

**Low-dose (0.01%) atropine eye drops to reduce progression of myopia in children: a multi-centre placebo-controlled randomised trial in the United Kingdom (CHAMP-UK) – study protocol.**

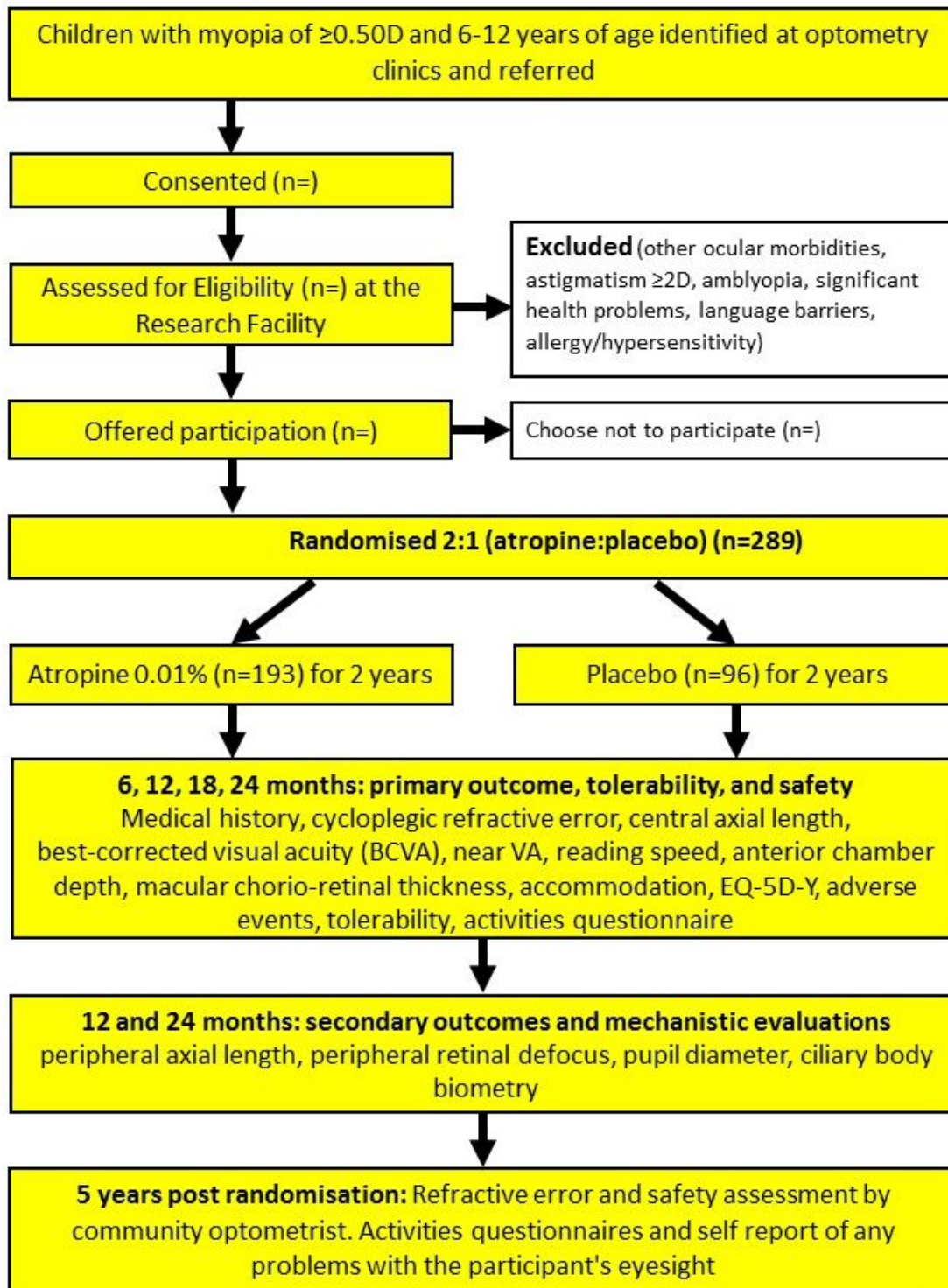
Journal:	<i>British Journal of Ophthalmology</i>
Manuscript ID	bjophthalmol-2019-314819.R1
Article Type:	Clinical science
Date Submitted by the Author:	n/a
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Keywords:	Treatment Medical, Optics and Refraction, Drugs, Clinical Trial, Child health (paediatrics)

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Figure 1.



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6 **Low-dose (0.01%) atropine eye drops to reduce progression of myopia in**  
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8 **children: a multi-centre placebo-controlled randomised trial in the United**  
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10 **Kingdom (CHAMP-UK) – study protocol.**  
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45 **Word count:** 2997

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51 **Acronym:** The Childhood Atropine for Myopia Progression in the UK study (CHAMP-UK)

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54 **Trial registration:** ISRCTN99883695, NCT03690089.  
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4 **Precis**  
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6 We report the protocol of a multicentre placebo-controlled randomised trial that will evaluate the  
7 efficacy, safety and mechanism of action of low dose atropine (0.01%) eye drops to reduce myopia  
8 progression in children with myopia.  
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Confidential: For Review Only

**Abstract**

**Background/Aims:** To report the protocol of a trial designed to evaluate the efficacy, safety and mechanism of action of low dose atropine (0.01%) eye drops for reducing progression of myopia in UK children.

**Methods:** Multicentre, double-masked, superiority, placebo-controlled, randomised trial. We will enrol children aged 6-12 years with myopia of  $-0.50$  diopters or worse in both eyes.

We will recruit 289 participants with an allocation ratio of 2:1 (193 atropine; 96 placebo) from five centres. Participants will instil one drop in each eye every day for two years and attend a research centre every six months. The vehicle and preservative will be the same in both study arms.

The primary outcome is spherical equivalent refractive error of both eyes measured by autorefractor under cycloplegia at two years (adjusted for baseline). Secondary outcomes include axial length, best corrected distance visual acuity, near visual acuity, reading speed, pupil diameter, accommodation, adverse event rates and allergic reactions, quality of life (EQ-5D-Y), and tolerability at two years. Mechanistic evaluations will include: peripheral axial length, peripheral retinal defocus, anterior chamber depth, iris colour, height and weight, activities questionnaire, ciliary body biometry and chorio-retinal thickness. Endpoints from both eyes will be pooled in combined analysis using generalised estimating equations to allow for the correlation between eyes within participant. Three years after cessation of treatment, we will also evaluate refractive error and adverse events.

**Conclusions:** CHAMP-UK will be the first randomised trial reporting outcomes of low-dose atropine eye drops for children with myopia in a UK population.

## Introduction

In Europe, myopia prevalence has risen dramatically over the last few decades. [1-6] Myopia appears to be occurring at a younger age and its severity has increased by an average of approximately 1 dioptre (D) among European-derived populations in one generation. [4, 7] In UK the proportion of myopic children has doubled in the last 50 years.[6] In the USA myopia prevalence increased from 25% to 42% in a generation.[7]

In the UK, most people with myopia have normal visual acuity when appropriately corrected, but myopia still has significant public health consequences from a variety of perspectives, educational, financial and psychological, as well as the risks of visual impairment. [8-10] Myopia is a risk factor for myopic maculopathy, retinal detachment, cataract and glaucoma in adult life, [9, 11, 12] and the risk increases with the degree of myopia. All these conditions are more challenging to treat than myopia itself, and reducing the risk for any of them requires interventions to slow myopia progression and thus decrease a child's severity of myopia in the long term rather than correct it optically with spectacles. Children with myopia also require frequent eye tests and change of spectacles that are funded primarily by taxpayers in the UK. Strategies to control progression of myopia are particularly meaningful in the context of World Health Organisation (WHO) initiatives to eliminate preventable causes of blindness.

Atropine at low concentration has been shown to be safe and effective in slowing myopia progression in children of Chinese ethnicity, [13-17] but its safety and effectiveness in European-derived populations has not been adequately assessed in a controlled trial. Therefore, the objective of the current study is to evaluate the efficacy, safety, and mechanism of action of low dose atropine (0.01%) in UK children with myopia. This paper describes the protocol of the randomised controlled trial called CHAMP-UK (The Childhood Atropine for Myopia Progression in the UK study).

## Materials and methods

This is a multicentre, randomised, double-masked, placebo-controlled, superiority trial, with 2:1 allocation of intervention and control (atropine:placebo).

*Participants:* Children will be eligible to participate in the study if they are 6-12 years of age at the time of consent, with myopia of  $-0.50$  D or greater (spherical equivalent refractive error) in both eyes, best-corrected distance visual acuity (BCDVA) 0.20 logMAR or better in both eyes, and no other significant ocular or systemic morbidities (see Table 1). Children with myopia  $\geq -10.00$  D or astigmatism  $\geq 2.00$  D in either eye will be excluded.

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**Table 1:** Inclusion and Exclusion Criteria:

### Inclusion Criteria

1. 6-12 years of age (at the time of consenting)
2. Myopia of  $-0.50$  D or greater (spherical equivalent refractive error) in both eyes
3. Best-corrected distance visual acuity (BCDVA) 0.20 logMAR or better in both eyes

### Exclusion Criteria

1. Children with other ocular morbidities\*
2. Myopia of  $-10$  D or greater in either eye
3. Astigmatism of 2 D or higher in either eye
4. Amblyopia
5. Significant health problems that can compromise the ability to attend research visits or complete the trial
6. Other factors that may compromise the ability to attend the research appointments
7. Parents or children with poor understanding of the English language
8. Children enrolled in other interventional trials
9. Allergy or hypersensitivity to atropine or excipients
10. Prior or current use of atropine or orthokeratology contact lenses

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\* *Ocular co-morbidities refer to any disease that can potentially compromised vision or that may require ocular surgery, including diabetes mellitus.*

*Setting:* Clinical research facilities from the following five academic departments of medical or optometry schools and NHS Trusts in Northern Ireland, England and Scotland:

- Wellcome Trust-Wolfson Northern Ireland Clinical Research Facility (NICRF): Belfast Health and Social Care Trust, Queen's University Belfast and Ulster University, Northern Ireland

- Department of Optometry at Glasgow Caledonian University and NHS Greater Glasgow and Clyde, Scotland



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3 - School of Optometry at Aston University and Birmingham Women and Children's NHS Foundation  
4 Trust, England  
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6 - Department of Vision and Hearing Sciences at Anglia Ruskin University and Cambridge University  
7 Hospitals NHS Hospital Trust, England  
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9 - Moorfields Eye Hospital NHS Foundation Trust, England  
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12 *Study Interventions:* The intervention group will receive preserved 0.01% atropine sulfate eye drops,  
13 administered at home once daily in both eyes for two years. The control group will receive placebo  
14 eye drops on the same dosing schedule, with the same preservative (benzalkonium chloride 0.01%  
15 w/v in sterile water) and pH. Atropine and placebo bottles will be identical, and thus participants  
16 and investigators will be masked to study group assignment.  
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22 *Outcomes:* The primary outcome is spherical equivalent refractive error (SER) (i.e. myopia severity)  
23 of both eyes after 24 months measured by autorefractor under cycloplegia, adjusted for baseline.  
24 Participants will have 1-2 drops of 1% cyclopentolate HCl instilled in each eye at least 20 minutes  
25 before autorefraction and another drop will be instilled if full cycloplegia has not been achieved.  
26 The autorefractor derives an average of five readings to produce the SER. The spherical equivalent  
27 refractive error (i.e. the spherical power plus half the cylindrical power) will be recorded for each  
28 eye.  
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34 Secondary outcomes of both eyes include the following: axial length (measured with partial coherent  
35 interferometry), best corrected distance visual acuity (unioocular and binocular), unioocular and  
36 binocular near visual acuity (ETDRS), reading speed (Wilkins), pupil diameter (by autorefractometer),  
37 accommodation (using a near target), spectacle correction power, adverse event rates and allergic  
38 reactions, quality of life (EQ-5D-Y), [18] and tolerability at two years.  
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43 The mechanism through which atropine inhibits myopia remains uncertain. Possible sites of action  
44 include the lens and ciliary muscle, the sclera, choroid and the retina. [11] A variety of tests will be  
45 done, according to their availability in the research units, to explore the effect of atropine on ocular  
46 anatomy and function. In experimental models, atropine reduces myopia and ocular elongation via a  
47 non-accommodative mechanism. [19] However, an accommodative effect cannot be ruled out in  
48 humans. [20] We will study the effect of atropine on the lens and ciliary muscle by measuring the  
49 anterior chamber depth with laser biometry and assessing changes in ciliary muscle using anterior-  
50 segment optical coherence tomography (OCT).  
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56 Altering the amount of defocus in the peripheral retina appears to be one mechanism influencing  
57 growth rate of the eye in myopia progression. [21] Whether topical atropine also has some  
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influence in the peripheral retina is unknown. Changes in peripheral eye length and off-axis measures of refraction are two ways in which the peripheral properties of the eye can be assessed. In the posterior eye, subfoveal choroid thinning is correlated with the degree of myopia. Changes in subfoveal choroidal thickness may occur in response to imposed retinal defocus. [22-24] We will study choroidal thickness using OCT. This will enable differences in choroidal thickness resulting from atropine use to be compared with normal myopic growth. We will also explore the influence of parental myopia using parents' current spectacle prescription power, and time spent on outdoor and indoor activities (Table 2).

Table 2. Diary used to record time spent indoor or outdoor.

Activity, to nearest whole hour	Monday	Tuesday	Wednesday	Thursday	Friday	Saturday	Sunday
Hours indoors: classroom, music etc							
Hours outside: sport, training, walking / cycling to or from school							
Hours reading, homework, video / computer / tablet / mobile phone use							
Hours outside: sport, training, walking / cycling to or from school							

Compliance will be assessed using electronic monitoring with a MEMS device (AARDEX Group Ltd, Switzerland). The MEMS Cap is a plastic container with a screw top in which the eye drop bottle is stored until needed for drop instillation. [25] When the top is unscrewed the device electronically records the date and time, and this is downloaded, analysed and taken as a surrogate for having administered the medication.

At five years after randomisation, we will post a questionnaire to participants' parents and ask for details of any possible complications and adverse events after trial completion. We will also request information from their children's optometrists regarding their eye health, visual acuity, and refractive error data.

*Sample size:* We anticipate that the underlying progression in the control group and the effect of atropine eye drops in a UK population will be smaller than the reported effect in Chinese populations, [17] but assuming that atropine reduces the progression of myopia by at least 40%, using  $SD=0.7$ , an intra-class correlation coefficient between the two eyes of 0.9 and a variation

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inflation factor of 1.9, we will need 97 participants in each group. Considering a dropout rate of 15%  
and that 10% of recruited children will be Chinese, we will need a total of 289 participants: 193  
atropine, 96 placebo (152 atropine, 76 placebo inflated by a variance inflation factor of 1.9) to detect  
this difference in the non-Chinese UK population with 90% power.

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*Allocation:* Randomisation will be computer-generated using a minimisation algorithm to ensure  
balanced allocation of participants across the two treatment groups, and that each participant's  
allocation is fully concealed from everyone involved in recruiting them to the trial. Minimisation will  
be by centre, ethnic background (white/non-white), and severity of myopia (less than -3.00 D in  
either eye / -3.00 D or greater in the eye with more severe myopia). The unit of randomisation will  
be the participant (not the eye). The randomisation list will be generated by sealed envelope  
(sealedenvelope.com) and group allocation will only be visible to those with administrator access in  
the trial management team in NICTU. Local researchers will access the automated randomisation  
system to obtain the kit number for each participant.

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*Adverse events/Safety reporting:* Serious adverse events (SAEs) related to participants' participation  
in the trial are reported in accordance with the guidance from The European Clinical Trials Directive  
2001/20/EC ([https://ec.europa.eu/health/human-use/clinical-trials/directive\\_en](https://ec.europa.eu/health/human-use/clinical-trials/directive_en)).

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*Timeline of procedures:* Table 3 displays the timing of the trial's outcome measurements.  
Participants will attend a research centre every six months (+/- two weeks) across the 2-year follow-  
up as illustrated in Figure 1 (see Table 3 for details). Following completion of the 2-year trial,  
participants will then again be contacted at the 5-year time point (three years after cessation of eye  
drops) to evaluate their refractive error and possible adverse events.

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*Recruitment strategy:* We will ask community optometrists and paediatric ophthalmologists to  
inform parents of children with myopia about the trial. We will aim to recruit approximately 60  
participants per centre.

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*Data collection:* The chief investigator (CI) and NICTU will provide training to site staff on trial  
processes and procedures, including the completion of the clinical research form (CRF) and data  
collection through investigator meetings and site initiation visits. All data for an individual  
participant will be collected by the local principal investigator (PI) or designee and recorded in the  
CRF for the study. Participant identification on the CRF will be through their unique trial identifier,  
allocated at the time of recruitment. Data will be collected and recorded on the CRF and

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3 questionnaires by the local PI or designee. Case report forms and questionnaires will be submitted  
4 to the NICTU in a timely manner.  
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8 Data management: Study data will be entered onto a web-based Clinical Trial Database and  
9 processed electronically as per NICTU standard operating procedures and the study specific data  
10 management plan. Data queries will be generated for site staff as required to clarify data or request  
11 missing information. All queries will be responded to and resolved within the study database. Any  
12 amended information will then be entered in the study database. All study documentation (including  
13 participant medical records) and data will be archived as per regulatory requirements and those  
14 responsible for archiving will be noted on the sponsor delegation framework.  
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21 *Statistical analysis:* The primary analysis will be performed based on the intention to treat principle.  
22 A p-value <0.05 is considered as statistically significant. Baseline characteristics will be summarised  
23 as mean and standard deviation (SD), median and inter-quartile range (IQR) or numbers and  
24 proportions (%) as appropriate, depending on the scale of measurement and distribution.  
25 Endpoints from both eyes will be pooled in combined analysis using generalised estimating  
26 equations to allow for the correlation between eyes within participant. Difference in the myopia  
27 progression and other continuous outcomes between the atropine and control groups will also be  
28 tested for significance using independent t-test. Analysis of covariance will be performed to adjust  
29 for baseline characteristics and other covariates. Fisher's exact test will be used to test the  
30 difference in the proportions between the groups for the categorical variables. Exploratory subgroup  
31 analyses will be performed on the primary outcome using 99% confidence intervals and interaction  
32 terms (treatment group by subgroup) for the following subgroups: age (6-9 and 10-12 years at  
33 randomisation), ethnic background (white versus non-white), and severity of myopia (less than -3D  
34 in either eye versus -3D or greater myopia). Sensitivity analyses will assess the impact of missing  
35 data for the primary outcome by imputing extreme values (lowest and highest). A detailed Statistical  
36 Analysis Plan, including exploratory analyses, will be completed before the final analysis is started.  
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49 *Monitoring:* On-site monitoring will be an on-going activity from the time of initiation until trial  
50 closeout and will comply with the principles of Good Clinical Practice (GCP). On-site monitoring visits  
51 during the trial will check the accuracy of entries on CRFs against the source documents, the  
52 adherence to the protocol, study procedures and GCP. The local PI or designee will ensure that  
53 access to all trial related documents including source documents (to confirm their consistency with  
54 CRF entries) are available during monitoring visits. The extent of source data verification will be  
55 documented in the monitoring plan.  
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4 *Ethics and governance:* The trial will comply with the principles of GCP, the requirements and  
5 standards set out by the EU Directive 2001/20/EC and the applicable regulatory requirements in the  
6 UK, the Medicines for Human Use (Clinical Trials) Regulations 2004 and subsequent amendments  
7 and the UK Policy Framework for Health and Social Care Research. CHAMP-UK has been reviewed  
8 and approved by a Research Ethics Committee (18/NI/0164). Local NHS Research and Development  
9 (R&D) approvals will be obtained prior to commencing the trial at the participating sites. An  
10 independent data and safety monitoring committee will oversee the trial. A Clinical Trial  
11 Authorisation was obtained from the Medicines for Human Use Regulatory Authority (MHRA) before  
12 the start of the trial.  
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21 *Protocol Compliance:* A protocol deviation is defined as an incident that deviates from the normal  
22 expectation of a particular part of the trial process. Any deviations from the protocol will be fully  
23 documented on the protocol deviation form in the CRF.  
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26 A serious breach is defined as a deviation from the trial protocol or GCP which is likely to effect to a  
27 significant degree:

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29 (a) The safety or physical or mental integrity of the subjects of the trial; or  
30 (b) The scientific value of the trial  
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32 Protocol compliance will be monitored by the NICTU.  
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36 *Protocol Amendments:* Changes to the protocol will require regulatory authority/ethics committee  
37 approval/favourable opinion prior to implementation.  
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41 *Participant Confidentiality:* All study reports and communication regarding the study will identify the  
42 participants by the assigned unique trial identifier only. Computers where information will be stored  
43 will be password-protected. Participant confidentiality will be maintained at every stage and  
44 identifying data will not be made publicly available to the extent permitted by the applicable laws  
45 and regulations.  
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51 *Post-trial Care:* Administration of study eye drops will stop after 24 months of trial participation. At  
52 the end of the trial, participants' myopia will be managed in accordance with standard clinical  
53 practice.  
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58 *Dissemination Policy:* The final study data report will be provided by the Trial Statistician. It is  
59 anticipated that the study findings will be published in peer reviewed journals and these articles will  
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3 be led by the CI. The CI will secure a searchable compendium of these publications and make the  
4 results readily accessible to the public and healthcare professionals. In addition, study findings may  
5 be presented at both national and international meetings and to appropriate patient groups.  
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9 *Data Sharing Statement:* Requests for data sharing will be reviewed on a case by case basis by the  
10 CI and Trial Management Group. We will share trial data with the CI of the MOSAIC trial  
11 (ISRCTN36732601) and the WA ATOM trial (ACTRN12617000598381) to facilitate prospective  
12 individual participant data meta-analysis once the results of the CHAMP-UK trial are accepted for  
13 publication. MOSAIC and WA-ATOM are placebo-controlled trials evaluating 0.01% atropine eye  
14 drops. WA-ATOM will enrol 150 children aged 6-16 years with progressive myopia. MOSAIC will  
15 enrol 250 children aged 6-16 years with progressive myopia (phase 1). All participants initially  
16 assigned to the placebo (n=83) crossover to the intervention arm of the study for Phase 2, and  
17 from month 24 to 36, instil 0.01% atropine eye drops in both eyes once nightly.  
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26 *Funder:* The trial is funded by a grant awarded by NIHR Efficacy and Mechanism Evaluation (project  
27 15/48/59). CHAMP-UK is registered (ISRCTN99883695, NCT03690089). The Belfast Health and Social  
28 Care Trust sponsors the trial and provides the necessary trial insurance.  
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### 33 **Discussion**

34 Myopia typically starts to develop in childhood and although the vision can be corrected with  
35 glasses, contact lenses or surgery, myopic eyes have an increased risk of developing co-morbidities  
36 such as glaucoma, cataract, retinal detachment, and choroidal neovascularisation at the macula. [9,  
37 11] Importantly, the risks of associated co-morbidity and visual loss are associated with the degree  
38 of myopia, and cannot be reduced with optical correction alone. Myopia is more prevalent in East  
39 Asia. Recent epidemiological studies show increasing rates among adolescents in European  
40 populations and suggest myopia is occurring at an earlier age than in previous generations. [3-5]  
41 Myopia usually progresses faster at younger ages, but myopia onset, progression, and stabilisation  
42 vary widely among individuals and are influenced by a range of variables including environment,  
43 lifestyle, parental refractive history and ethnicity. [4, 26]  
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51 A number of interventions to reduce the progression of myopia have been investigated. Multifocal  
52 lenses and under-correction of myopic refractive error have at best a weak effect on myopia  
53 correction. While orthokeratology and peripheral defocus contact lenses may have some effect on  
54 axial length progression, atropine, even at low doses, appears to be the most effective intervention  
55 in Asian children to reduce progression of myopic refractive error. [27,28] Low dose atropine is now  
56 widely used in some East Asian countries for treating children with myopia, but has not been tested  
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3 in European populations.

4 Atropine is an anti-cholinergic agent that is relatively selective for muscarinic receptors. Topical use  
5 of high-concentration atropine (0.5 to 1.0%) causes pupil dilatation by blocking the muscarinic  
6 receptors in the pupillary sphincter musculature and reduces or paralyzes contraction of the ciliary  
7 muscle. Both of these result in adverse effects, e.g., photophobia and blurred near (reading) vision  
8 that are highly undesirable in a school-age population. Additionally, cessation of topical high-  
9 concentration atropine for myopia control has been associated with rapid myopia progression  
10 towards original, un-treated levels (“myopia rebound”). [15]  
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12 The effectiveness of low dose atropine (three different concentrations) in children of Chinese race  
13 has been evaluated. [13-16] The lowest tested dose of 0.01% was associated with better tolerability  
14 and efficacy, with minimum rebound effect. [14] A systematic review and network meta-analysis  
15 has confirmed muscarinic antagonists as the most effective interventions for myopia control in  
16 children of Chinese ethnicity. [27] However, there is limited evidence from European populations on  
17 atropine effectiveness in controlling myopia progression. We will also evaluate the possible  
18 mechanisms of action of atropine, and gather information regarding central and peripheral axial  
19 length, accommodation, lens position, ciliary body biometry, chorio-retinal thickness and daily  
20 activities.  
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3 **Acknowledgements**

4 **Funder:** NIHR Efficacy and Mechanism Evaluation (project 15/48/59).

5  
6 Prof Congdon is supported by the Ulverscroft Foundation (UK)

7  
8 **Trial sponsor:** Belfast Health and Social Care Trust.

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## References

1. Rudnicka AR, Kapetanakis VV, Wathern AK, et al. Global variations and time trends in the prevalence of childhood myopia, a systematic review and quantitative meta-analysis: implications for aetiology and early prevention. *Br J Ophthalmol*. 2016;100:882-90
2. Breslin KM, O'Donoghue L, Saunders KJ. A prospective study of spherical refractive error and ocular components among Northern Irish schoolchildren (the NICER study). *Invest Ophthalmol Vis Sci*. 2013;54:4843-50.
3. Williams KM, Hysi PG, Nag A, et al. Age of myopia onset in a British population-based twin cohort. *Ophthalmic Physiol Opt*. 2013;33:339-45
4. Williams KM, Bertelsen G, Cumberland P, et al; European Eye Epidemiology (E(3)) Consortium. Increasing prevalence of myopia in Europe and the impact of education. *Ophthalmol*. 2015;122:1489-97.
5. Williams KM, Verhoeven VJ, Cumberland P, et al. Prevalence of refractive error in Europe: the European Eye Epidemiology (E(3)) Consortium. *Eur J Epidemiol*. 2015;30:305-15.
6. McCullough SJ, O'Donoghue L, Saunders KJ. Six year refractive change among White children and young adults: evidence for significant increase in myopia among White UK children. *PLoS ONE*. 2016; e0146332.
7. Vitale S, Ellwein L, Cotch MF, et al. Prevalence of refractive error in the United States, 1999-2004. *Arch Ophthalmol*. 2008;126:1111-9
8. Ma X, Zhou Z, Yi H, et al. Effect of providing free glasses on children's educational outcomes in China: cluster randomized controlled trial. *BMJ*. 2014;349:g5740
9. Zheng YF, Pan CW, Chay J, et al. The economic cost of myopia in adults aged over 40 years in Singapore. *Invest Ophthalmol Vis Sci*. 2013;54:7532-7
10. Tideman JW, Snabel MC, Tedja MS, et al. Association of axial length with risk of uncorrectable visual impairment for Europeans with myopia. *JAMA Ophthalmol*. 2016;134:1355-63.
11. Flitcroft DI. The complex interactions of retinal, optical and environmental factors in myopia aetiology. *Prog Retin Eye Res*. 2012;31:622-60.
12. Marcus MW, de Vries MM, Junoy Montolio FG, et al. Myopia as a risk factor for open-angle glaucoma: a systematic review and meta-analysis. *Ophthalmol*. 2011;118:1989-94.
13. Chia A, Chua WH, Cheung YB, et al. Atropine for the treatment of childhood myopia: safety and efficacy of 0.5%, 0.1%, and 0.01% doses (Atropine for the Treatment of Myopia 2). *Ophthalmol*. 2012;119:347-54.
14. Chia A, Chua WH, Wen L, et al. Atropine for the treatment of childhood myopia: changes after stopping atropine 0.01%, 0.1% and 0.5%. *Am J Ophthalmol*. 2014;157:451-7.
15. Chia A, Lu QS, Tan D. Five-year clinical trial on atropine for the treatment of myopia 2: Myopia control with atropine 0.01% eyedrops. *Ophthalmol*. 2016;123:391-9.
16. Yam JC, Jiang Y, Tang SM, et al. Low-Concentration Atropine for Myopia Progression (LAMP) study: A randomized, double-blinded, placebo-controlled Trial of 0.05%, 0.025%, and 0.01% atropine eye drops in myopia control. *Ophthalmol*. 2019;126:113-24
17. Li SM, Wu SS, Kang MT, et al. Atropine slows myopia progression more in Asian than white children by meta-analysis. *Optom Vis Sci*. 2014;91:342-50.
18. Wille N, Badia X, Bonsel G, Burström K, Cavrini G, Devlin N, Egmar AC, Greiner W, Gusi N, Herdman M, Jelsma J, Kind P, Scalone L, Ravens-Sieberer U. Development of the EQ-5D-Y: a child-friendly version of the EQ-5D. *Qual Life Res*. 2010;19:875-86.
19. McBrien NA, Moghaddam HO, Reeder AP. Atropine reduces experimental myopia and eye enlargement via a nonaccommodative mechanism. *Invest Ophthalmol Vis Sci*. 1993;34:205-15.
20. Schmid KL, Strang NC. Differences in the accommodation stimulus response curves of adult myopes and emmetropes: a summary and update. *Ophthalmic Physiol Opt*. 2015;35:613-21.

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21. Smith EL III, Hung LF, Huang J. Relative peripheral hyperopic defocus alters central refractive development in infant monkeys. *Vision Res.* 2009;49:2386–92
22. Chiang ST, Phillips JR, Backhouse S. Effect of retinal image defocus on the thickness of the human choroid. *Ophthalmic Physiol Opt.* 2015;35:405-13
23. Read SA, Collins MJ, Vincent SJ, et al. Choroidal thickness in myopic and nonmyopic children assessed with enhanced depth imaging optical coherence tomography. *Invest Ophthalmol Vis Sci.* 2013;54:7578-86.
24. Read, Scott A., Alonso-Caneiro D, Vincent SJ, et al. Longitudinal changes in choroidal thickness and eye growth in childhood. *Investigative Ophthalmology and Visual Science* 2015;56:3103-12.
25. Boland MV, Chang DS, Frazier T, et al. Electronic monitoring to assess adherence with once-daily glaucoma medications and risk factors for nonadherence: the automated dosing reminder study. *JAMA Ophthalmol.* 2014;132:838-44.
26. Donovan L, Sankaridurg P, Ho A, et al. Myopia progression in Chinese children is slower in summer than in winter. *Optom Vis Sci.* 2012;89:1196-202.
27. Walline JJ, Lindsley K, Vedula SS, et al. Interventions to slow progression of myopia in children. *Cochrane Database Syst Rev.* 2011;(12):CD004916.
28. Huang J, Wen D, Wang Q, et al. Efficacy comparison of 16 interventions for myopia control in children: A network meta-analysis. *Ophthalmol.* 2016;123:697-708.

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4 **Legends**  
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6 **Figure 1:** Study flow chart with outcome timeline.  
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**Table 3. Schedule of Assessments**

	<b>Baseline</b>	<b>6 months</b>	<b>12 months</b>	<b>18 months</b>	<b>24 months</b>
Adverse events		✓	✓	✓	✓
Tolerability		✓	✓	✓	✓
EQ-5D-Y questionnaire	✓	✓	✓	✓	✓
Activities questionnaire- to be sent home with participant for completion	✓	✓	✓	✓	✓
Best corrected VA (logMAR ETDRS)	✓	✓	✓	✓	✓
Near VA (near logMAR ETDRS)	✓	✓	✓	✓	✓
Iris colour	✓				
Reading speed (Wilkins Rate of Reading Test)	✓	✓	✓	✓	✓
Pupil diameter prior to cycloplegia (autorefractor)	✓	✓	✓	✓	✓
Accommodation (autorefractor)	✓	✓	✓	✓	✓
Peripheral retinal defocus (autorefractor)	✓		✓		✓
Anterior chamber depth (laser biometer)	✓	✓	✓	✓	✓
Cycloplegic refractive error (autorefractor)	✓	✓	✓	✓	✓
Ciliary body biometry (AS-OCT)*	✓		✓		✓
Central axial length (laser biometer)	✓	✓	✓	✓	✓
Peripheral axial length (laser biometer)	✓		✓		✓
Chorio-retinal thickness (SD-OCT)	✓	✓	✓	✓	✓

\*: if instrumentation is available

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