A systematic review and meta-analysis of clinical outcomes of vitrectomy with or without intravitreal bevacizumab pretreatment for severe diabetic retinopathy

Li-Quan Zhao, Huang Zhu, Pei-Quan Zhao, Yi-Qian Hu

ABSTRACT

Aims To examine possible benefits of intravitreal bevacizumab (IVB) pretreatment in vitrectomy for severe diabetic retinopathy.

Methods A comprehensive literature search was performed using the Cochrane Collaboration methodology to identify randomised controlled trials and comparative studies of vitrectomy with or without IVB pretreatment for severe or complicated diabetic retinopathy. Meta-analyses were performed for intraoperative (including intraoperative bleeding, endodiathermy, iatrogenic retinal tears and mean surgical time) and postoperative outcome parameters (including best-corrected visual acuity, recurrent vitreous haemorrhage, reabsorption time of blood and other complications).

Results Six randomised controlled trials and one comparative study were identified and used for comparing vitrectomy alone (142 eyes, control group) with vitrectomy with IVB pretreatment (139 eyes). The intraoperative findings showed that the incidence of intraoperative bleeding and frequency of endodiathermy were statistically significantly less in the IVB pretreatment group (p < 0.01) than in the vitrectomy alone group. The IVB pretreatment group took significantly less surgical time than the control group (p = 0.003). Postoperative results indicated that reabsorption time of blood was significantly shorter (p = 0.04), incidence of recurrent VH was almost significantly less (p = 0.05), and final best-corrected visual acuity was significantly better (p = 0.003) in the IVB group than in the control group. Other complications, including final retinal detachment, and reoperation, were statistically insignificant.

Conclusion IVB pretreatment in vitrectomy can achieve excellent clinical outcomes for severe diabetic retinopathy. It potentially facilitates surgeons’ manoeuvres and reduces intra- and postoperative complications.

INTRODUCTION

Recently, numerous studies have reported clinical outcomes of intravitreal bevacizumab (IVB) as an adjunct to vitrectomy in the management of diabetic retinopathy.1–10 Bevacizumab can induce regression of retinal neovascularisation in patients with diabetes; therefore, it was suggested that a presurgical administration of IVB may reduce intraoperative bleeding during vitrectomy in proliferative diabetic retinopathy (PDR).1 3–6 However, the presurgical administration of IVB remains controversial. Some studies reported that bevacizumab pretreatment for diabetic vitrectomy did not influence rates of postoperative vitreous haemorrhage or final visual acuity.7 10

Although many surgeons perform IVB before vitrectomy in patients with diabetes, systematic or larger sample size studies demonstrating its benefits in facilitation of surgery and clinical outcomes are limited. Therefore, it is necessary to review in greater depth the available studies to understand the benefits of IVB pretreatment. In an attempt to detect benefits in safety and efficacy as the primary comparative criteria, we performed a system review and meta-analysis of existing RCTs and high-quality comparative studies of vitrectomy with or without IVB pretreatment for the treatment of severe or complicated diabetic retinopathy.

MATERIALS AND METHODS

This meta-analysis was performed according to a predetermined protocol described previously.11 12

Literature search

Two reviewers independently searched the following electronic databases: PubMed, EMBASE and the Cochrane Controlled Trials Register up to 30 April 2010. For maximum sensitivity, we used free text and thesaurus terms including ‘vitrectomy,’ ‘diabetic retinopathy’ and ‘bevacizumab.’ Full articles were retrieved, when titles and/or abstracts met this objective. A manual cross-reference search of the bibliographies of relevant articles was conducted. All published studies comparing vitrectomy alone versus vitrectomy with IVB pretreatment for diabetic retinopathy were included, if they met the inclusion criteria. The search included RCTs and high-quality comparative studies. For an inclusion in the meta-analysis, the patients in the selected studies had to present with severe diabetic retinopathy and be older than 18 years of age. At least one or more clinical outcomes representing intraoperative and/or postoperative outcome parameters must be assessed and published. There was no language restriction on the publications.

Discordance about study inclusion between the two reviewers was resolved through discussion until 100% agreement was reached on the final interpretation of the data.

Quality assessment of retrieved articles

The selected studies were appraised by two reviewers, who independently assessed their quality
using the methods recommended in the *Cochrane Handbook for Systematic Reviews of Interventions*. Methodological quality included allocation concealment, method of allocation to treatment, masking of outcome assessment and completeness of follow-up.

**Outcome measure**

The intraoperative outcome parameters included incidence of intraoperative bleeding, frequency of endodiathermy, iatrogenic retinal tears and mean surgical time. Postoperative outcome parameters included best-corrected visual acuity (BCVA), recurrent vitreous haemorrhage (VH), reabsorption time of blood, early and late elevation of intraocular pressure (IOP), final retinal detachment (RD) and repeat vitrectomy.

**Data extraction and analysis**

The studies were tabulated and methodologically evaluated to assess homogeneity. Any heterogeneity between the studies would not be justified to pool the assessed outcomes. A customised data-extraction form, as described in the *Cochrane Handbook for Systematic Reviews of Interventions*, was used to record the duration of the trial, sample size, dropouts, the system and ocular baseline features, the type of surgery and differential treatment, the dosage and time of bevacizumab injection.

**Statistical analysis**

Quantitative data were analysed using the Cochrane Review Manager (RevMan) version 4.2 software. Summary estimates, including 95% CIs, were calculated. For continuous outcome data (eg, mean surgical time), means and standard deviations were used to calculate a weighted mean difference (WMD). For dichotomous outcomes (eg, proportions of intraoperative bleeding), the OR was calculated.

Statistical heterogeneity was tested using the Q statistic of the \( \chi^2 \) value test and I\(^2\) test. Fixed-effects models were used, unless significant evidence of statistical heterogeneity or clinical diversity was found. For results showing significant heterogeneity (I\(^2\)>50%), a random-effects meta-analysis was performed by the DerSimonian–Laird method. Outcome measures were assessed on an intent-to-treat basis. A p value of <0.05 was considered statistically significant. A sensitivity analysis was performed by excluding the non-randomised study.

Publication bias was assessed by visually inspecting a funnel plot.

**RESULTS**

Six RCTs and one comparative study published between 2008 and 2010 met the inclusion criteria. In di Lauro’s study, the group of IVB 7 days prior to vitrectomy was selected to the IVB group. The present meta-analysis involved 142 eyes treated with vitrectomy alone and 139 eyes receiving vitrectomy with IVB pretreatment. The selection of seven studies is summarised in tables 1, 2. Each study revealed that there were no significant differences in preoperative demographic features and other factors that may have influenced surgery, such as previous panretinal photocoagulation.

**Intraoperative outcome parameters**

Five studies reported data for intraoperative bleeding. The studies applied differential scales for the evaluation of intraoperative bleeding. Clinically significant intraoperative bleeding (moderate to severe) was included in this meta-analysis.

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**Table 1** Characteristics of studies of vitrectomy alone versus vitrectomy with IVB pretreatment included in the meta-analysis

<table>
<thead>
<tr>
<th>Trial (first author, year)</th>
<th>Location</th>
<th>Design of trial</th>
<th>System baselines</th>
<th>Ocular baselines</th>
<th>Jadad score</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yang, 2008</td>
<td>Taiwan</td>
<td>Prospective, non-randomised, comparative case study</td>
<td>NS: DM, Systematic hypertension, Systematic hypertension</td>
<td>NS: DM, DM, Age, Diabetes, DM, Panretinal photocoagulation</td>
<td>4</td>
</tr>
</tbody>
</table>

Table 2: Characteristics of surgical procedures of vitrectomy alone versus vitrectomy with intravitreal bevacizumab pretreatment included in the meta-analysis

<table>
<thead>
<tr>
<th>Dosage of intravitreal bevacizumab</th>
<th>Time prior to pars plana vitrectomy</th>
<th>Type of pars plana vitrectomy</th>
<th>Surgery: IPV vs IPV</th>
<th>Type of PDR</th>
<th>Complications of PDR</th>
<th>Postoperative outcome parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.25 mg (0.05 ml)</td>
<td>7–9 days</td>
<td>PPV</td>
<td>Done</td>
<td>Active PDR, VH and TRD</td>
<td>Active severe PDR; visible large new vessels within the proliferative zone (PV), new vessels within the non-proliferative zone (NPV), active TRD, angiomaticous RD, or TRD complicated with VH</td>
<td>Two studies reported data for the reabsorption time of blood or fluid. Analysis of these data showed that the mean reabsorption time was statistically significantly less in the IVB treatment group than in the vitrectomy alone group (WMD 14.15; 95% CI 1.17 to 27.09; p = 0.03) (figure 1A). A sensitivity analysis was performed to examine the effect of excluding the non-randomised study, and the statistical results did not change (WMD 23.70; 95% CI 18.01 to 29.39; p &lt; 0.00001). This comparison generated a significant heterogeneity (I² = 75.9%), and the result must be interpreted with caution.</td>
</tr>
<tr>
<td>1.25 mg (0.05 ml)</td>
<td>2 weeks</td>
<td>PPV</td>
<td>Done</td>
<td>Active PDR; VH and TRD</td>
<td>Active severe PDR; visible large new vessels within the proliferative zone (PV), new vessels within the non-proliferative zone (NPV), active TRD, angiomaticous RD, or TRD complicated with VH</td>
<td>Five studies reported data for the proportion of recurrent VH. The studies used different scales for recurrent VH evaluation. Clinical significant recurrent VH (severe recurrent VH with no fundus details) were included in this meta-analysis. Furthermore, all incidences of recurrent VH during follow-up were included. Analysis of these data showed that the proportion of recurrent VH was almost statistically significantly less in the treatment group than in the control group (OR 5.48; 95% CI 0.97 to 31.02; p = 0.05) (figure 2A). A sensitivity analysis was performed to examine the effect of excluding the non-randomised study, and the statistical results did not change (OR 7.52; 95% CI 0.97 to 58.40; p = 0.05). Owing to the poor definition and heterogeneity (I² = 93.6%, 73.2%, respectively) of criteria for the reabsorption time and recurrent VH, the above two results should be interpreted with caution.</td>
</tr>
<tr>
<td>2.5 mg (0.1 ml)</td>
<td>5–7 days</td>
<td>20-gauge PPV</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>Four studies reported data for the frequency of endodiathermy. Analysis of these data showed that the frequency of endodiathermy was statistically significantly less in the treatment group than in the control group (OR 15.06; 95% CI 1.00 to 15.79; p = 0.05) (figure 1B).</td>
</tr>
<tr>
<td>1.25 mg (0.05 ml)</td>
<td>5–7 days</td>
<td>20-gauge PPV</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>Three studies reported data for the number of iatrogenic retinal tears. Analysis of these data showed that the incidence of iatrogenic retinal tears was almost significantly less in the IVB treatment group (OR 3.72; 95% CI 1.00 to 13.79; p &lt; 0.0001) (figure 1C).</td>
</tr>
<tr>
<td>2.5 mg (0.1 ml)</td>
<td>3–5 days</td>
<td>20-gauge PPV</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>Five studies reported data for the mean surgical time. Three studies reported data for the reabsorption time of blood or fluid. Analysis of these data showed that the mean reabsorption time was statistically significantly less in the IVB treatment group than in the vitrectomy alone group (WMD 14.15; 95% CI 1.17 to 27.09; p = 0.03) (figure 1A). A sensitivity analysis was performed to examine the effect of excluding the non-randomised study, and the statistical results did not change (WMD 23.70; 95% CI 18.01 to 29.39; p &lt; 0.00001). This comparison generated a significant heterogeneity (I² = 75.9%), and the result must be interpreted with caution.</td>
</tr>
<tr>
<td>1.25 mg (0.05 ml)</td>
<td>7 days</td>
<td>20-gauge PPV</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>Owing to the poor definition and heterogeneity (I² = 93.6%, 73.2%, respectively) of criteria for the reabsorption time and recurrent VH, the above two results should be interpreted with caution.</td>
</tr>
<tr>
<td>1.25 mg (0.05 ml)</td>
<td>7 days</td>
<td>Standard PPV</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>Six studies reported data for BCVA. Analysis of these data showed that the mean BCVA was statistically significantly better in the treatment group than in the control group (WMD: 0.31; 95% CI 0.10 to 0.52; p = 0.003) (figure 2C).</td>
</tr>
<tr>
<td>1.25 mg (0.05 ml)</td>
<td>7 days</td>
<td>Standard PPV</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>Six studies reported data for postoperative complications. Analysis of these data showed no statistically significant differences in the early (&lt;1 week) elevation of IOP (≥25 mm Hg), late (≥1 week) elevation of IOP (≥25 mm Hg), incidence of final RD and proportion of repeat vitrectomy (table 5).</td>
</tr>
</tbody>
</table>

Analysis of these data showed that the incidence of intraoperative bleeding was statistically significantly less in the IVB pretreatment group than in the vitrectomy alone group (OR 8.85; 95% CI 2.08 to 37.61; p = 0.003) (figure 1A). A sensitivity analysis was performed to examine the effect of excluding the non-randomised study, whereby the statistical results did not change (OR 14.51; 95% CI 3.06 to 68.85; p = 0.0008). Owing to the poor definition and heterogeneity (I² = 73.9%) of criteria for the intraoperative bleeding, the result should be interpreted with caution. Four studies reported data for the frequency of endodiathermy. Analysis of these data showed that the frequency of endodiathermy was statistically significantly less in the treatment group than in the control group (OR 15.06; 95% CI 4.01 to 56.53; p < 0.0001) (figure 1B). Three studies reported data for the number of iatrogenic retinal tears. Analysis of these data showed that the incidence of iatrogenic retinal tears was almost significantly less in the IVB treatment group (OR 3.72; 95% CI 1.00 to 13.79; p = 0.05) (figure 1C). Five studies reported data for the mean surgical time. Three studies reported data for the reabsorption time of blood or fluid. Analysis of these data showed that the mean reabsorption time was statistically significantly less in the IVB treatment group than in the vitrectomy alone group (WMD 14.15; 95% CI 1.17 to 27.09; p = 0.03) (figure 1A). A sensitivity analysis was performed to examine the effect of excluding the non-randomised study, and the statistical results did not change (WMD 23.70; 95% CI 18.01 to 29.39; p < 0.00001). This comparison generated a significant heterogeneity (I² = 75.9%), and the result must be interpreted with caution. Three studies reported data for the number of iatrogenic retinal tears. Analysis of these data showed that the incidence of iatrogenic retinal tears was almost significantly less in the IVB treatment group (OR 3.72; 95% CI 1.00 to 13.79; p = 0.05) (figure 1C). Five studies reported data for the mean surgical time. Three studies reported data for the reabsorption time of blood or fluid. Analysis of these data showed that the mean reabsorption time was statistically significantly less in the IVB treatment group than in the vitrectomy alone group (WMD 14.15; 95% CI 1.17 to 27.09; p = 0.03) (figure 1A). A sensitivity analysis was performed to examine the effect of excluding the non-randomised study, and the statistical results did not change (WMD 23.70; 95% CI 18.01 to 29.39; p < 0.00001). This comparison generated a significant heterogeneity (I² = 75.9%), and the result must be interpreted with caution.
Publication bias
Based on a visual analysis of the funnel plots, no obvious evidence of publication bias was found (figure 3).

DISCUSSION
The present meta-analysis revealed that IVB injection before vitrectomy for PDR can reduce intraoperative bleeding and the frequency of endodiathermy, and shorten the mean surgical time. It can also shorten reabsorption time of blood after vitrectomy, decrease the incidence of recurrent VH and improve BCVA.

Bevacizumab is a recombinant humanised monoclonal anti-vascular endothelial growth factor antibody used to induce regression of neovascularisation and reduce permeability. It has increasingly been used to treat choroidal neovascularisation and...
diabetic macular oedema. It has also proven to be effective for the treatment of PDR complicated by VH. Results of fluorescein angiography revealed a reduction in leakage from the foci of neovascularisation and regression of the neovascular component of fibrovascular tissue in eyes with PDR within 1 week after IVB.

Based on these observations, it was suggested that IVB may reduce the incidence of intraoperative and postoperative haemorrhage in diabetic vitrectomy. Chen first reported that preoperative IVB was helpful in facilitating vitrectomy in severe PDR. Many clinical trials proved that IVB pretreatment ameliorates fundus conditions before vitrectomy. In Ahmadieh's study, owing to a significant resolution of VH and improvement of vision after IVB injection, nine eyes (25.7%) initially scheduled for vitrectomy obviated the need for surgical intervention. Laboratory experiment provided objective, quantifiable data that a lower number of erythrocytes retrieved from the vitrectomy cassette were observed in those patients treated with IVB 2 weeks before surgery. From surgeons' experiences, the regression of the vascular component of the fibrovascular complexes after IVB facilitates segmentation and delamination of membranes. This is because they are less adhesive to the underlying retina and readily separated from the retina. The haemodynamic changes in retinal

![Figure 2](http://bjo.bmj.com/)

**Figure 2** Forest plots of weighted mean difference (WMD) of reabsorption time of blood (A), OR of proportions of recurrent vitreous haemorrhage (B) and WMD of final mean BCVA (C) after surgery comparing vitrectomy alone to vitrectomy with intravitreal bevacizumab (IVB) pretreatment. PPV, pars plana vitrectomy.

### Table 3 Postoperative complications of vitrectomy alone versus vitrectomy with intravitreal bevacizumab pretreatment in the meta-analysis

<table>
<thead>
<tr>
<th>Complications</th>
<th>No of studies</th>
<th>Pars plana vitrectomy group</th>
<th>Intravitreal bevacizumab+pars plana vitrectomy group</th>
<th>Percentage rate difference (95% CI)</th>
<th>p Value for overall effect</th>
</tr>
</thead>
<tbody>
<tr>
<td>Early elevation of intraocular pressure (&lt;25 mm Hg)</td>
<td>3, 3, 6</td>
<td>13/66</td>
<td>12/87</td>
<td>1.08 (0.36 to 3.27)</td>
<td>0.89</td>
</tr>
<tr>
<td>Late elevation of intraocular pressure (&gt;25 mm Hg)</td>
<td>3, 3, 6</td>
<td>8/57</td>
<td>0/48</td>
<td>17.00 (0.91 to 319.22)</td>
<td>0.06</td>
</tr>
<tr>
<td>Final retinal detachment</td>
<td>3, 3, 4</td>
<td>5/51</td>
<td>2/54</td>
<td>2.44 (0.52 to 11.46)</td>
<td>0.26</td>
</tr>
<tr>
<td>Repeat vitrectomy</td>
<td>2, 4</td>
<td>3/42</td>
<td>1/46</td>
<td>2.69 (0.37 to 19.33)</td>
<td>0.32</td>
</tr>
</tbody>
</table>
Facilitating early visual rehabilitation for patients is another important clinical outcome. This positive effect may be due to minimal injury to the retinal tissue during surgery, a minimal extent of haemorrhage and posterior focal reproliferation, or clear visual pathway after surgery in bevacizumab-treated cases. The studies included did not report apparent macular oedema in patients with diabetes before and after vitrectomy, but surgical manoeuvre or surgical trauma, such as delamination, could influence macular physiology function. The effect of reducing diabetic macular oedema might play a role. Many clinical trials have shown that IVB can reduce diabetic macular oedema and result in a better visual recovery.

Di Lauro compared a 7-day with 20-day previtrectomy administration of bevacizumab. Clinical outcomes showed no statistically significant differences between the two groups, but intraoperative severe bleeding, frequency of endodiathermy, iatrogenic retinal break, silicone-oil tamponade, surgical mean time and the rate of recurrent VH featured more in the 20-day group. Concerns about potential harm of increasing vitreoretinal traction owing to rapid neovascular invasion with fibrosis contraction by IVB pretreatment must be raised. Many studies have noted complications of aggravation of fibrosis after IVB injection for active progressive PDR. One recent study found that tractional retinal detachments were worsened or newly created in 11 eyes (5.2%) of 211 IVB injections in severe PDR. The time interval between IVB injection and diagnosis of tractional retinal detachment was on average 15 days. Ishikawa performed IVB injections in eight eyes and found that two eyes, injected 7 days before the operation, had severe fibrosis, resulting in some surgical complications. The evidence available indicates that the effect of IVB on regression of new vessels is rapid, often evident after 24 h. It seems that 5–5 days or even less time is adequate for IVB to exert its antiangiogenic effect. A dosage of 1.25 mg is commonly used. Some studies have revealed that IVB at doses of 1.25 to 2.5 mg provide similar clinical outcomes in diabetic retinopathy. Clarification of optimal interval time and dosage of IVB still needs an RCT with a larger sample size.

The selected trials revealed that no local and systemic complications related to IVB were observed in the treatment groups. The postoperative final RD and repeat surgery showed no significant differences. These are possibly attributable to the natural process of PDR. Long-term follow-up studies (>6 months) are needed.

This meta-analysis may have some limitations. First, we cannot fully exclude publication bias. The number of included studies is insufficient to carry out a further statistical test, to detect publication bias through asymmetry plot. In addition, we did not attempt to gain access to unpublished results. Second, the studies included were heterogeneous in terms of study location, population, number of patients from different studies and basal condition. Access to individual-level data could certainly have improved the quality of adjustment as well as the precision of estimates. Third, there was a large disparity in study quality (relative methodological strengths and weaknesses), as reflected by the Jadad score. This review unfortunately did not comprise a sufficient number of studies of a high level Jadad score to justify a performance of subgroup meta-analyses.

In conclusion, preoperative IVB may represent a new strategy to make vitrectomy safer and more effective for severe PDR. RCTs with larger sample sizes or systematic studies are needed to better evaluate the long-term benefits and safety of bevacizumab pretreatment for severe PDR.
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