

**ONLINE-ONLY SUPPLEMENTARY MATERIAL****Online-only Tables:**

Supplementary Table 1. Inclusion and exclusion criteria

Supplementary Table 2. Raised IOP during 6 months of study

**Online-Only Figures:**

Supplementary Figure 1. Frequency of retreatment required during the first 6 months of the trial

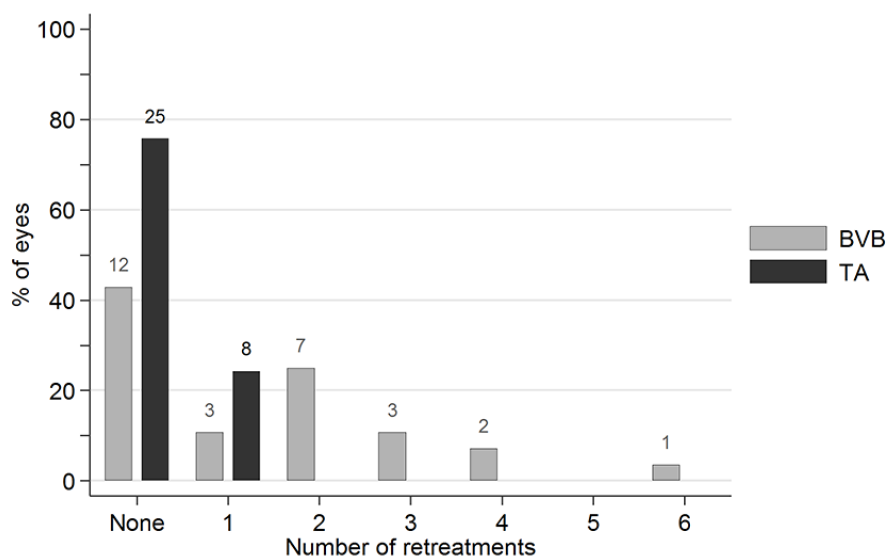
**Supplementary Table 1. Inclusion and exclusion criteria**

<p><b>Inclusion criteria</b></p> <p>Visually significant cataract requiring surgery AND Centre-involving diabetic macular oedema (DMO) at baseline OR Centre-involving DMO within 24 months of study enrolment Age more than 18 years</p>
<p><b>Exclusion criteria</b></p> <p>Macular oedema from causes other than diabetic retinopathy OR significant angiographic macular ischaemia Concurrent ocular inflammation / infection Loss of vision from other causes (e.g. age related macular degeneration, myopic macular degeneration) Intractable glaucoma OR pre-existing glaucomatous visual field defect Previous history of steroid response ( Intraocular pressure elevation to more than 35mmHg following steroid treatment ) Best corrected visual acuity less than 6/60 in the fellow eye Prior history of adverse reaction/allergy to triamcinolone acetate or anti-vascular endothelial growth factor (VEGF) drugs Previous intravitreal injection of triamcinolone acetate within 10 weeks OR intravitreal injection of anti-VEGF drugs within 3 weeks of study entry Previous macular argon laser photocoagulation within 3 months of study entry Patients requiring systemic steroids for other indications ( more than 5mg of prednisolone daily or equivalent) Pregnancy OR breastfeeding Patients with concurrent severe systemic infections/disease (e.g. septicaemia)</p>

**Supplementary Table 2. Raised IOP during 6 months of study**

ID	Group	IOP								Prior IVI	Retreated
		Baseline	Week 1	Month 1	Month 2	Month 3	Month 4	Month 5	Month 6		
A	TA	16	15	19	16	20	<b>26</b>	15	<b>38</b>	None	Nil
B	TA	13	19	15	DNA	19	<b>27</b>	14	16	None	Nil
C	TA	13	12	12	14	14	13	<b>22</b>	11	None	Month 6
D	TA	15	<b>23</b>	15	14	20	<b>25</b>	17	10	None	Nil
E	BVB	19	15	16	16	14	<b>23</b>	11	14	1x BVB	Nil
F	BVB	14	19	18	17	13	12	16	<b>22</b>	None	Nil
G	BVB	20	10	13	11	14	<b>23</b>	18	15	None	Nil

**BVB**, bevacizumab; **ID**, patient identification; **IOP**, intraocular pressure; **IVI**, previously received intravitreal injections within 24 months prior to study entry/surgery; **TA**, Triamcinolone; **DNA**, patient did not attend that visit



Graph includes data up to and including 6 months for all available subjects to date  
Annotations indicate the actual number of eyes in each retreatment group

**Supplementary Figure 1. Frequency of retreatment required during the first 6 months of the trial**

**BVB**, bevacizumab; **TA**, Triamcinolone.