

ARTICLE OPEN

Check for updates

Presenting clinical characteristics of open globe injuries in ocular trauma: baseline analysis of cases in the ASCOT national clinical trial

Suzie Cro^{1™}, Giles Partington¹, Victoria R. Cornelius¹, Philip J. Banerjee², Tapiwa Margaret Zvobgo³, Edward J. Casswell³, Syed Shahid³, Catey Bunce^{3,4}, Elizabeth Robertson³, Caroline Murphy⁵, Joanna Kelly⁵ and David G. Charteris³

© The Author(s) 2022

BACKGROUND/OBJECTIVES: The Adjunctive Steroid Combination in Ocular Trauma (ASCOT) trial is a unique pragmatic, multicentre, patient and assessor masked, randomised controlled trial. We evaluate the clinical characteristics and pathology of this large trial cohort of patients with open globe injuries undergoing vitreoretinal surgery, including the associations between patient characteristics and their baseline vision.

SUBJECTS/METHODS: We (i) summarise demographics, injury history and ocular history of the 280 participants recruited into the ASCOT trial using descriptive statistics; (ii) analyse the national and seasonal variation across England and Scotland in these participant characteristics; and (iii) explore the associations between participant demographic, trauma history, ocular history and presenting baseline visual acuity (measured using the Early Treatment Diabetic Retinopathy Study, ETDRS) using multivariable regression analyses.

RESULTS: The majority of participants with open globe penetrating injuries were of white ethnicity (233, 84%), male (246, 88%), with a median age of 43 years (IQR 30–55 years). There was considerable variability in presenting visual acuity with 75% unable to read any letters on the ETDRS chart, whilst the median ETDRS letter score was 58 (IQR 24–80) for those who could read \geq 1 letter. The most common causes of injury were workplace related (31%) or interpersonal violence (24%). Previous eye surgery, visual axis corneal scar, lens status, hyphaemia and vitreous haemorrhaging were found to be associated with presenting vision as measured by the ETDRS chart.

CONCLUSION: The ASCOT trial provides valuable insights into the spectrum of pathology of patients with open globe eye injuries undergoing vitreoretinal surgery. The identified causes of injury and clinical presentation of the cases will help in training and resource planning to deal with these often challenging surgical cases.

TRIAL REGISTRATION: EudraCT No. 014-002193-37. HTA Project 12/35/64.

Eye; https://doi.org/10.1038/s41433-022-02206-z

INTRODUCTION

Ocular trauma is a leading cause of monocular visual impairment and monocular blindness worldwide [1]. Globally, 55 million people experience serious ocular trauma every year [2]. It has been estimated that worldwide 1.6 million people are blind as a result of ocular trauma, with 2.3 million having bilateral low vision and 19 million with unilateral vision [2]. The prevalence of ocular trauma has been shown to vary considerably by country, gender, age and socio-economic status [3–6]. Blindness from ocular trauma is a serious health problem that has extensive psychological impacts on patients and their relatives, and major socioeconomic costs [7].

Frequently the posterior segment of the eye, comprising the vitreous humour, retina, choroid, and optic nerve, is affected by injuries that result in visual loss and to prevent severe loss of vision, posterior segment (vitreoretinal) surgery is necessary. In a one-year

population-based prospective study of ocular trauma in Scotland, among cases of serious eye injury blinding outcomes occurred in 26% [8]. Whilst surgical techniques have improved, outcomes remain unsatisfactory. This is mainly due to the development of the intraocular scarring response proliferative vitreoretinopathy (PVR). PVR is the main cause of visual loss in eyes with open globe injuries (defined as a full thickness wound of the eye wall) and is estimated to occur in 10–45% of all open globe injuries [9–12].

Experimental studies have suggested that steroid (triamcinolone acetonide, TA) treatment can reduce the severity of PVR [13]. The primary aim of the Adjunctive Steroid Combination in Ocular Trauma (ASCOT) trial is to assess whether, adjunctive intravitreal and sub-Tenon's triamcinolone acetonide given at the time of surgery improves visual acuity at 6 months compared with standard surgery. ASCOT is a phase 3 randomised controlled trial which recruited a total of 280 eligible patients nationally across 27 sites in England and

¹Imperial Clinical Trials Unit, Imperial College London, London, UK. ²Frimley Health NHS Foundation Trust, London, UK. ³Moorfields Eye Hospital NHS Foundation Trust, London, UK. ⁴The Royal Marsden NHS Foundation Trust, London, UK. ⁵King's Clinical Trials Unit at Kings Health Partners, King's College London, London, UK. ^{Sem}ail: s.cro@imperial.ac.uk

Scotland [14, 15]. The trial began recruitment in December 2014 and completed recruitment in March 2020; final participant follow-up was completed in September 2020.

ASCOT is the only large scale prospective multi-centre national trial to have been undertaken on traumatic open globe injuries and therefore, for the first time, provides a unique opportunity to evaluate the detailed clinical characteristics of a large cohort of patients with open globe injuries. A previous national study of serious eye trauma in Scotland reported an incidence of 1.96/ 100,000 but did not report any detail on clinical presentations [8]. The aims of this analysis are to evaluate the clinical characteristics and pathology of this large trial cohort of patients with open alobe injuries undergoing vitreoretinal surgery, including the associations between patient characteristics and their baseline vision. Specific objectives are (1) to describe demographics, injury history and ocular history of participants recruited into the ASCOT trial, (2) to analyse the national and seasonal variation across England and Scotland in these participant characteristics, and (3) to explore associations between participant demographic, trauma history, ocular history and presenting baseline visual acuity.

SUBJECTS AND METHODS

Full details of the ASCOT study design have been published [14]. Briefly, ASCOT is a pragmatic, multi-centre, double-masked, randomised controlled trial. The study population included adult patients (aged 18 years or over) with a full thickness, open globe injury undergoing vitrectomy with the ability to give informed consent. Exclusion criteria were patients with pre-existing uncontrolled uveitis, previous steroid-induced glaucoma, pregnant or breastfeeding females, those with previous known adverse reactions to triamcinolone, inability to attend regular follow-up, current or planned use of corticosteroid use of a dose above physiological levels. All penetrating injuries attending eye emergency clinics were screened and patients were recruited at vitreoretinal outpatient or emergency clinics at 27 participating vitreoretinal surgical centres throughout the UK. Eligible participants were randomised (1:1) to standard surgery plus adjunctive triamcinolone acetonide or standard surgery alone for the operated eye. Operating surgeons were masked until the end of surgery; participants and study investigators were masked to treatment allocation throughout. For patients with bilateral injuries, data is reported for the worse injured eye, which was considered as the study eye for randomisation and operated on according to randomisation (the better eye received standard treatment). The ASCOT study protocol was approved by the Central London NHS Research Ethics Committee (14/LO/1428).

Data collection and outcomes

In this report, we analyse participant baseline demographics, trauma history and ocular history including baseline biomicroscopic ocular exam results prior to surgery. The baseline eye exam included the Early Treatment Diabetic Retinopathy Study (ETDRS) letter score measured using the standard validated ETDRS chart at a starting distance of 4 m in the eye operated on. If 20 or more letters were read at 4 m, 30 was added to the total letter score. If less than 20 letters were read correctly at 4 m then the distance was moved to 1 m and the number of letters read at 1 m formed the total score. If no letters were seen at 1 m then the score is 0 and test for Counting Fingers, Hand Movements and Light Perception.

Statistical analysis

The target sample size for ASCOT was 300 patients (150 per arm) over a 3-anda-half-year recruitment period. As actual recruitment rate was slower than projected, the recruitment period was extended to a total of 75 months; 280 eligible patients were recruited and all are included within this analysis.

Baseline characteristics are summarised as mean (standard deviation— SD) for approximately normally distributed continuous variables or median (Interquartile range—IQR) where skewed. Categorical variables are summarised by frequencies and percentages. Descriptive data summaries are based on observations only and the number of missing observations are reported. The Pearson's chi-squared test was used to assess seasonal variation in the injury numbers by month, and cause of injury.

To explore the associations between baseline visual acuity (measured using an ETDRS chart at a starting distance of 4 m) and participant

characteristics of interest, multivariable regression analyses were conducted. The clinically important characteristics of interest, informed by previous studies and study team clinical opinions were: previous primary repair, trauma classification, extent of trauma, previous eye surgery, macular disease, other historical ocular conditions, visual axial corneal scarring, hyphaemia, lens state, vitreous haemorrhage, endophthalmitis, retinal status and PVR. Given the high distribution of participants with an ETDRS score of 0 letters indicating very low/no vision, a zero-inflated Negative Binomial model was used to explore associations. This model incorporates two parts; (i) a logistic regression model that models the probability of zero/very low vision (ETDRS = 0) and (ii) a Negative Binomial model that models the amount of vision as measured by the ETDRS letter count score. The Negative Binomial model (part ii) included all the clinically important variables of interest. The logistic regression model (part i) used a subset of the clinically important variables that were first identified to be associated with zero vision (see Supplementary file 1, Table S3).

Retinal detachment at baseline and associations between the clinically important characteristics of interest were explored in an additional analysis using a logistic regression model. Estimates from the logistic regression models are summarised as Odds Ratios (OR). Estimates from the zero-inflated Negative binomial model are presented as Incidence Rate Ratios (IRR) which represent the relative risk of a higher ETDRS score for the associated variable. Estimates are accompanied with 95% confidence intervals. A *p* value <0.20 was used to indicate a potential association and a result of interest. As exploration of missing data patterns revealed <10% of the sample would be excluded due to missing data, the analysis included all participants with complete data under the assumption that any missing data were missing completely at random. All statistical analysis was performed using Stata/IC version 15.0 or above.

RESULTS

Baseline characteristics

Table 1 summarises the baseline demographics, trauma history and ocular history for the study cohort of 280 patients. The majority of participants were of white ethnicity (233, 84%), male (246, 88%), with a median age of 43 years (IQR 30–55 years). The median age of the females was 63.0 years (IQR 44.6–75.6) and for the males was 42.2 years (IQR 29.9–3.3). The causes of the ocular trauma are indicated in Table 1; 14% (38) were classified as "other injury" and these are listed in Supplementary Table S1.

The majority of participants (209, 75%) had a score of zero on the ETDRS chart meaning they presented with very low vision of counting fingers or worse. The median ETDRS score among those with an ETDRS score greater than zero (71, 25%), was 58 letters (IQR 24–80) demonstrating high variability in presenting vision.

Regional variability

The majority (148, 53%) of ASCOT participants were recruited from hospitals in London, a quarter came from the Midlands & Northern England (70, 25%), 55 (20%) were recruited in Southern England and 7 from Scotland (3%). London had a higher proportion of Black, Asian and Minority Ethnic (BAME) participants (24%) than any other region (Southern England: 4%, Midlands and North England: 11% and Scotland: 0%). This variation reflects the regional ethnic diversity in the wider population, at the most recent 2011 census 40% of London residents identified as BAME versus 8% in Southern England, 11% in the Midlands and North England and 4% in Scotland [16, 17]. Supplementary Table S2 summarises baseline characteristics by region (London, Southern England, Midlands & Northern England, and Scotland).

There was little difference between the causes of injury across the regions. In London, nearly half (70, 47%) of all injuries were categorised as penetrating, with 46 (31%) as rupture. However, all other regions had substantially higher amounts of ruptures than penetrations. Similarly, injuries in London were more frequently to zone 1 of the eye (corneal injury) than to zone 2 (scleral anterior) (39% vs 35% [58 vs 52]), whereas the reverse was true in other regions. There was a higher history of ocular co-morbidities for patients in London (21, 14%) and Midlands & North England (7, 10%).

S. Cro et al	١.
--------------	----

Table 1. continued

Table 1.	Baseline demographics,	trauma history	/ and	ocular	histor	v.

Variable (n missing)N%Gender (n, %) (n = 0)3412%Female24688%Region (n, %) (n = 0)14853%South England5525%South England7025%Soctand7025%Soctand7025%South England7025%South England7025%South England7025%South England7025%South England7025%Black207%Asian186%Other62%Mixed13749%Eye injured (n, %) (n = 0)1%Kight13749%Left13849%Both662%How was the eye injured (n, %) (n = 0)4%Interpersonal violence6624%Sports injury104%Interpersonal violence6624%Domestic gardening83%Domestic gardening135%Iatrogenic135%Iatrogenic135%Severity of trauma: classification (n, %) (n = 1)2%Sceral poterior to muscle insertion1104%IoPB2125%34%Domestic gardening135%Severity of trauma: classification (n, %) (n = 0)2%Severity of trauma: extent (n, %) (n = 1)2%Sceral poterior to muscle insertion1104% <t< th=""><th>Table 1. Dasenne demographics, tradma history</th><th>und ocun</th><th>in miscory.</th></t<>	Table 1. Dasenne demographics, tradma history	und ocun	in miscory.
Female 34 12% Male 246 88% Region (n, %) [n = 0]	Variable [n missing]	N	%
Male 246 88% Region (n, %) [n = 0]	Gender (<i>n</i> , %) [<i>n</i> = 0]		
Region (n, %) [n = 0] London 148 53% South England 55 20% Midlands & North England 70 25% Scottand 70 25% Ethnicity (n, %) [n = 0]	Female	34	12%
London 148 53% South England 55 20% Midlands & North England 70 25% Scottand 70 25% Ethnicity (n, %) [n = 0] """"""""""""""""""""""""""""""""""""	Male	246	88%
South England 55 20% Midlands & North England 70 25% Scotland 70 25% Ethnicity (n, %) [n = 0] """"""""""""""""""""""""""""""""""""	Region (n, %) [n = 0]		
Midlands & North England 70 25% Scotland 7 3% Ethnicity (n, %) [n = 0]	London	148	53%
Scotland73%Ethnicity (n, %) [n = 0]23383%Black207%Asian186%Other62%Mixed31%Eye injured (n, %) [n = 0]13749%Eye injured (n, %) [n = 0]13749%Both52%How was the eye injured (n, %) [n = 0]13731%Workplace incident8831%Road traffic accident114%Interpersonal violence6624%Sports injury104%Other injury ⁸ 3713%Other domestic ^b 218%Domestic gardening83%Domestic gardening31%Fall238%Previous primary repair (n, %) [n = 1]20573%Severity of trauma: classification (n, %) [n = 2]31%Nutpure11340%10FPenetrating10337%1%Ferlorating114%10FIOFB5319%1%Severity of trauma: extent (n, %) [n = 2]26%26%Scleral posterior to muscle insertion7226%Severity of trauma: RAPD present? - Yes (n, %)4315%Incal posterior to muscle insertion14050%Glauccoma (n, %) [n = 1]41%Previous eye surgery (study eye) (n, %) [n = 0]145%Macular disease (n, %) [n = 0]145%Macular disease	South England	55	20%
Ethnicity (n, %) [n = 0] White 233 83% Black 20 7% Asian 18 6% Other 6 2% Mixed 3 1% Eye injured (n, %) [n = 0] 137 49% Left 138 49% Both 5 2% How was the eye injured (n, %) [n = 0] Workplace incident 88 31% Road traffic accident 11 4% Interpersonal violence 66 24% Sports injury 10 4% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% Iatrogenic 3 1% Fall 23 8% Previous primary repair (n, %) [n = 1] 23 8% Previous primary repair (n, %) [n = 2] 2% 3% Porture 113 40% Previous primary repair (n, %) [n = 2] 2% 3% Soleral posterior to muscle insertion 110 4%	Midlands & North England	70	25%
White 233 83% Black 20 7% Asian 18 6% Other 6 2% Mixed 3 1% Eye injured (n, %) [n = 0] 137 49% Left 138 49% Both 5 2% How was the eye injured (n, %) [n = 0] Workplace incident 88 31% Road traffic accident 11 4% Interpersonal violence 66 24% Sports injury 10 4% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% Iatrogenic 3 1% Fall 23 8% Previous primary repair (n, %) [n = 1] 20 7% Severity of trauma: classification (n, %) [n = 2] 2% Cornea 95 34% DFB 53 19% Severity of trauma: RAPD present? - Yes (n, %) 15%	Scotland	7	3%
Black 20 7% Asian 18 6% Other 6 2% Mixed 3 1% Eye injured (n, %) [n = 0]	Ethnicity (n , %) [$n = 0$]		
Asian 18 6% Other 6 2% Mixed 3 1% Eye injured (n, %) [n = 0]	White	233	83%
Other 6 2% Mixed 3 1% Eye injured (n, %) [n = 0]	Black	20	7%
Mixed 3 1% Eye injured (n, %) [n = 0] 137 49% Left 138 49% Both 5 2% How was the eye injured (n, %) [n = 0] 38 31% Workplace incident 88 31% Road traffic accident 11 4% Interpersonal violence 66 24% Sports injury 10 4% Other injury ^a 37 13% Domestic gardening 8 3% Domestic DIY 13 5% Iatrogenic 3 1% Fall 23 8% Previous primary repair (n, %) [n = 1] 205 73% Severity of trauma: classification (n, %) [n = 0] 37 13% Penetrating 103 37% IOFB 34 10% Severity of trauma: extent (n, %) [n = 2] 26% Severity of trauma: extent (n, %) [n = 2] 26% Severity of trauma: RAPD present? - Yes (n, %) 43 15%	Asian	18	6%
Eye injured $(n, %) [n = 0]$ Right 137 49% Left 138 49% Both 5 2% How was the eye injured $(n, %) [n = 0]$ """"""""""""""""""""""""""""""""""""	Other	6	2%
Right 137 49% Left 138 49% Both 5 2% How was the eye injured $(n, \%) [n = 0]$	Mixed	3	1%
Left 138 49% Both 5 2% How was the eye injured $(n, \%) [n = 0]$ ************************************	Eye injured (n , %) [$n = 0$]		
Both 5 2% How was the eye injured (n, %) [n = 0]	Right	137	49%
How was the eye injured $(n, \%)$ $[n = 0]$ Workplace incident 88 31% Road traffic accident 11 4% Interpersonal violence 66 24% Sports injury 10 4% Other injury ^a 37 13% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair $(n, %)$ $[n = 1]$ 205 73% Severity of trauma: classification $(n, %)$ $[n = 0]$ Was 13 Rupture 113 40% Perforating 103 37% IOFB 33 19% Severity of trauma: extent $(n, \%)$ $[n = 2]$ Vorea 26% Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 13 15% $[n = 2]$ Not documented 140 </td <td>Left</td> <td>138</td> <td>49%</td>	Left	138	49%
How was the eye injured $(n, \%)$ $[n = 0]$ Workplace incident 88 31% Road traffic accident 11 4% Interpersonal violence 66 24% Sports injury 10 4% Other injury ^a 37 13% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair $(n, %)$ $[n = 1]$ 205 73% Severity of trauma: classification $(n, %)$ $[n = 0]$ Was 13 Rupture 113 40% Perforating 103 37% IOFB 33 19% Severity of trauma: extent $(n, \%)$ $[n = 2]$ Vorea 26% Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 13 15% $[n = 2]$ Not documented 140 </td <td>Both</td> <td>5</td> <td>2%</td>	Both	5	2%
Workplace incident 88 31% Road traffic accident 11 4% Interpersonal violence 66 24% Sports injury 10 4% Other injury ^a 37 13% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair (n, %) [n = 1] 205 73% Severity of trauma: classification (n, %) [n = 0] 40% Nepture 113 40% Penetrating 103 37% Perforating 11 4% IOFB 3 19% Severity of trauma: extent (n, %) [n = 2] 26% Severity of trauma: RAPD present? - Yes (n, %) 43 15% [n = 2] Not documented 140 50% Glaucoma (n, %) [n = 1] 4 1% Previous eye surgery (study eye) (n, %) [n = 2] 26% Visual axis corneal scar (n, %) [n = 0] 72 26%			
Road traffic accident 11 4% Interpersonal violence 66 24% Sports injury 10 4% Other injury ^a 37 13% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair (n, %) [n = 1] 205 73% Severity of trauma: classification (n, %) [n = 0] 7% Rupture 113 40% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent (n, %) [n = 2] 7 26% Cornea 95 34% Scleral posterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes (n, %) 43 15% Glaucoma (n, %) [n = 1] 1 0%		88	31%
Interpersonal violence 66 24% Sports injury 10 4% Other injury ^a 37 13% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DlY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair (n, %) [n = 1] 205 73% Severity of trauma: classification (n, %) [n = 0] 7% Rupture 113 40% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent (n, %) [n = 2] V V Cornea 95 34% Scleral posterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes (n, %) 43 15% [n = 2] Not documented 140 50% Glaucoma (n, %) [n = 1] 1			
Sports injury 10 4% Other injury ^a 37 13% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair (n, %) [n = 1] 205 73% Severity of trauma: classification (n, %) [n = 0] 7% Rupture 113 40% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent (n, %) [n = 2] $Cornea$ 95 Cornea 95 34% Scleral posterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes (n, %) 43 15% [n = 2] Not documented 140 50% Glaucoma (n, %) [n = 1] 1 0% Other historic ocular conditions (n, %) [n = 2] 29 10% Visual axis corneal scar (n, %) [n = 0]			
Other injury ^a 37 13% Other domestic ^b 21 8% Domestic gardening 8 3% Domestic DIY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair $(n, %) [n = 1]$ 205 73% Severity of trauma: classification $(n, %) [n = 0]$ 73 37% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent $(n, %) [n = 2]$ V Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 12 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ Not documented 140 50% Glaucoma $(n, \%) [n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%) [n = 0]$ 149 53% Macular disease $(n, \%) [n = 0]$ 52 19% <td></td> <td></td> <td></td>			
Other domestic b218%Domestic gardening83%Domestic DIY135%latrogenic31%Fall238%Previous primary repair $(n, \%) [n = 1]$ 2073%Severity of trauma: classification $(n, \%) [n = 0]$ 73%Rupture11340%Penetrating10337%Perforating114%IOFB5319%Severity of trauma: extent $(n, \%) [n = 2]$ 72Cornea9534%Scleral posterior to muscle insertion11040%Severity of trauma: RAPD present? - Yes $(n, \%)$ 15% $[n = 2]$ Not documented14050%Glaucoma $(n, \%) [n = 1]$ 41%Previous eye surgery (study eye) $(n, \%) [n = 0]$ 14953%Macular disease $(n, \%) [n = 1]$ 10%Other historic ocular conditions $(n, \%) [n = 2]$ 2910%Visual axis corneal scar $(n, \%) [n = 0]$ 7226%Uveitis $(n, \%) [n = 0]$ 5219%14%Hyphaemia level $(n, \%) [n = 0]$ 5018%>50%4416%16%Iris state $(n, \%) [n = 4]$ 11140%Incomplete13549%			
Domestic gardening83%Domestic DIY135%latrogenic31%Fall238%Previous primary repair $(n, \%) [n = 1]$ 237%Severity of trauma: classification $(n, \%) [n = 0]$ 7%Rupture11340%Penetrating10337%Perforating114%IOFB5319%Severity of trauma: extent $(n, \%) [n = 2]$ 72Cornea9534%Scleral anterior to muscle insertion11040%Severity of trauma: RAPD present? - Yes $(n, \%)$ 15% $[n = 2]$ 14050%Macular disease $(n, \%) [n = 1]$ 41%Previous eye surgery (study eye) $(n, \%) [n = 0]$ 14953%Macular disease $(n, \%) [n = 1]$ 10%Other historic ocular conditions $(n, \%) [n = 2]$ 2910%Visual axis corneal scar $(n, \%) [n = 0]$ 7226%Uveitis $(n, \%) [n = 0]$ 5219%Hyphaemia level $(n, \%) [n = 0]$ 5018%>50%4416%Iris state $(n, \%) [n = 4]$ 11140%Incomplete13549%			
Domestic DIY 13 5% latrogenic 3 1% Fall 23 8% Previous primary repair $(n, \%) [n = 1]$ 205 73% Severity of trauma: classification $(n, \%) [n = 0]$ 73% Rupture 113 40% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent $(n, \%) [n = 2]$ Cornea 95 34% Scleral anterior to muscle insertion 110 40% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ Not documented 140 50% Glaucoma $(n, \%) [n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%) [n = 0]$ 149 53% Macular disease $(n, \%) [n = 1]$ 1 0% Other historic ocular conditions $(n, \%) [n = 2]$ 29 10% Visual axis corneal scar $(n, \%) [n = 0]$ 52 19% Hyphaemia level $(n, \%) [n = 0]$ 186 66%			
latrogenic31%Fall238%Previous primary repair $(n, \%)$ $[n = 1]$ 20573%Severity of trauma: classification $(n, \%)$ $[n = 0]$ 73%Rupture11340%Penetrating10337%Perforating114%IOFB5319%Severity of trauma: extent $(n, \%)$ $[n = 2]$ 72Cornea9534%Scleral anterior to muscle insertion11040%Scleral posterior to muscle insertion7226%Severity of trauma: RAPD present? - Yes $(n, \%)$ 4315% $[n = 2]$ 14050%Glaucoma $(n, \%)$ $[n = 1]$ 41%Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 14953%Macular disease $(n, \%)$ $[n = 1]$ 10%Other historic ocular conditions $(n, \%)$ $[n = 2]$ 2910%Visual axis corneal scar $(n, \%)$ $[n = 0]$ 7226%Uveitis $(n, \%)$ $[n = 0]$ 5219%Hyphaemia level $(n, \%)$ $[n = 0]$ 5219%Hyphaemia level $(n, \%)$ $[n = 0]$ 18666%<50%			
Fall238%Previous primary repair $(n, \%)$ $[n = 1]$ 20573%Severity of trauma: classification $(n, \%)$ $[n = 0]$			
Previous primary repair $(n, \%)$ $[n = 1]$ 205 73% Severity of trauma: classification $(n, \%)$ $[n = 0]$ 113 40% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent $(n, \%)$ $[n = 2]$ 53 19% Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ Not documented 140 50% Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 186 66% <50%	-		
Severity of trauma: classification $(n, \%)$ $[n = 0]$ Rupture 113 40% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent $(n, \%)$ $[n = 2]$ 53 19% Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ Not documented 140 50% Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 50 18% <50%			
Rupture 113 40% Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent $(n, \%) [n = 2]$ 53 34% Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ Not documented 140 50% Glaucoma $(n, \%) [n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%) [n = 0]$ 149 53% Macular disease $(n, \%) [n = 1]$ 1 0% 0% Other historic ocular conditions $(n, \%) [n = 2]$ 29 10% Visual axis corneal scar $(n, \%) [n = 0]$ 52 19% Hyphaemia level $(n, \%) [n = 0]$ 50 18% <50%		205	7370
Penetrating 103 37% Perforating 11 4% IOFB 53 19% Severity of trauma: extent $(n, \%) [n = 2]$ 53 34% Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ Not documented 140 50% Glaucoma $(n, \%) [n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%) [n = 0]$ 149 53% Macular disease $(n, \%) [n = 1]$ 1 0% Other historic ocular conditions $(n, \%) [n = 2]$ 29 10% Visual axis corneal scar $(n, \%) [n = 0]$ 72 26% Uveitis $(n, \%) [n = 0]$ 52 19% Hyphaemia level $(n, \%) [n = 0]$ 52 19% No hyphaemia 186 66% <50%		112	400/-
Perforating 11 4% IOFB 53 19% Severity of trauma: extent $(n, \%) [n = 2]$			
IOFB 53 19% Severity of trauma: extent $(n, \%)$ $[n = 2]$ 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ 140 50% Sclaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 72 26% Sclow 50 18% 50% <50%			
Severity of trauma: extent $(n, \%)$ $[n = 2]$ Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ 140 50% Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%			
Cornea 95 34% Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% $[n = 2]$ 140 50% Sdaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%		22	19%
Scleral anterior to muscle insertion 110 40% Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% Not documented 140 50% Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%	•	05	2.40/
Scleral posterior to muscle insertion 72 26% Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% Not documented 140 50% Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 50 18% <50%			
Severity of trauma: RAPD present? - Yes $(n, \%)$ 43 15% Not documented 140 50% Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%			
[n = 2] Not documented 140 50% Glaucoma $(n, %)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, %)$ $[n = 0]$ 149 53% Macular disease $(n, %)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, %)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, %)$ $[n = 0]$ 72 26% Uveitis $(n, %)$ $[n = 0]$ 52 19% Hyphaemia level $(n, %)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%	•		
Not documented 140 50% Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%		43	15%
Glaucoma $(n, \%)$ $[n = 1]$ 4 1% Previous eye surgery (study eye) $(n, \%)$ $[n = 0]$ 149 53% Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%		140	50%
Previous eye surgery (study eye) $(n, \%) [n = 0]$ 149 53% Macular disease $(n, \%) [n = 1]$ 1 0% Other historic ocular conditions $(n, \%) [n = 2]$ 29 10% Visual axis corneal scar $(n, \%) [n = 0]$ 72 26% Uveitis $(n, \%) [n = 0]$ 52 19% Hyphaemia level $(n, \%) [n = 0]$ 52 19% No hyphaemia 186 66% <50%			
Macular disease $(n, \%)$ $[n = 1]$ 1 0% Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 19% No hyphaemia 186 66% <50%			
Other historic ocular conditions $(n, \%)$ $[n = 2]$ 29 10% Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 18% <50%	, , , , , , , , , , , , , , , , , , , ,		
Visual axis corneal scar $(n, \%)$ $[n = 0]$ 72 26% Uveitis $(n, \%)$ $[n = 0]$ 52 19% Hyphaemia level $(n, \%)$ $[n = 0]$ 52 18% <50%			
Uveitis (n, %) $[n = 0]$ 5219%Hyphaemia level $(n, \%) [n = 0]$ 18666%<50%			
Hyphaemia level (n, %) $[n = 0]$ No hyphaemia 186 66% <50%			
No hyphaemia 186 66% <50%		32	1970
<50%	<i></i>	196	66%
>50% 44 16% Iris state (n, %) [n = 4] 111 40% Normal 111 40% Incomplete 135 49%			
Iris state (n, %) [n = 4] Normal 111 40% Incomplete 135 49%			
Normal 111 40% Incomplete 135 49%		44	16%
Incomplete 135 49%			400/
Incarcerated 30 11%			
	Incarcerated	30	11%

N	%
70	25%
96	35%
2	1%
20	7%
89	32%
182	66%
139	76%
43	24%
5	2%
-	
137	49%
	14%
	38%
	59%
	1%
	24%
07	24%
42.5	
43.5	(30.9, 55.8)
20	(7.61)
20	(7, 61)
11	(8,15)
39	15%
195	77%
20	8%
0	(0.0, 1.0)
	(0.0, 100.0)
209	75%
71	25%
19	9%
114	55%
71	34%
5	2%
53	29.1
58	(24, 80)
	(1, 100)
rest 1 d.p.	so may not
	70 96 2 20 89 182 137 38 105 85 1 67 43.5 20 137 38 105 85 1 67 43.5 20 11 39 195 20 71 53 53 58

Percentages have been rounded throughout to nearest 1 d.p. so may not always sum to 100%.

IOFB intraocular foreign body, *TRD* tractional retinal detachment, *RRD* rhegmatogenous retinal detachment, *VH* vitreous haemorrhage, *RAPD* relative afferent pupillary defect, *PVR* proliferative vitreoretinopathy, *IOP* intraocular pressure, *ACIOL* anterior chamber intraocular lens, *PCIOL* Posterior chamber intraocular lens, *IQR* Interquartile range, *ETDRS* Early Treatment Diabetic Retinopathy Study.

^aOther Injury includes any injuries that do not fall into any other category (e.g. freak accidents/childhood play), or where not enough information was given to categorise. For a full listing of other injury causes see supplementary file.

supplementary file. ^bOther Domestic includes all other injuries within a domestic setting not covered by Domestic Gardening, Domestic DIY or Falls. For a full listing of other injury causes see supplementary file. 3

4

More than 25% of patients in London and Scotland had an ETDRS score above zero (zero indicating very low vision of counting fingers or worse).

Seasonal variation

The dates of ocular injuries were investigated for seasonal trends. Only injuries from 2014 were included as the date of injury was not collected accurately prior to this time, resulting in 255 participants included within this analysis. The most common month for injury was April and injuries in April were consistently high across each year of recruitment. January, February, June and November had slightly fewer injuries occurring than in other months (see Fig. 1). There was however no evidence to suggest significant variation in the number of injuries by month beyond that expected by chance (p = 0.293, $\chi^2 = 13$, 11 degrees of freedom).

Workplace incidents and interpersonal violence accounted for the majority of ocular injuries and there were fluctuations in the months these injuries occurred (see Fig. 2A–E). Interpersonal violence injuries peaked in December to January and April to May, with lows in February and June. Workplace injuries occurred most often in April, closely followed by August and September, with lulls in January, February, May and October. There was no evidence to suggest cause of injury differed significantly across month of injury (p = 0.534, $\chi^2 = 97.1$, 99 degrees of freedom).

Associations between trauma and ocular history and presenting vision

Table 2 displays the regression model coefficients and 95% Cl's for the multivariable regression analyses that explore the associations between patient history, presenting clinical characteristics and presenting vision (ETDRS).

Factors associated with zero vision (an ETDRS score of 0 letters indicating very low/no vision) were visual axis corneal scar (p = 0.063), hyphaema (p = 0.018), vitreous haemorrhage (p < 0.001) and PVR (p = 0.001) (see Table 2), indicating that if the trauma causes these then very poor vision at presentation is more likely. Having hyphaema greatly increased the odds of zero/very low vision for both those with <50% hyphaema OR = 5.24 (95% CI 1.34, 20.5) and those with >50% hyphaema OR = 6.55 (95% CI 0.78, 55.2) relative to those without hyphaema. Vitreous haemorrhage with no fundal view had much higher odds of zero/very low vision in comparison to without visual haemorrhage OR = 8.45 (95% CI 3.07, 23.2), whereas visual haemorrhage with fundus had much lower odds of zero/very low vision OR = 0.29 (95% CI 0.11, 0.76) compared to without visual haemorrhage.

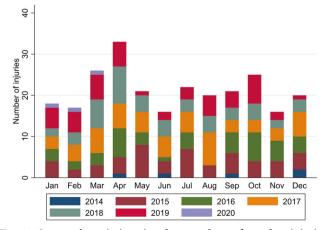


Fig. 1 Seasonal variation in the number of ocular injuries. Number of injuries shown by year (represented by different colours) and month.

For individuals who could read at least one letter on the ETDRS chart (ETDRS score >0), previous eye surgery (p = 0.001), visual axis corneal scar (p = 0.048), lens status (p = 0.128) and vitreous haemorrhaging (p = 0.189) were identified to be associated with presenting ETDRS score. The expected ETDRS score for patients who had had previous eye surgery was decreased by a factor of 0.41 times (95% Cl 0.25, 0.69) relative to no previous surgery among participants with ETDRS > 0. In comparison to a clear lens, having an anterior chamber intraocular lens (ACIOL) was associated with better vision, IRR = 2.05 (95% CI 0.75, 5.60, p = 0.159), whereas those with cataracts IRR = 0.74 (95% CI 0.48, 1.14), and aphakic IRR = 0.69 (95% CI 0.42, 1.16,) had lower ETDRS scores among participants with ETDRS > 0. The ETDRS score with vitreous haemorrhaging and no fundal view was 0.61 times (95% CI 0.33, 1.15) with no vitreous haemorrhage among participants with ETDRS > 0. The ETDRS score with visual axis corneal scar was 1.59 times higher (95% CI 1.00, 2.51) in comparison to without among participants with ETDRS > 0—however, those with visual axis corneal scare were more likely to have no vision at presentation than those without (although not statistically significantly so).

Associations with retinal detachment are shown in Table 3. Hyphaema was shown to be strongly associated with retinal detachment (p = 0.033); over 50% hyphaema was associated with higher odds of retinal detachment relative to none, OR = 3.29 (95% CI 1.32, 8.16). The location of injury also had an association (p = 0.015) with retinal detachment with injuries to scleral-posterior to muscle insertion having higher odds of retinal detachment 3.43 (95% CI 1.44, 8.19) relative to injuries extending to the cornea only.

DISCUSSION

The ASCOT study was a prospective randomised controlled trial designed to test the efficacy of intraocular and periocular triamcinolone given at the time of vitrectomy surgery in cases of open globe injury. It recruited nationally in the UK from 27 sites. The baseline data therefore represents the cases considered suitable for inclusion by the principal investigators at the individual sites and is not a comprehensive epidemiological study on open globe injuries. Nevertheless, it provides detailed data on a large cohort of patients undergoing vitrectomy surgery.

The study population were overwhelmingly (88%) male, consistent with other analyses of eye trauma epidemiology [8, 18, 19]. The median age of the study population was 43 demonstrating the difference between trauma patients and the majority of other patients undergoing intraocular surgery. There was considerable variability in presenting visual acuity. Five patients had no perception of light (NPL) vision at baseline indicating that amongst UK vitreoretinal surgeons there is some willingness to operate on patients with very limited vision. Internationally other reports have documented patients with NPL vision following penetrating trauma undergoing vitreoretinal surgery although the prognosis for these cases remains very poor [20].

There was a broad range of causal mechanisms of the injury in the study cases. The most common setting was the workplace (31%)followed by a significant number of the injuries (24%) due to interpersonal violence. In the UK, regulations relating to eye protection in the workplace are included in 1992 legislation on personal protective equipment at work, which came into effect in 1993 (recently updated in 2022 to also cover more causal employment relationships in addition to contracted employees) [21, 22]. The regulations clarify how suitable eye care should be provided to employees who may be exposed to a risk. Despite these regulations, the results of this study emphasise the need for further efforts to improve eye protection in the working environment. The majority of cases presented with a penetration (37%) or a globe rupture (40%) with smaller numbers having a perforation (4%) or an intraocular foreign body (19%).

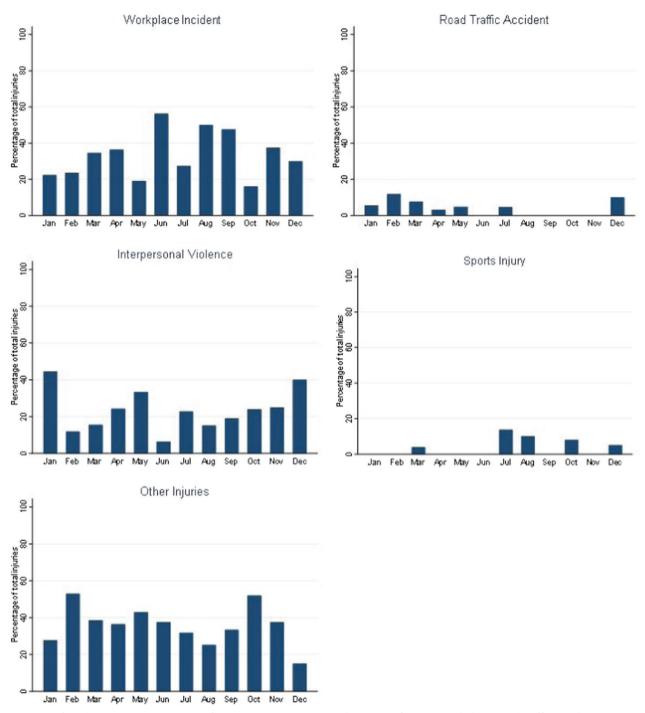


Fig. 2 Causes of ocular injuries. Proportion of injuries (as a percentage) by cause of injury (workplace, road traffic accident, interpersonal violence, sports injury and other) and month of injury.

The baseline clinical characteristics provide a useful insight into the range of pathology likely to be encountered by vitreoretinal surgeons. The initial view of the posterior segment is often compromised with 26% presenting with a visual axis corneal scar and 34% a hyphaema. Anterior segment pathology is likely with only 25% having a clear lens (35% had a cataract which may require combined surgery) and 60% having iris abnormalities. Vitreous haemorrhage was present in 66% with three-quarters of these being fundus obscuring. Approximately half of all patients had retinal detachment at the time of vitrectomy surgery with PVR already present in half of these. However, all patients in ASCOT were listed for vitrectomy, rather than being observed. The rapid development of PVR is notable, as the mean interval between injury and surgery was 20 days. The fovea was detached in over half (60%) of detachments.

In the largest retrospective review to date of open globe injuries (848 patients), Andreoli et al. report that approximately 1/3 of patients (29%) proceed to vitrectomy surgery [23]. In a subsequent report of an expanded cohort of 894 open globe injury patients, retinal detachment complicated 29% of patients. Vitreous haemorrhage was significantly associated with the presence of retinal detachment where 85% of eyes with retinal detachment

5

Table 2. Associations with presenting vision as measured by the ETDRS.

Variable	Zero-inflated negative binomial			Zero-Infla	Zero-Inflated Logistic Regression (ETDRS $=$ 0)		
	IRR ^a	95% CI	p value	OR ^b	95% CI	p value	
Previous primary repair							
No (reference)							
Yes	1.1	0.63, 1.91	0.742				
Classification							
Rupture (reference)			0.801				
Penetration	0.92	0.60, 1.43					
Perforation	0.54	0.07, 4.10					
IOFB	1.13	0.63, 2.04					
Extension zone							
Cornea (reference)			0.274				
Scleral-anterior to muscle insertion	1.37	0.92, 2.04					
Scleral-posterior to muscle insertion	1.13	0.67, 1.92					
Previous eye surgery (eye operating on)							
No (reference)			0.001				
Yes	0.41	0.25, 0.69					
Macular disease							
No (reference)			0.98				
Yes	0	0					
Other eye history							
No (reference)			0.249				
Yes	0.74	0.45, 1.23					
Visual axis corneal scar							
No (reference)			0.048				
Yes	1.59	1.0, 2.51		2.33	0.95, 5.67	0.063	
Hyphaema	110 5	110/ 210 1		2.00	01207 5107	0.000	
None (reference)			0.469			0.018	
<50%	1.14	0.60, 2.17	01105	5.24	1.34, 20.54	0.010	
>50%	0.48	0.14, 1.64		6.55	0.78, 55.24		
Lens status	0.10	0.11, 1.01		0.55	0.70, 55.21		
Clear (reference)			0.128				
Cataract	0.74	0.48, 1.14	0.120				
ACIOL	2.05	0.75, 5.60					
PCIOL	1.11	0.60, 2.06					
Aphakic	0.69	0.80, 2.08					
Vitreous haemorrhage	0.09	0.41, 1.10					
-			0.189			<0.001	
None (reference)	0.61	0.22 1.15	0.169	9 4 5	2 07 22 20	<0.001	
VH with no fundal view	0.61	0.33, 1.15		8.45	3.07, 23.20		
VH with fundus	1.11	0.79, 1.57		0.29	0.11, 0.76		
Endophtamlitis							
No (reference)		0.20 5.11	0.000				
Yes	1.4	0.39, 5.11	0.606				
Retinal status			0.205				
Attached (reference)	<i>.</i>		0.305				
TRD	0.64	0.27, 1.53					
RRD	1.26	0.85, 1.87					
Proliferative vitreoretinopathy (PVR)							
No (reference)							
Yes	1.38	0.71, 2.69	0.338	7.2	2.30, 22.61	0.001	

ACIOL anterior chamber itraocular lens, PCIOL posterior chamber introcular lens, IOFB intraocular foreign body, TRD tractional retinal detachment, RRD rhegmatogenous retinal detachment. VH vitreous haemorrhage, PVR proliferative vitreoretinopathy, ETDRS Early Treatment Diabetic Retinopathy Study. ^aIRR (Incidence rate ratio) represents the relative risk of a higher ETDRS score for each associated variable relative to the reference group for that variable. ^bOR (Odds ratio) represents the odds of ETRDS score ≤ 0 , for the associated variable relative to the reference group for that variable.

had an associated vitreous haemorrhage compared with a 32% incidence of vitreous haemorrhage in eyes without retinal detachment ($p \le 0.001$). Additionally, they reported a significant higher risk of retinal detachment in older, male patients with

poorer presenting vision and higher zone of injury (Zone 3, Scleral posterior to muscle insertion) [24].

Endophthalmitis was rare (2%) on presentation. Pre-existing eye disease was uncommon: four of the 280 patients had glaucoma

Table 3. Associations with retinal detachment.

Variable	OR ^a	95% CI	p value
Previous primary repair			
No (reference)			0.235
Yes	1.66	0.72, 3.86	
Classification			
Rupture (reference)			0.966
Penetration	0.9	0.47, 1.74	
Perforation	0.8	0.19, 3.35	
IOFB	1.08	0.40, 2.93	
Extension zone			
Cornea (reference)			0.015
Scleral-anterior to muscle insertion	1.45	0.71, 2.98	
Scleral-posterior to muscle insertion	3.43	1.44, 8.19	
Previous eye surgery (study eye)			
No (reference)			0.096
Yes	1.75	0.91, 3.37	
Macular disease			
No (reference)			
Yes	N/A		
Other eye history			
No (reference)			0.907
Yes	1.06	0.42, 2.68	
Visual axis corneal scar			
No (reference)			0.943
Yes	1.02	0.52, 2.01	
Hyphaema			
None (reference)			0.033
<50%	1.83	0.82, 4.08	
>50%	3.29	1.32, 8.16	
Lens status			
Clear (reference)			0.715
Cataract	1.1	0.51, 2.40	
ACIOL	N/A		
PCIOL	0.69	0.19, 2.47	
Aphakic	1.31	0.57, 3.01	
Vitreous haemorrhage			
None (reference)			0.117
VH with no fundal view	0.47	0.22, 1.00	
VH with fundus	0.5	0.20, 1.23	
Endophthamitis			
No (reference)			
Yes	0.43	0.04, 4.81	0.495

ACIOL anterior chamber itraocular lens, PCIOL Posterior chamber introcular lens, IOFB Intraocular foreign body. TRD tractional retinal detachment, RRD rhegmatogenous retinal detachment, VH vitreous haemorrhage.

 $^{\rm a}{\rm OR}$ (odds ratio) represents the odds of ETRDS score >0, for the associated variable relative to the reference group for that variable.

and one had had previous macular disease recorded—this may reflect the younger age range of patients sustaining open globe eye trauma.

The presenting clinical characteristics which were associated with better presenting vision were often related to ocular media

Eye

opacities. The presence of corneal scarring, cataract, hyphaema and vitreous haemorrhage conferred less good presenting vision. Eyes which had previously undergone surgery also had less good vision possibly relating to more complex pre-existing intraocular pathology. Eyes with ACIOLs had better visual acuity – this may reflect a more longstanding and stable situation prior to the need for vitreoretinal intervention, however, this should be interpreted with caution as there were only two patients with ACIOLs. The presence of PVR (and by implication retinal detachment) on presentation was also associated with less good presenting vision. PVR is associated with poor vision and poor outcomes from surgery both with and without previous ocular trauma [9, 25]. 7

The associations with retinal detachment at presentation are also notable. The zone of injury clearly influences the likelihood of retinal and vitreous involvement in the injury and as expected more posterior injuries were more likely to have retinal detachment on presentation. The association with hyphaema could potentially reflect the potential of intraocular blood to promote PVR through fibrogenic growth factors.

Recruitment for the study was centred in London, particularly Moorfields Eye Hospital which was the lead site. The patient cohort is therefore not a comprehensive study of open globe injury in the UK. To compare regions, cases were grouped together to provide cohorts of a meaningful size for analysis. The cases recruited in London were generally less severe (a greater proportion of corneal, zone 1 (cornea), injuries) than elsewhere with more ocular co-morbidities. These observations must, however, be treated with caution as they may be influenced by recruitment bias in the trial centres.

There was a trend for recruitment to be higher in spring (March and April) and October although this was not statistically significant with the patient numbers in the study. Notably workplace injuries had a trend to more common in spring and summer with interpersonal violence being more common in January and December. It is possible that a reduction in work and an increase in in social interaction results in the reversal of causation seen in December and January. Although a bias in case recruitment may influence the monthly cases numbers overall, it is unlikely to have affected the variation in injury causation seen.

CONCLUSION

The ASCOT study provides detailed clinical data on an extensive cohort of patients with open globe penetrating injuries undergoing vitreoretinal surgery. Although not a comprehensive epidemiological study, this study provides valuable insights into the spectrum of pathology encountered by vitreoretinal surgeons. We have documented the common settings for injury—in the workplace and through interpersonal violence, highlighted a trend in seasonal variation and reported the clinical presentations of the cases—predominantly penetrating or globe rupture and often with media opacities limiting the fundal view. These observations can help training and in planning the resources needed to deal with often challenging surgical cases.

SUMMARY

What was known before

- Ocular trauma is a leading cause of monocular visual impairment and blindness worldwide, affecting 55 million people every year.
- Ocular trauma is a serious health problem that has extensive, variable, physical and psychological impacts on patients and their relatives.

 The primary aim of the Adjunctive Steroid Combination in Ocular Trauma (ASCOT) trial is to assess whether, adjunctive intravitreal and sub-Tenon's triamcinolone acetonide given at the time of surgery improves visual acuity at 6 months compared with standard surgery.

What this study adds

- Common settings for open globe injury for the ASCOT cohort include the workplace and through interpersonal violence, with a trend in seasonal variation.
- There is considerable variability in presenting visual acuity for patients with open globe eye injuries undergoing vitreoretinal surgery.
- Clinical presentations of cases are predominantly penetrating or globe rupture and often with media opacities limiting the fundal view.

DATA AVAILABILITY

The datasets generated and analysed during the current study are available from the corresponding author on reasonable request.

REFERENCES

- Esmaeli B, Elner SG, Schork MA, Elner VM. Visual outcome and ocular survival after penetrating trauma. A clinicopathologic study. Ophthalmology. 1995;102:393–400.
- Négrel AD, Thylefors B. The global impact of eye injuries. Ophthalmic Epidemiol. 1998;5:143–69.
- Chua D, Wong W, Lamoureux EL, Aung T, Saw S-M, Wong TY. The prevalence and risk factors of ocular trauma: the Singapore Indian Eye Study. Ophthalmic Epidemiol. 2011;18:281–7.
- Marta A, Silva N, Correia N, Pessoa B, Ferreira N, Beirão M, et al. A 15-year retrospective epidemiologic study of ocular trauma in the north of Portugal. Eur J Ophthalmol. 2021;31:1079–84.
- Kousiouris P, Klavdianou O, Douglas KAA, Gouliopoulos N, Chatzistefanou K, Kantzanou M, et al. Role of socioeconomic status (SES) in globe injuries: a review. Clin Ophthalmol. 2022;16:25–31.
- Wong TY, Klein BEK, Klein R. The prevalence and 5-year incidence of ocular trauma: The Beaver Dam Eye Study. Ophthalmology. 2000;107:2196–202.
- Schein OD, Hibberd PL, Shingleton BJ, Kunzweiler T, Frambach DA, Seddon JM, et al. The spectrum and burden of ocular injury. Ophthalmology. 1988;95:300–5.
- 8. Morris DS, Willis S, Minassian D, Foot B, Desai P, MacEwen CJ. The incidence of serious eye injury in Scotland: a prospective study. Eye. 2014;28:34–40.
- Cardillo JA, Stout JT, LaBree L, Azen SP, Omphroy L, Cui JZ, et al. Post-traumatic proliferative vitreoretinopathy. The epidemiologic profile, onset, risk factors, and visual outcome. Ophthalmology. 1997;104:1166–73.
- Spiegel D, Nasemann J, Nawrocki J, Gabel VP. Severe ocular trauma managed with primary pars plana vitrectomy and silicone oil. Retina. 1997;17:275–85.
- 11. Framme C, Roider J. Epidemiology of open globe injuries. Klin Monatsbl Augenheilkd. 1999;215:287–93.
- 12. Mittra RA, Mieler WF. Controversies in the management of open-globe injuries involving the posterior segment. Surv Ophthalmol. 1999;44:215–25.
- Tano Y, Chandler D, Machemer R. Treatment of intraocular proliferation with intravitreal injection of triamcinolone acetonide. Am J Ophthalmol. 1980;90:810–6.
- Banerjee PJ, Cornelius VR, Phillips R, Lo JW, Bunce C, Kelly J, et al. Adjunctive intraocular and peri-ocular steroid (triamcinolone acetonide) versus standard treatment in eyes undergoing vitreoretinal surgery for open globe trauma (ASCOT): study protocol for a phase III, multi-centre, double-masked randomised controlled trial. Trials. 2016;17:339.
- 15. Lo JW, Bunce C, Charteris D, Banerjee P, Phillips R, Cornelius VR. A phase III, multicentre, double-masked randomised controlled trial of adjunctive intraocular and peri-ocular steroid (triamcinolone acetonide) versus standard treatment in eyes undergoing vitreoretinal surgery for open globe trauma (ASCOT): statistical analysis plan. Trials. 2016;17:383.
- UK Government. Regional ethnic diversity. 2020. https://www.ethnicity-facts-figures. service.gov.uk/uk-population-by-ethnicity/national-and-regional-populations/ regional-ethnic-diversity/latest?msclkid=59a55652aae111ecbb8b64c181b4b732# areas-of-england-and-wales-by-ethnicity. Accessed 23 Mar 2022.

- Government S. Ethnicity. 2021 https://www.scotlandscensus.gov.uk/censusresults/at-a-glance/ethnicity/?msclkid=711b982faae211eca6f208109d6ec2e1. Accessed 23 Mar 2022.
- Kuhn F, Morris R, Witherspoon CD, Mann L. Epidemiology of blinding trauma in the United States Eye Injury Registry. Ophthalmic Epidemiol. 2006;13:209–16.
- Desai P, MacEwen CJ, Baines P, Minassian DC. Incidence of cases of ocular trauma admitted to hospital and incidence of blinding outcome. Br J Ophthalmol. 1996;80:592–6.
- Fujikawa A, Mohamed YH, Kinoshita H, Matsumoto M, Uematsu M, Tsuiki E, et al. Visual outcomes and prognostic factors in open-globe injuries. BMC Ophthalmol. 2018;18:138.
- UK Health and Safety Commission. The personal protective equipment at work regulations 1992. UK Statutory Instruments. 1992; No. 2966. Avaialble at: https:// www.legislation.gov.uk/uksi/1992/2966/introduction/made. Accessed: 2 Sep 2022
- 22. UK Health and Safety Commission. The personal protective equipment at work (Amendment) regulations 2022. UK Statutory Instruments. 2022; No. 8.Avaialble at: https://www.legislation.gov.uk/uksi/1992/2966/introduction/made. Accessed: 2 Sep 2022.
- 23. Andreoli MT, Yiu G, Hart L, Andreoli CM. B-scan ultrasonography following open globe repair. Eye. 2014;28:381–5.
- Stryjewski TP, Andreoli CM, Eliott D. Retinal detachment after open globe injury. Ophthalmology. 2014;121:327–33.
- Patel NN, Bunce C, Asaria RH, Charteris DG. Resources involved in managing retinal detachment complicated by proliferative vitreoretinopathy. Retina. 2004;24:883–7.

ACKNOWLEDGEMENTS

Authors contributing to this paper are supported by the United Kingdom Clinical Research Collaboration-registered King's Clinical Trials Unit at King's Health Partners. We thank Beverley White-Alao and staff in the wider KCTU who supported the electronic data capture and randomisation systems and data management for the ASCOT study.

AUTHOR CONTRIBUTIONS

DC is the chief investigator for ASCOT. SC, GP, VC, CB and DC wrote the statistical analysis plan for this manuscript. SC, GP and VC conducted the statistical analysis for this manuscript. SC, GP and DC wrote the first draft of the manuscript. SC, GP, VC, PB, MZ, EC, SS, CB and DC interpreted the data and critically reviewed the manuscript. PB, VC, and CB are co-investigators for ASCOT. DC, VC, PB and CB designed the ASCOT study. ER was the ASCOT trial manager. CM and JK were involved in the trial design, trial and data management. All authors read and approved the final manuscript.

FUNDING

The ASCOT trial is funded by a project grant from the National Institute for Health Research Health Technology Assessment (HTA) programme (HTA 12/35/64). CB's post is part funded by the National Institute for Health Research (NIHR) Biomedical Research Centre at The Royal Marsden NHS Foundation Trust and the Institute of Cancer Research, London. The views expressed are those of the author(s) and not necessarily those of the NIHR or the Department of Health and Social Care. SC is funded by an NIHR advanced research fellowship (NIHR300593). The views expressed are those of the authors and not necessarily those of the NIHR or the Department of Health and Social Care.

COMPETING INTERESTS

The authors declare no competing interests.

ADDITIONAL INFORMATION

Supplementary information The online version contains supplementary material available at https://doi.org/10.1038/s41433-022-02206-z.

Correspondence and requests for materials should be addressed to Suzie Cro.

Reprints and permission information is available at http://www.nature.com/ reprints

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations. **Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http:// creativecommons.org/licenses/by/4.0/.

© The Author(s) 2022