebo  Adverse effects data at 12 months available for 98/157 (62%) participants  Unclear whether figures represent percentages or absolute number of people with adverse effect in each group	Reported as not significant P value not reported	↔	Not significant

#### Further information on studies

#### Comment:

#### Clinical guide:

Despite evidence suggesting that 5-fluorouracil plus heparin can prevent proliferative vitreoretinopathy (PVR) in people with high-risk features for PVR undergoing vitrectomy surgery for rhegmatogenous retinal detachment,<sup>[57]</sup> it is unclear whether fluorouracil plus heparin is effective at improving retinal re-attachment rates in people with established PVR. The prevention of PVR will be addressed in full in future updates of this review.

## GLOSSARY

**Classification of PVR** Grade A PVR denotes vitreous haze and pigment clumping (of retinal pigment epithelium cells) in the vitreous cavity (although this grade is rarely used). Grade B PVR shows areas of surface retinal wrinkling with rolled edges to retinal tears. Grade C PVR consists of fixed full thickness retinal folds involving 1 to 3 quadrants: Grade C1–C3. Grade D was classified as a total RRD with either a wide (D1), narrow (D2), or closed (D3) funnel configuration because of fixed retinal folds.<sup>[35]</sup> Grade D was removed from the 1991 classification update,<sup>[36]</sup> and Grade C was divided into anterior and posterior PVR, which is then subdivided based on the number of hours involved (CA1–12 and CP1–12), and on type of fibrosis and contracture present (focal, diffuse, or subretinal, and anterior, circumferential, and/or anterior displacement [anterior loop traction]).

**Aphakic** An aphakic eye has neither a natural crystalline lens nor an artificial lens.

**Macular pucker** refers to the distorted anatomical appearance of the macular retina caused by localised epiretinal fibrotic membrane formation. It can result in distorted and reduced central vision.

**Phakic** A phakic eye has an intact natural crystalline lens.

**Pneumatic retinopexy** A small volume of gas, primarily expansile gas, is injected into the vitreous cavity and used to close the retinal break(s). No attempt is made to relieve vitreoretinal traction. Once closure of retinal breaks is achieved, the physiological retinal pigment epithelium pump removes subretinal fluid resulting in retinal reattachment. Before or after gas injection, laser or cryotherapy is usually applied to the retinal breaks (retinopexy) to create a permanent choroidoretinal adhesion.

**Proliferative vitreoretinopathy (PVR)** after a retinal detachment may occur either spontaneously before surgery or after treatment. PVR refers to the growth of avascular fibrocellular membranes within the vitreous cavity and on the front and back surfaces of the retina. These membranes, which are essentially scar tissues, occur in the mildest form as fine fibrous membranes on the retinal surface without visible retinal distortion or merely rolling of the edges of retinal breaks. In more severe forms, the membranes cause fixed retinal folds, preventing closure of retinal breaks and exerting traction on the retina. Retinal folds may also result in recurrence of retinal detachment, even after an initially successful retinal detachment procedure, because of spontaneous reopening of otherwise successfully treated retinal breaks, or because of the development of new retinal breaks. Epiretinal membranes on the surface of the macula causing macular pucker and ocular hypotony secondary to PVR involving the ciliary body may also occur. PVR may result in disappointing visual results.<sup>[59]</sup>

**Pseudophakic** A pseudophakic eye has had the natural lens removed and replaced with an artificial intraocular lens implant.

**Retinal dialysis** is a separation of the retina where it inserts into the pars plana at the ora serrata.

**Retinal operculum** This is a separated flap of retina avulsed from the retinal surface by vitreoretinal separation, leaving a retinal hole.

**Retinal-flap tear** This is a tear in the retina associated with local vitreoretinal traction, separation, or both; the flap of the tear remains attached to the vitreous and connected by its base to the anterior edge of the retinal tear.

**Scleral buckling surgery** A buckling element or explant, usually made of either solid silicone or silicone sponge, is sutured to the sclera externally to indent the sclera and underlying retinal pigment epithelium towards the detached retina at the site of the retinal break(s), to close the break and relieve vitreoretinal traction. Buckles can be either segmental or encircling. Once closure of retinal breaks is achieved, the physiological retinal pigment epithelium pump removes subretinal fluid resulting in retinal reattachment. This process can be assisted by subretinal fluid drainage at the time of surgery, which also allows break closure if subretinal fluid is deep. During surgery, laser or cryotherapy is usually applied to the retinal breaks (retinopexy) to create a permanent choroidoretinal adhesion.<sup>[60]</sup>

**Silicone oil tamponade** is used in vitrectomy as an alternative to gas. Silicone oil is also now available in a heavier-than-water preparation, allowing inferior retinal tamponade without head-down posturing.

**logMAR chart** A tool for measuring visual acuity, similar to but more precise than a Snellen chart. The chart is typically read at 4 m and scored from the total number of letters read. A score of 1.0 is equivalent to Snellen acuity 6/60 and indicates that all 5 letters on the top line, but no others, were read. A score of 0.1 is equivalent to Snellen acuity 6/6.

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

**Laser photocoagulation** Refers to the transpupillary application of laser (usually argon laser) to retinal breaks or predisposing rhegmatogenous retinal detachment lesions. It can be delivered either by a slit lamp-mounted laser system or by using a laser connected to an indirect ophthalmoscope. Contiguous laser burns are placed around the lesion in 2 to 3 rows leading to areas of full-thickness chorioretinal adhesion within 2 to 3 days of treatment. Laser photocoagulation can be carried out under local anaesthetic. Because it is delivered through the pupil, posterior retinal lesions can be treated without the need to open the conjunctiva.

**Low-quality evidence** 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.

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

**Posterior vitreous detachment (PVD)** is the separation of the vitreous gel from its posterior attachment to the retina. PVD is associated with ageing of the vitreous characterised by liquefaction of the vitreous gel itself. Liquefaction occurs at an earlier age in myopic eyes than in emmetropic and hypermetropic eyes, and can be accelerated by inflammation caused by surgery, trauma, or uveitis. Vitreous liquefaction leads to vitreous gel instability, which triggers PVD. PVD is present in autopsy studies in less than 10% of people aged under 50 years, in at least one eye in 27% of people aged 60 to 65 years, and in 63% of people aged over 70 years. It usually occurs as an acute event with rapid evolution of vitreoretinal separation from the posterior to anterior retina.<sup>[58]</sup>

**Snellen visual acuity** The Snellen chart usually includes letters, numbers, or pictures printed in lines of decreasing size, which are read or identified from a fixed distance; distance visual acuity is usually measured from a distance of 6 m (20 feet). The Snellen visual acuity is written as a fraction: 6/18 means that from 6 m away the best line that can be read is a line that could normally be read from a distance of 18 m away.

**Stickler's syndrome (hereditary arthro-ophthalmopathy)** is a hereditary disease of type 2 collagen resulting in abnormal vitreous, myopia, and a variable degree of orofacial abnormalities, deafness, and arthropathies.

**Very low-quality evidence** Any estimate of effect is very uncertain.

**Visual acuity testing** This is carried out with charts using letters or standard pictures or symbols. Modern tests that incorporate crowding and logMAR (logarithm of the minimum angle of resolution) size scaling are more accurate. One line of letters or symbols (usually 4 or 5) constitutes 0.1 logMAR units and roughly approximates to one line on a Snellen chart, although this conversion factor is inaccurate and should only be used as a crude guide to interpretation. Given the variability in test performance within individuals, a change in 0.2 logMAR units is often quoted as being the smallest clinically important change, although some studies use a change of 0.1 logMAR or greater, which might be considered clinically more marginal. Change of less than 0.1 logMAR unit is not clinically important and could be accounted for by test–retest variability.

**Vitrectomy** The vitreous is removed internally using a cutting aspirating instrument relieving vitreoretinal traction directly. A tamponade agent, usually gas or silicone oil, is used to close the break(s). Closure is assisted by postoperative positioning to place the tamponade bubble against the break(s) in an optimum way. Gases can be short- ( $\text{SF}_6$ ), medium- ( $\text{C}_2\text{F}_6$ ), or long-acting ( $\text{C}_3\text{F}_8$ ), and last a variable period of time depending on concentration and gas fill before being absorbed. Once closure of retinal breaks is achieved, the physiological retinal pigment epithelium pump removes subretinal fluid resulting in retinal reattachment. This process can be assisted by subretinal fluid drainage at the time of surgery. During surgery, laser or cryotherapy is usually applied to the retinal breaks (retinopexy) to create a permanent choroidoretinal adhesion.<sup>[61]</sup> Vitrectomy surgery can be carried out with different gauge (di-

ameter) instruments ranging from 27 gauge to 20 gauge. Twenty-gauge vitrectomy is often referred to as a standard vitrectomy and 23, 25, and 27 gauge as narrow-gauge. Many narrow-gauge surgeries are done through transconjunctival incisions without prior conjunctival opening and without suturing. Proposed advantages of narrow-gauge surgery include faster postoperative recovery and less postoperative pain and inflammation, with more precise surgical manoeuvres. Disadvantages include more flexible instrumentation and the risk of postoperative hypotony with wound leak. Typically surgeons have a preference for one gauge over others, and also availability of equipment can limit gauge choice. There is no evidence as yet that one gauge has a higher success rate than any other for RRD.

## SUBSTANTIVE CHANGES

**Corticosteroids during vitrectomy surgery for proliferative vitreoretinopathy:** one RCT added. <sup>[53]</sup> Categorisation unchanged (unknown effectiveness).

**Scleral buckling versus primary vitrectomy:** two systematic reviews added. <sup>[40] [41]</sup> Categorisation unchanged (unknown effectiveness).

## REFERENCES

- Yanoff M, Duker JJ. *Ophthalmology*. 2nd ed. St Louis, MO: Mosby, 2004: 982–989.
- Kanski JK. *Clinical ophthalmology*. Edinburgh, UK: Butterworth-Heinemann, 2003: 349–388.
- Yanoff M, Duker JJ. *Ophthalmology*. 2nd ed. St Louis, MO: Mosby, 2004: 990.
- Tanner V, Harle D, Tan J, et al. Acute posterior vitreous detachment: the predictive value of vitreous pigment and symptomatology. *Br J Ophthalmol* 2000;84:1264–1268. [\[PubMed\]](#)
- Rowe JA, Erie JC, Baratz KH, et al. Retinal detachment in Olmsted County, Minnesota, 1976 through 1995. *Ophthalmology* 1999;106:154–159. [\[PubMed\]](#)
- Laatikainen L, Tolppanen EM, Harju H. Epidemiology of rhegmatogenous retinal detachment in a Finnish population. *Acta Ophthalmol (Copenh)* 1985;63:59–64. [\[PubMed\]](#)
- Törnquist R, Stenkula S, Törnquist P. Retinal detachment. A study of a population-based patient material in Sweden 1971–1981. I. Epidemiology. *Acta Ophthalmol (Copenh)* 1987;65:213–222. [\[PubMed\]](#)
- Haimann MH, Burton TC, Brown CK. Epidemiology of retinal detachment. *Arch Ophthalmol* 1982;100:289–292. [\[PubMed\]](#)
- Haut J, Massin M. Frequency of incidence of retina detachment in the French population. Percentage of bilateral detachment. *Arch Ophthalmol Rev Gen Ophthalmol* 1975;35:533–536. [In French] [\[PubMed\]](#)
- Wilkes SR, Beard CM, Kurland LT, et al. The incidence of retinal detachment in Rochester, Minnesota, 1970–1978. *Am J Ophthalmol* 1982;94:670–673. [\[PubMed\]](#)
- Polkinghorne PJ, Craig JP. Northern New Zealand Rhegmatogenous Retinal Detachment Study: epidemiology and risk factors. *Clin Experiment Ophthalmol* 2004;32:159–163. [\[PubMed\]](#)
- Wilkinson CP, Rice TA. *Michel's retinal detachment*. 2nd ed. St Louis, MO: Mosby, 1997: 29.
- Wilkinson CP, Rice TA. *Michel's retinal detachment*. 2nd ed. St Louis, MO: Mosby, 1997: 93.
- Wilkinson CP. Evidence-based analysis of prophylactic treatment of asymptomatic retinal breaks and lattice degeneration. *Ophthalmology* 2000;107:12–16. [\[PubMed\]](#)
- Wilkinson CP, Rice TA. *Michel's retinal detachment*. 2nd ed. St Louis, MO: Mosby, 1997: 49.
- Shea M, Davis MD, Kamel I. Retinal breaks without detachment, treated and untreated. *Mod Probl Ophthalmol* 1974;12:97–102. [\[PubMed\]](#)
- Foos RY, Allen RA. Retinal tears and lesser lesions of the peripheral retina in autopsy eyes. *Am J Ophthalmol* 1967;64:643–655. [\[PubMed\]](#)
- Byer NE. What happens to untreated asymptomatic retinal breaks, and are they affected by posterior vitreous detachment? *Ophthalmology* 1998;105:1045–1049. [\[PubMed\]](#)
- Burton TC. The influence of refractive error and lattice degeneration on the incidence of retinal detachment. *Trans Am Ophthalmol Soc* 1989;87:143–155. [\[PubMed\]](#)
- Byer NE. Lattice degeneration of the retina. *Surv Ophthalmol* 1979;23:213–248. [\[PubMed\]](#)
- Okun E. Gross and microscopic pathology in autopsy eyes. III. Retinal breaks without detachment. *Am J Ophthalmol* 1961;51:369–391. [\[PubMed\]](#)
- Vote BJ, Casswell AG. Retinal dialysis: are we missing diagnostic opportunities? *Eye* 2004;18:709–713. [\[PubMed\]](#)
- Johnston PB. Traumatic retinal detachment. *Br J Ophthalmol* 1991;75:18–21. [\[PubMed\]](#)
- Ashrafzadeh MT, Schepens CL, Elzeneiny II, et al. Aphakic and phakic retinal detachment. I. Preoperative findings. *Arch Ophthalmol* 1973;89:476–483. [\[PubMed\]](#)
- Eye Disease Case Control Study Group. Risk factors for idiopathic rhegmatogenous retinal detachment. *Am J Epidemiol* 1993;137:749–757. [\[PubMed\]](#)
- Mitry D, Charteris DG, Yorston D, et al; Scottish RD Study Group. The epidemiology and socioeconomic associations of retinal detachment in Scotland: a two-year prospective population-based study. *Invest Ophthalmol Vis Sci* 2010;51:4963–4968. [\[PubMed\]](#)
- Byer NE. Rethinking prophylactic treatment of retinal detachment. In: Stirpe M, ed. *Advances in vitreoretinal surgery*. Acta Third International Congress on Vitreoretinal Surgery. Rome, 12–14 September 1991. New York, NY: Ophthalmic Communications Society, 1992: 399–411.
- Davis MD. The natural history of retinal breaks without detachment. *Trans Am Ophthalmol Soc* 1973;71:343–372. [\[PubMed\]](#)
- Dralands L, Larminier F, Cornelis H, et al. Evolution of lesions of the retinal periphery in the fellow eye of a retinal detachment. *Bull Mem Soc Fr Ophthalmol* 1981;92:73–77. [In French] [\[PubMed\]](#)
- Hovland KR. Vitreous findings in fellow eyes of aphakic retinal detachment. *Am J Ophthalmol* 1978;86:350–353. [\[PubMed\]](#)
- Folk JC, Arrindell EL, Klugman MR. The fellow eye of patients with phakic lattice retinal detachment. *Ophthalmology* 1989;96:72–79. [\[PubMed\]](#)
- Bhagwandien AC, Cheng YY, Wolfs RC, et al. Relationship between retinal detachment and biometry in 4262 cataractous eyes. *Ophthalmology* 2006;113:643–649. [\[PubMed\]](#)
- Tuft SJ, Minassian D, Sullivan P. Risk factors for retinal detachment after cataract surgery: a case-control study. *Ophthalmology* 2006;113:650–656. [\[PubMed\]](#)
- Wilkinson CP, Rice TA. *Michel's retinal detachment*. 2nd ed. St Louis, MO: Mosby, 1997: 935–977.
- Retina Society Terminology Committee. The classification of retinal detachment with proliferative vitreoretinopathy. *Ophthalmology* 1983;90:121–125. [\[PubMed\]](#)
- Machemer R, Aaberg TM, Freeman HM, et al. An updated classification of retinal detachment with proliferative vitreoretinopathy. *Am J Ophthalmol* 1991;112:159–165. [\[PubMed\]](#)
- Wilkinson CP, Rice TA. *Michel's retinal detachment*. 2nd ed. St Louis, MO: Mosby, 1997: 960–961.
- Tornambe PE, Hilton GF. Pneumatic retinopathy. A multicenter randomized controlled clinical trial comparing pneumatic retinopathy with scleral buckling. The Retinal Detachment Study Group. *Ophthalmology* 1989;96:772–783. [\[PubMed\]](#)
- Mulvihill A, Fulcher T, Datta V, et al. Pneumatic retinopathy versus scleral buckling: a randomised controlled trial. *Ir J Med Sci* 1996;165:274–277. [\[PubMed\]](#)
- Soni C, Hainsworth DP, Almony A. Surgical management of rhegmatogenous retinal detachment: a meta-analysis of randomized controlled trials. *Ophthalmology* 2013;120:1440–1447. [\[PubMed\]](#)
- Sun Q, Sun T, Xu Y, et al. Primary vitrectomy versus scleral buckling for the treatment of rhegmatogenous retinal detachment: a meta-analysis of randomized controlled clinical trials. *Curr Eye Res* 2012;37:492–499. [\[PubMed\]](#)
- Brazitikos PD, Androudi S, Christen WG, et al. Primary pars plana vitrectomy versus scleral buckle surgery for the treatment of pseudophakic retinal detachment: a randomized clinical trial. *Retina* 2005;25:957–964. [\[PubMed\]](#)
- Sharma YR, Karunaniithi S, Azad RV, et al. Functional and anatomic outcome of scleral buckling versus primary vitrectomy in pseudophakic retinal detachment. *Acta Ophthalmol Scand* 2005;83:293–297. [\[PubMed\]](#)
- Ahmadiéh H, Moradian S, Faghihi H, et al. Anatomic and visual outcomes of scleral buckling versus primary vitrectomy in pseudophakic and aphakic retinal detachment: six-month follow-up results of a single operation – report no. 1. *Ophthalmology* 2005;112:1421–1429. [\[PubMed\]](#)
- Heimann H, Bartz-Schmidt KU, Bornfeld N, et al. Scleral buckling versus primary vitrectomy in rhegmatogenous retinal detachment: a prospective randomized multicenter clinical study. *Ophthalmology* 2007;114:2142–2154. [\[PubMed\]](#)
- Koriyama M, Nishimura T, Matsubara T, et al. Prospective study comparing the effectiveness of scleral buckling to vitreous surgery for rhegmatogenous retinal detachment. *Jpn J Ophthalmol* 2007;51:360–367. [\[PubMed\]](#)
- Schwartz SG, Flynn HW Jr, Lee WH, et al. Tamponade in surgery for retinal detachment associated with proliferative vitreoretinopathy. In: *The Cochrane Library*, Issue 3, 2013. Chichester, UK: John Wiley & Sons, Ltd. Search date 2009.
- McCuen B, Azen SP, Boone DC, et al; Silicone Study Group. Vitrectomy with silicone oil or perfluoropropane gas in eyes with severe proliferative vitreoretinopathy: results of a randomized controlled trial. *Silicone Study Report 2. Arch Ophthalmol* 1992;110:780–792. [\[PubMed\]](#)
- Barr CC, Lai MY, Lean JS, et al. Postoperative intraocular pressure abnormalities in the Silicone Study. *Silicone Study Report 4. Ophthalmology* 1993;100:1629–1635. [\[PubMed\]](#)
- Cox MS, Azen SP, Barr CC, et al. Macular pucker after successful surgery for proliferative vitreoretinopathy. *Silicone Study Report 8. Ophthalmology* 1995;102:1884–1891. [\[PubMed\]](#)
- Lean JS, Boone DC, Azen SP, et al; Silicone Study Group. Vitrectomy with silicone oil or sulfur hexafluoride gas in eyes with severe proliferative vitreoretinopathy: results of a randomized controlled trial. *Silicone Study Report 1. Arch Ophthalmol* 1992;110:770–779. [\[PubMed\]](#)
- Ahmadiéh H, Feghhi M, Tabatabaei H, et al. Triamcinolone acetonide in silicone-filled eyes as adjunctive treatment for proliferative vitreoretinopathy: a randomized clinical trial. *Ophthalmology* 2008;115:1938–1943. [\[PubMed\]](#)

53. Koerner F, Koerner-Stiefbold U, Garweg JG. Systemic corticosteroids reduce the risk of cellophane membranes after retinal detachment surgery: a prospective randomized placebo-controlled double-blind clinical trial. *Graefes Arch Clin Exp Ophthalmol* 2012;250:981–987.[\[PubMed\]](#)
54. Wiedemann P, Hilgers RD, Bauer P, et al. Adjunctive daunorubicin in the treatment of proliferative vitreoretinopathy: results of a multicenter clinical trial. Daunomycin Study Group. *Am J Ophthalmol* 1998;126:550–559.[\[PubMed\]](#)
55. Kumar A, Nainiwal S, Choudhary I, et al. Role of daunorubicin in inhibiting proliferative vitreoretinopathy after retinal detachment surgery. *Clin Experiment Ophthalmol* 2002;30:348–351.[\[PubMed\]](#)
56. Charteris DG, Aylward GW, Wong D, et al. A randomized controlled trial of combined 5-fluorouracil and low-molecular-weight heparin in management of established proliferative vitreoretinopathy. *Ophthalmology* 2004;111:2240–2245.[\[PubMed\]](#)
57. Asaria RH, Kon CH, Bunce C, et al. Adjuvant 5-fluorouracil and heparin prevents proliferative vitreoretinopathy: results from a randomized, double-blind controlled clinical trial. *Ophthalmology* 2001;108:1179–1183.[\[PubMed\]](#)
58. Wilkinson CP, Rice TA. Michel's retinal detachment. St Louis, MO: Mosby, 1997: 29–33.
59. Yanoff M, Duker JJ. *Ophthalmology*. 2nd ed. St Louis, MO: Mosby, 2004: 1002–1003.
60. Yanoff M, Duker JJ. *Ophthalmology*. 2nd ed. St Louis, MO: Mosby, 2004: 787.
61. Kanski JK. *Clinical ophthalmology*. Edinburgh, UK: Butterworth-Heinemann, 2003: 383–384.

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**GRADE** Evaluation of interventions for Retinal detachment.

Important outcomes	Studies (Participants)	Outcome	Comparison	Re-attachment rate, Re-operation rate, Visual acuity					GRADE	Comment
				Type of evidence	Quality	Consistency	Directness	Effect size		
<i>What are the effects of different surgical interventions in people with rhegmatogenous retinal detachment?</i>										
	2 (218) <sup>[38]</sup> <sup>[39]</sup>	Re-attachment rate	Scleral buckling versus pneumatic retinopexy	4	-1	0	0	0	Moderate	Quality point deducted for incomplete reporting
	2 (218) <sup>[38]</sup> <sup>[39]</sup>	Visual acuity	Scleral buckling versus pneumatic retinopexy	4	-1	0	0	0	Moderate	Quality point deducted for incomplete reporting
	at least 5 (at least 780) <sup>[40]</sup>	Re-attachment rate	Scleral buckling versus primary vitrectomy in pseudophakic or aphakic rhegmatogenous retinal detachment (RRD)	4	-1	-1	0	0	Low	Quality point deducted for significant heterogeneity between trials; consistency point deducted for conflicting results
	7 (867) <sup>[40]</sup> <sup>[41]</sup>	Visual acuity	Scleral buckling versus primary vitrectomy in pseudophakic or aphakic rhegmatogenous retinal detachment (RRD)	4	-1	0	-1	0	Low	Quality point deducted for incomplete reporting of results; directness point deducted for inclusion of co-intervention
	1 (150) <sup>[42]</sup>	Re-operation rate	Scleral buckling versus primary vitrectomy in pseudophakic or aphakic rhegmatogenous retinal detachment (RRD)	4	-1	0	0	0	Moderate	Quality point deducted for sparse data
	3 (523) <sup>[40]</sup>	Re-attachment rate	Scleral buckling versus primary vitrectomy in phakic rhegmatogenous retinal detachment (RRD)	4	0	0	-2	0	Low	Directness points deducted for unclear outcome in one RCT and inclusion of co-intervention in one RCT
	at least 3 (at least 477) <sup>[40]</sup> <sup>[41]</sup>	Visual acuity	Scleral buckling versus primary vitrectomy in phakic rhegmatogenous retinal detachment (RRD)	4	-1	0	-2	0	Very low	Quality point deducted for incomplete reporting of results; directness points deducted for unclear clinical relevance and because some people had secondary procedures
<i>What are the effects of interventions to treat rhegmatogenous retinal detachment associated with proliferative vitreoretinopathy?</i>										
	1 (265 eyes) <sup>[48]</sup>	Re-attachment rate	Silicone oil tamponade versus long-acting gas tamponade	4	-1	0	0	0	Moderate	Quality point deducted for methodological issues (no statistical assessment for one comparison and poor follow-up at 36 months)
	1 (265 eyes) <sup>[48]</sup>	Visual acuity	Silicone oil tamponade versus long-acting gas tamponade	4	0	0	0	0	High	
	1 (97 eyes) <sup>[51]</sup>	Re-attachment rate	Silicone oil tamponade versus short-acting gas tamponade	4	-1	0	0	0	Moderate	Quality point deducted for sparse data
	1 (97 eyes) <sup>[51]</sup>	Visual acuity	Silicone oil tamponade versus short-acting gas tamponade	4	-1	0	0	0	Moderate	Quality point deducted for sparse data
	2 (295) <sup>[52]</sup> <sup>[53]</sup>	Re-attachment rate	Corticosteroids versus no corticosteroids/placebo/standard care	4	-1	0	-1	0	Low	Quality point deducted for incomplete reporting of results; directness point deducted for the small number of comparators

Important outcomes	Re-attachment rate, Re-operation rate, Visual acuity								
	Studies (Participants)	Outcome	Comparison	Type of evidence	Quality	Consistency	Directness	Effect size	GRADE
2 (295) <sup>[52]</sup> <sup>[53]</sup>	Visual acuity	Corticosteroids versus no corticosteroids/placebo/standard care	4	-1	0	-1	0	Low	Quality point deducted for incomplete reporting of results; directness point deducted for the small number of comparators
1 (220) <sup>[53]</sup>	Re-operation rate	Corticosteroids versus no corticosteroids/placebo/standard care	4	-1	0	-1	0	Low	Quality point deducted for incomplete reporting of results; directness point deducted for the small number of comparators
2 (307) <sup>[54]</sup> <sup>[55]</sup>	Re-attachment rate	Daunorubicin versus no daunorubicin/placebo/standard care	4	-1	0	0	0	Moderate	Quality point deducted for incomplete reporting
2 (294) <sup>[54]</sup> <sup>[55]</sup>	Visual acuity	Daunorubicin versus no daunorubicin/placebo/standard care	4	-1	0	0	0	Moderate	Quality point deducted for incomplete reporting
1 (286) <sup>[54]</sup>	Re-operation rate	Daunorubicin versus no daunorubicin/placebo/standard care	4	0	0	0	0	High	
1 (148) <sup>[56]</sup>	Re-attachment rate	Fluorouracil plus heparin versus placebo	4	-2	0	0	0	Low	Quality points deducted for sparse data and incomplete reporting
1 (157) <sup>[56]</sup>	Visual acuity	Fluorouracil plus heparin versus placebo	4	-1	0	0	0	Moderate	Quality point deducted for sparse data

We initially allocate 4 points to evidence from RCTs, and 2 points to evidence from observational studies. To attain the final GRADE score for a given comparison, points are deducted or added from this initial score based on preset criteria relating to the categories of quality, directness, consistency, and effect size. Quality: based on issues affecting methodological rigour (e.g., incomplete reporting of results, quasi-randomisation, sparse data [ $<200$  people in the analysis]). Consistency: based on similarity of results across studies. Directness: based on generalisability of population or outcomes. Effect size: based on magnitude of effect as measured by statistics such as relative risk, odds ratio, or hazard ratio.