## 1 Appendix C: Review Protocols

|   | Details   |
|---|---|
| Review question   | What information do people with cataracts and their carers find useful, and what format (for example written or verbal) do they prefer it to be provided in?  |
| Objectives  | To determine the information needs of people at diagnosis of cataracts and their carers, in order to inform the content, utility and applicability of literature (verbal or written) on cataracts.  |
| Type of review  | Qualitative   |
| Language  | English   |
| Study design  | Interviews, surveys, mixed-methods designs  |
| Status  | Articles published from 2000 onwards  |
| Population  | Adults (18 years and over) diagnosed with non-trauma related cataracts or their carers  |
| Intervention  | Education, information booklet/leaflets, videos   |
| Comparator  | Not relevant  |
| Outcomes  | Themes surrounding patients' or carers' educational or informational needs for example, information on prognosis, self-management, treatment options  |
| Other criteria for inclusion/exclusion of studies           | <ul> <li>Include studies that used qualitative methods for data collection (including focus<br/>group and individual interviews, observation, and document analysis) and qualitative<br/>methods for data analysis (including thematic analysis or any other appropriate<br/>qualitative analysis method that enabled analysis of text and observations and<br/>narrative presentation of findings).</li> </ul> |
|   | <ul> <li>Exclude non-qualitative research and studies for example, narrative reviews,<br/>commentaries, editorials/letters, opinion pieces, case studies/reports.</li> </ul>  |
|   | <ul> <li>Exclude studies that collected data using qualitative methods but<br/>analysed/presented the data using only quantitative methods.</li> </ul>  |
| Review strategies (data extraction, quality                 | The quality of included papers will be assessed using appropriate study design specific checklists.   |
| assessment, data analysis)                                  | Data will be analysed qualitatively (thematic synthesis methods) and overall confidence in the evidence for each outcome will be assessed using the CERQual tool.   |
| Subgroup analyses (treatment effect modifiers)              | None specified  |
| Baseline characteristics to be extracted in evidence tables | None specified  |

|  | Details   |
|--|---|
| Review question  | What are the indicators for referral for cataract surgery?  |
| Objectives   | To identify indicators for referral for cataract surgery by optometrists/general practitioners  |
| Type of review   | Prognostic  |
| Language   | English   |
| Study design   | Studies of prioritisation criteria/appropriateness frameworks/scores including validation studies, surveys  |
| Status   | Articles published from 2000 onwards  The guideline committee agreed that the search date should start from 2000 which coincides with the publication of 'Action on Cataracts' that provided guidance on the referral criteria for cataract surgery in the UK. This impacted upon waiting times and patients' expectations. |
| Population   | Studies that included indicators for referral for phacoemulsification cataract surgery in adults (18 years and older) with non-trauma related cataracts   |
| Intervention   | Prioritisation criteria/appropriateness frameworks/scores/referral policies   |
| Comparator   | Not relevant  |
| Outcomes   | <ul> <li>Indicators for referral for cataract surgery</li> <li>Conversion rate i.e. proportion of people referred and went on to have cataract surgery</li> <li>Patient reported outcome measures (PROMs)</li> <li>Health-related quality of life</li> <li>Resource use and cost</li> </ul>                                 |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Exclude: narrative reviews, commentaries, editorials/letters, opinion pieces, case studies/reports</li> <li>Exclude: non-OECD countries</li> </ul>   |
| Review strategies (data extraction, quality assessment, data analysis) | The quality of included papers will be assessed using appropriate study design specific checklists. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.   |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of different techniques for undertaking biometry?   |
| Objectives   | To compare the effectiveness of:  |
|  | ultrasound biometry (immersion and contact) and optical biometry to measure axial length  |
|  | keratometry (manual and automated) and topography to measure corneal curvature  |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials (RCTs)   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing biometry prior to phacoemulsification cataract surgery with intraocular lens (IOL) implantation   |
| Interventions  | Ultrasound biometry (axial length)  |
|  | <ul> <li>Immersion ultrasound. Examples: immersion A-scan, ultrasound A-scan, A-scan<br/>ultrasonography (Canon KU-1 IOL measurer), immersion B-guided</li> </ul>   |
|  | <ul> <li>Contact/applanation ultrasound (contact A-mode). Examples: Grieshaber Biometric<br/>System, VPLUS A/B scanner</li> </ul>   |
|  | Keratometry (corneal curvature)   |
|  | Manual  |
|  | Automated  Framples: IQL Mester, sutekerstemeter/Tenson KB, 7100, partial scherence.  |
|  | Examples: IOL Master, autokeratometer/Topcon KR- 7100, partial coherence interferometry keratometer, videokeratography  |
| Comparators  | Optical biometry (axial length)   |
|  | <ul> <li>Examples: partial coherence laser interferometry (optical or ocular) coherence<br/>biometry, laser Doppler interferometry, IOL Master (Carl Zeiss), LENSTAR LS 900,<br/>optical low-coherence reflectometry (OLCR) optical biometer, laser interference<br/>biometry</li> </ul>                |
|  | Topography (corneal curvature)  |
|  | Examples: Pentacam Scheimpflug, Orbscan Topography System   |
| Outcomes   | Deviation from predicted refractive outcome expressed as a spherical equivalent   |
|  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Exclude: narrative reviews, case studies/reports/series, reliability studies, diagnostic<br/>accuracy studies, non-comparative studies. Studies on healthy eyes/people, animals,<br/>other ocular conditions besides cataracts</li> </ul>  |
|  | <ul> <li>Exclude: combination surgical procedures i.e. cataract surgery in tandem with other<br/>surgical procedures e.g. phacotrabeculectomy, canaloplasty, Descemet's stripping<br/>automated endothelial keratoplasty, keratorefractive</li> </ul>   |
|  | • Exclude: studies comparing biometry techniques and no biometry only, standard care that is not specified or clinical prediction   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | People with high myopia (different axial lengths)   |
| Baseline characteristics to be extracted in evidence tables            | Axial length  |

|  | Details   |
|--|---|
| Review question  | What are the most appropriate formulae to optimise intraocular lens biometry calculation?   |
| Objectives   | To determine the most appropriate formulae to optimise intraocular lens biometry calculation in adults undergoing phacoemulsification cataract surgery.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials (RCTs)   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing biometry prior to phacoemulsification cataract surgery with intraocular lens implantation   |
| Interventions  | Formulae used in intraocular lens biometry calculations   |
|  | • Examples: Hoffer Q, Sanders/Retzlaff/Kraff (SRK/T), Holladay II, Olsen, Colebrander, Haigis   |
| Comparators  | All formulae vs. each other   |
| Outcomes   | Deviation from predicted refractive outcome expressed as a spherical equivalent   |
|  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | Exclude: SRK I, SRK II, Binkhorst II, Holladay I as these are no longer clinically relevant   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | <ul> <li>People with prior history of corneal refractive surgery e.g. LASIK, LASEK, radial keratotomy or photorefractive keratectomy</li> <li>Axial length (linked to specific formulae)</li> </ul>   |
|  | Piggy back lenses i.e. 2 intraocular lenses inserted instead of 1   |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of strategies used to select intraocular lens constants in order to optimise biometry calculation?  |
| Objectives   | To determine the effectiveness of different strategies used to select intraocular lens constants in order to optimise biometry calculation.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials (RCTs)   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing biometry prior to phacoemulsification cataract surgery with intraocular lens implantation   |
| Interventions  | Optimisation methods of IOL constants   |
|  | <ul> <li>Examples: surgeon-specific lens constants, axial length-specific lens constants,<br/>keratometry-specific lens constants</li> </ul>  |
| Comparators  | Different optimisation methods vs. each other   |
| Outcomes   | <ul><li>Deviation from predicted refractive outcome expressed as a spherical equivalent</li><li>Resource use and cost</li></ul>   |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | Axial length measurement i.e. different biometry techniques   |

|  | Details   |
|--|---|
| Review question  | What other factors should be considered such as, who should undertake biometry and when should preoperative biometry be assessed?   |
| Objectives   | To identify other factors that should be considered to minimise the risk of biometry and postoperative refractive errors.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Any   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing biometry prior to phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Who should undertake biometry   |
|  | When should preoperative biometry be assessed   |
|  | Second eye prediction refinement  |
| Comparator   | Not relevant  |
| Outcomes   | <ul> <li>Deviation from predicted refractive outcome expressed as a spherical equivalent</li> <li>Resource use and cost</li> </ul>  |
| Other criteria for   | Exclude: letters, editorials, commentaries, case reports  |
| inclusion/exclusion of studies   | <ul> <li>Exclude: combination surgical procedures i.e. cataract surgery in tandem with other<br/>surgical procedures e.g. trabeculectomy, canaloplasty, Descemet's stripping<br/>automated endothelial keratoplasty, keratorefractive; studies that do not specify the<br/>type of cataract surgery</li> </ul>                        |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE and/or CERQual as appropriate. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What are the procedural causes of wrong lens implant errors?  |
| Objectives   | To determine the procedural causes of wrong lens implant errors   |
| Type of review   | Qualitative   |
| Language   | English   |
| Study design   | Any   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Wrong lens implant errors   |
| Comparator   | Not relevant  |
| Outcomes   | Procedural causes of wrong lens implant errors  |
|  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included studies will be assessed using relevant risk of bias tools. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE and/or CERQual as appropriate. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What strategies should be adopted to reduce the risk of wrong lens implant errors?  |
| Objectives   | To identify strategies to minimise the risk of wrong lens implant errors  |
| Type of review   | Qualitative   |
| Language   | English   |
| Study design   | Any   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Strategies to minimise risk of wrong lens implant errors e.g. surgical checklists   |
| Comparator   | Not relevant  |
| Outcomes   | Wrong lens implant error rates  |
|  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included studies will be assessed using relevant risk of bias tools. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE and/or CERQual as appropriate. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of risk stratification techniques to reduce surgical complications?   |
| Objectives   | To determine the effectiveness of preoperative risk stratification techniques in reducing surgical complications and errors   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | <ul> <li>Prognostic validation studies</li> <li>Randomised controlled trials (RCTs)</li> <li>Observational studies</li> </ul>   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Preoperative risk stratification systems  |
| Comparator   | No preoperative risk stratification system  |
| Outcomes   | <ul> <li>Surgical complications rates e.g. dropped nucleus/posteriorly dislocated crystalline<br/>lenses, pseudophakic bullous keratopathy/endothelial cell loss, posterior capsule<br/>rupture/vitreous loss/prolapse, retinal detachment, endophthalmitis, posterior and<br/>anterior capsular tears, conversion to manual extracapsular cataract extraction,<br/>postoperative refractive astigmatism, suprachoroidal haemorrhage, chronic macular<br/>oedema</li> </ul> |
|  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Exclude: studies that examine the risk of cataracts following other ocular surgical<br/>procedures e.g. trabeculectomy; risk factors for developing cataracts; risk of<br/>complications from procedures/types of device, studies on procedural safety<br/>surgical checklists e.g. WHO, case reports/case studies</li> </ul>  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included studies will be assessed using relevant risk of bias tools. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE and/or CERQual as appropriate.   |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details  |
|--|--|
| Review question  | What are the risk factors associated with increased surgical complications in cataract surgery?  |
| Objectives   | To determine the risk factors that are associated with an increase in surgical complications and errors  |
| Type of review   | Prognostic   |
| Language   | English  |
| Study design   | <ul><li> Prognostic studies</li><li> Observational studies</li></ul>   |
| Status   | No date restrictions   |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation  |
| Intervention   | Not relevant   |
| Comparator   | Not relevant   |
| Outcomes   | Surgical complications rates e.g. dropped nucleus/posteriorly dislocated crystalline lenses, pseudophakic bullous keratopathy/endothelial cell loss, posterior capsule rupture/vitreous loss/prolapse, retinal detachment, endophthalmitis, posterior and anterior capsular tears, conversion to manual extracapsular cataract extraction, postoperative refractive astigmatism, suprachoroidal haemorrhage, chronic macular oedema  Resource use and cost |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Exclude: studies that examine the risk of cataracts following other ocular surgical<br/>procedures e.g. trabeculectomy; risk factors for developing cataracts; risk of<br/>complications from procedures/types of device, studies on procedural safety<br/>surgical checklists e.g. WHO, case reports/case studies</li> </ul>   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included studies will be assessed using relevant risk of bias tools. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE and/or CERQual as appropriate.  |
| Subgroup analyses (treatment effect modifiers)                         | None specified   |
| Baseline characteristics to be extracted in evidence tables            | None specified   |

|  | Details  |
|--|--|
| Review question  | What is the effectiveness of laser-assisted phacoemulsification cataract surgery compared with standard ultrasound phacoemulsification cataract surgery?   |
| Objectives   | To compare the effectiveness of laser-assisted phacoemulsification cataract surgery with standard ultrasound phacoemulsification cataract surgery.   |
| Type of review   | Intervention   |
| Language   | Any  |
| Study design   | Randomised controlled trials (RCTs); randomisation of individuals or eyes. Single eye studies and studies which include both eyes.   |
| Status   | No date restrictions   |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery and posterior chamber intraocular lens (IOL) implantation   |
| Intervention   | Laser-assisted phacoemulsification cataract surgery (e.g. Catalys, LENSAR, LenSx, Victus)  |
| Comparator   | Standard ultrasound phacoemulsification cataract surgery   |
| Outcomes   | <ul> <li>Visual acuity (aided and unaided; best corrected visual acuity [BCVA]): report mean at longest follow-up timepoint</li> <li>Intraoperative complications</li> <li>Resource use and costs (e.g. total duration of procedure, number of operating rooms/practitioners)</li> <li>Postoperative complications</li> <li>Refractive outcome (deviations e.g. deviation from the predictive refractive outcome)</li> <li>Refractive outcome (predictions)</li> <li>Patient satisfaction</li> </ul> |
| Other criteria for inclusion/exclusion of studies                      | None   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.  |
| Subgroup analyses (treatment effect modifiers)                         | <ul> <li>Type of intraocular lenses e.g. standard monofocal lenses</li> <li>Surgical technique: grid fragmentation pattern specific cataract surgery</li> </ul>  |
| Baseline characteristics to be extracted in evidence tables            | None specified   |

|   | Details   |
|---|---|
| Review question   | What is the optimal type and administration of anaesthesia for cataract surgery?  |
| Objectives  | To determine the optimal type and administration of anaesthesia for phacoemulsification cataract surgery  |
| Type of review  | Intervention  |
| Language  | English   |
| Study design  | Randomised controlled trials (RCTs)   |
| Status  | No date restrictions  |
| Population  | Adults (18 years and over) undergoing phacoemulsification cataract surgery for non-trauma related cataracts and intraocular lens (IOL) implantation   |
| Intervention  | Methods:  • peribulbar/periocular block  • retrobulbar block  • sub-Tenon's anaesthesia  • topical (drops) ± intracameral (diluted with saline)  Drugs:  • Lidocaine/xylocaine  • Bupivacaine   |
| Comparator  | <ul> <li>Different methods vs. each other</li> <li>Different drugs vs. each other</li> <li>Warming of drug vs. no warming of drug</li> </ul>  |
| Outcomes  | <ul> <li>Intraoperative pain</li> <li>Pain on administration of anaesthesia</li> <li>Surgical complication rates</li> <li>Anaesthetic-related complications</li> <li>Patient satisfaction</li> <li>Resource use and costs</li> </ul>  |
| Other criteria for inclusion/exclusion of studies           | <ul> <li>Exclude: studies on general ophthalmic conditions</li> <li>Exclude: letters, editorials, commentaries, narrative reviews, observational studies, case reports</li> <li>Exclude: combination surgical procedures i.e. cataract surgery in tandem with other surgical procedures e.g. trabeculectomy, canaloplasty, Descemet's stripping automated endothelial keratoplasty, keratorefractive; studies that do not specify the type of cataract surgery</li> <li>Exclude: 2-chloroprocaine, articaine (not licensed for ophthalmic use in UK)</li> <li>Exclude: studies on concomitant intravenous sedation/preoperative anxiolytic regimens as this may mask pain perception and will also be covered in separate sedation review question e.g. benzodiazepines (bromazepam, alprazolam, diazepam), sedatives (propofol or remifentanil)</li> <li>Exclude: studies on new viscoelastic substances as this will be covered in the hyaluronidase review question (e.g. sodium hyaluronate 1.5% and lidocaine 1%)</li> </ul> |
| Review strategies   | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.   |
| Subgroup analyses (treatment effect modifiers)              | <ul><li>Axial lengths</li><li>People on anticoagulants</li></ul>  |
| Baseline characteristics to be extracted in evidence tables | None specified  |

|   | Details   |
|---|---|
| Review question   | What is the effectiveness of sedation as an adjunct to local anaesthesia during cataract surgery?   |
| Objectives  | To determine the effectiveness of sedation as an adjunct to local anaesthesia during phacoemulsification cataract surgery.  |
| Type of review  | Intervention  |
| Language  | English   |
| Study design  | Randomised controlled trials (RCTs)   |
| Status  | No date restrictions  |
| Population  | Adults (18 years and over) undergoing phacoemulsification cataract surgery for non-trauma related cataracts and intraocular lens (IOL) implantation   |
| Intervention  | Sedation (midazolam, fentanyl, propofol)  |
| Comparator  | No sedation   |
| Outcomes  | <ul> <li>Intraoperative pain</li> <li>Pain on administration of anaesthesia</li> <li>Surgical complication rates</li> <li>Anaesthetic-related complications</li> <li>Patient satisfaction</li> <li>Resource use and costs</li> </ul>  |
| Other criteria for inclusion/exclusion of studies           | <ul> <li>Exclude: studies on general ophthalmic conditions</li> <li>Exclude: letters, editorials, commentaries, narrative reviews, observational studies, case reports</li> <li>Exclude: combination surgical procedures i.e. cataract surgery in tandem with other surgical procedures e.g. trabeculectomy, canaloplasty, Descemet's stripping automated endothelial keratoplasty, keratorefractive; studies that do not specify the type of cataract surgery</li> <li>Exclude: studies on different types of sedation only</li> </ul> |
| Review strategies   | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.   |
| Subgroup analyses (treatment effect modifiers)              | Method of anaesthetic administration  |
| Baseline characteristics to be extracted in evidence tables | None specified  |

|   | Details   |
|---|---|
| Review question   | What is the effectiveness of hyaluronidase as an adjunct to local anaesthesia during cataract surgery?  |
| Objectives  | To determine the effectiveness of hyaluronidase as an adjunct to local anaesthesia during phacoemulsification cataract surgery.   |
| Type of review  | Intervention  |
| Language  | English   |
| Study design  | Randomised controlled trials (RCTs)   |
| Status  | No date restrictions  |
| Population  | Adults (18 years and over) undergoing phacoemulsification cataract surgery for non-trauma related cataracts and intraocular lens (IOL) implantation   |
| Intervention  | Hyaluronidase/hyalase/hyaluronic acid   |
| Comparator  | No hyaluronidase/hyaluronic acid  |
| Outcomes  | <ul> <li>Intraoperative pain</li> <li>Surgical complication rates</li> <li>Anaesthetic-related complications</li> <li>Patient satisfaction</li> <li>Adverse effects of treatment e.g. allergies</li> <li>Volume of anaesthetic</li> <li>Resource use and costs</li> </ul>   |
| Other criteria for inclusion/exclusion of studies           | <ul> <li>Exclude: studies on general ophthalmic conditions</li> <li>Exclude: letters, editorials, commentaries, narrative reviews, observational studies, case reports</li> <li>Exclude: combination surgical procedures i.e. cataract surgery in tandem with other surgical procedures e.g. trabeculectomy, canaloplasty, Descemet's stripping automated endothelial keratoplasty, keratorefractive; studies that do not specify the type of cataract surgery</li> </ul> |
| Review strategies   | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.   |
| Subgroup analyses (treatment effect modifiers)              | None specified  |
| Baseline characteristics to be extracted in evidence tables | None specified  |

|   | Details   |
|---|---|
| Review question   | In what circumstances should general anaesthesia be considered in phacoemulsification cataract surgery?   |
| Objectives  | To determine in what circumstances general anaesthesia should be considered in phacoemulsification cataract surgery.  |
| Type of review  | Intervention  |
| Language  | English   |
| Study design  | Any   |
| Status  | No date restrictions  |
| Population  | Adults (18 years and over) undergoing phacoemulsification cataract surgery for non-trauma related cataracts and intraocular lens (IOL) implantation   |
| Intervention  | General anaesthesia   |
| Comparator  | Not relevant  |
| Outcomes  | Indications for general anaesthesia in phacoemulsification cataract surgery   |
| Other criteria for inclusion/exclusion of studies           | <ul> <li>Exclude: studies on general ophthalmic conditions</li> <li>Exclude: letters, editorials, commentaries</li> <li>Exclude: combination surgical procedures i.e. cataract surgery in tandem with other surgical procedures e.g. trabeculectomy, canaloplasty, Descemet's stripping automated endothelial keratoplasty, keratorefractive; studies that do not specify the type of cataract surgery</li> </ul> |
| Review strategies   | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included studies will be assessed using relevant risk of bias tools. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE and/or CERQual as appropriate.   |
| Subgroup analyses (treatment effect modifiers)              | None specified  |
| Baseline characteristics to be extracted in evidence tables | None specified  |

|  | Details   |
|--|---|
| Review question  | Are different lens design (aspheric vs. spheric, plate vs. loop) effective in improving postoperative vision (refractive outcomes, optical aberrations) in cataract surgery?  |
| Objectives   | To determine the effectiveness of different lens design (aspheric vs. spheric, plate vs. loop) in improving postoperative vision (refractive outcomes, optical aberrations) in cataract surgery.  |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
|  | If none available then comparative cohort   |
| Status   | Articles published from 1990 onwards  |
|  | 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK   |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Different monofocal lenses:   |
|  | Aspheric vs. spheric  |
|  | Plate vs. loop vs. 3 piece  |
| Comparator   | As listed in the interventions  |
| Outcomes   | Visual acuity   |
|  | Visual function   |
|  | Patient reported dysphotopsia (count data)  |
|  | Night vision problems   |
|  | Contrast sensitivity  |
|  | Depth of focus  |
|  | Near vision   |
|  | Lens centration   |
|  | Quality of life   |
|  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details  |
|--|--|
| Review question  | Are different lens design (square-edged vs. round-edge, plate vs. loop) and material (hydrophilic acrylic, hydrophobic acrylic, collagen, hydroxyethyl methacrylate-based vs. silicone-based) effective in preventing posterior capsule opacification in cataract surgery?                               |
| Objectives   | To determine the effectiveness of different lens design (square-edged vs. round-edge, plate vs. loop) and material (hydrophilic acrylic, hydrophobic acrylic, collagen, hydroxyethyl methacrylate-based vs. silicone-based) effective in preventing posterior capsule opacification in cataract surgery. |
| Type of review   | Intervention   |
| Language   | English  |
| Study design   | Randomised controlled trials then comparative cohort   |
| Status   | Articles published from 1990 onwards<br>1990 chosen as a date to correspond with the widespread adoption of<br>phacoemulsification in the UK   |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation  |
| Intervention   | Different monofocal lenses:  • Square-edge vs. round-edge  • Plate vs. loop  • Hydrophilic acrylic, hydrophobic acrylic, collagen, hydroxyethyl methacrylate-based vs. silicone-based  |
| Comparator   | As listed in the interventions   |
| Outcomes   | <ul> <li>Rates of posterior capsule opacification</li> <li>Visual acuity</li> <li>Contrast sensitivity</li> <li>Quality of life</li> <li>Resource use and cost</li> </ul>  |
| Other criteria for inclusion/exclusion of studies                      | None specified   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.  |
| Subgroup analyses (treatment effect modifiers)                         | None specified   |
| Baseline characteristics to be extracted in evidence tables            | None specified   |

|  | Details   |
|--|---|
| Review question  | Are tinted lenses effective in preventing the incidence and progression of age-related macular degeneration compared with colourless lenses in cataract surgery?  |
| Objectives   | To determine the effectiveness of tinted lenses in preventing the progression of agerelated macular degeneration compared with colourless lenses in cataract surgery.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials then comparative observational   |
| Status   | Articles published from 1995 onwards  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Different monofocal/multifocal lenses:  |
|  | Tinted vs. colourless   |
|  | Different colours   |
| Comparator   | As listed in the interventions  |
| Outcomes   | <ul> <li>Incidence of age-related macular degeneration</li> <li>Rates of progression of age-related macular degeneration</li> <li>Visual acuity</li> <li>Colour vision</li> <li>Sleep problems</li> <li>Depression</li> <li>Quality of life</li> <li>Resource use and cost</li> </ul>                   |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | Diagnosed AMD (cataract or non-cataract eye)  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | Wat is the optimal strategy to facilitate simultaneous distance and near vision following   |
|  | cataract surgery?   |
| Objectives   | To compare the effectiveness of the following strategies to facilitate simultaneous distance and near vision in cataract surgery:   |
|  | multifocal intraocular lenses   |
|  | standard monofocal intraocular lenses with different focal points in each eye   |
|  | <ul> <li>standard monofocal intraocular lenses with the same focal point in both eyes plus<br/>glasses/contact lenses (optical correction)</li> </ul>   |
| Type of review   | Intervention  |
| Language   | Any   |
| Study design   | Randomised controlled trials (RCTs); unilateral and bilateral implantation  |
| Status   | Articles published from 1990 onwards  |
| Cidido   | 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK   |
| Population   | Adults (18 years and older) undergoing phacoemulsification cataract surgery and intraocular lens (IOL) implantation in one or both eyes   |
| Interventions  | <ul> <li>Any type of non-accommodative multifocal intraocular lenses (including toric<br/>multifocal lenses)</li> </ul>   |
|  | Examples: AcrySof IQ ReSTOR SN6AD3, ReSTOR SN6AD1, ReSTOR SN60D3, ReZoom NXG1, Gradiol (concept-gradient refractive index optics), Mplus X, MS 714 PB Diff, Sulcoflex 653F, TECNIS ZM900, ZMA00   |
|  | <ul> <li>Implantation of 1 or 2 monofocal intraocular lenses with the aim of optimising near<br/>vision in 1 eye and distance vision in the other</li> </ul>  |
|  | <ul> <li>Standard monofocal intraocular lenses with the same focal point in both eyes plus<br/>glasses/contact lenses (optical correction)</li> <li>Examples: Akreos AO, ZA9003</li> </ul>  |
| Comparators  | <ul> <li>All 3 listed interventions vs. each other</li> <li>Different types of multifocal lenses vs. each other</li> </ul>  |
| Outcomes   | Unaided near, intermediate and distance visual acuity   |
|  | Contrast sensitivity  |
|  | Complications: glare and other optical aberrations  |
|  | Visual function/Quality of life   |
|  | Best corrected visual acuity (BCVA): near, intermediate and distance  |
|  | Patient satisfaction  |
|  | Resource use and costs  |
| Other criteria for inclusion/exclusion of studies                      | Exclude: studies examining accommodating multifocal lenses only   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be pooled in pairwise and/or network meta-analyses. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | Lens technology (bifocal, trifocal, multifocal, refractive, diffractive)  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |
|  |   |

|  | Details   |
|--|---|
| Review question  | What is the optimal strategy to address pre-existing regular astigmatism in people undergoing cataract surgery?   |
| Objectives   | To determine the optimal strategy to address pre-existing astigmatism in people undergoing cataract surgery.  |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | <ul><li>Randomised controlled trials</li><li>If none available then comparative cohort</li></ul>  |
| Status   | Articles published from 1990 onwards 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation with pre-existing astigmatism   |
| Intervention   | <ul> <li>Corneal (limbal) relaxing incisions</li> <li>On-axis surgery (incision is made on steepest axis to flatten it)</li> <li>Astigmatic keratotomy</li> <li>Opposite clear corneal incisions (OCCI)</li> <li>Toric intraocular lens</li> </ul>  |
| Comparator   | <ul><li>Standard monofocal/multifocal lenses with no correction</li><li>Each other</li></ul>  |
| Outcomes   | <ul> <li>Visual acuity</li> <li>Level of astigmatism</li> <li>Patient satisfaction</li> <li>Quality of life</li> <li>Resource use and cost (including time taken)</li> </ul>  |
| Other criteria for inclusion/exclusion of studies                      | OECD only   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | <ul><li>Surgeon experience</li><li>Astigmatic corrections pre-operatively</li></ul>   |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of interventions (for example, prophylactic laser surgery) to prevent retinal detachment in people with myopia undergoing cataract surgery?   |
| Objectives   | To determine the effectiveness of interventions (for example, prophylactic laser surgery) to prevent retinal detachment in people with myopia undergoing cataract surgery.  |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) with myopia undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | <ul> <li>Prophylactic interventions prior to cataract surgery (not at the time of surgery)</li> <li>Retinal LASER surgery</li> <li>Cryotherapy</li> </ul>   |
| Comparator   | No specific intervention  |
| Outcomes   | <ul> <li>Rates of retinal detachment</li> <li>Time to event data</li> <li>Health-related quality of life</li> <li>Resource use and cost</li> </ul>  |
| Other criteria for inclusion/exclusion of studies                      | Exclude: non-OECD countries   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | <ul><li>Myopia ranges: 3 dioptres, &gt;3 dioptres</li><li>Age</li></ul>   |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of bilateral simultaneous (rapid sequential) cataract surgery compared with unilateral eye surgery?   |
| Objectives   | To determine the effectiveness of bilateral simultaneous (rapid sequential) cataract surgery compared with unilateral eye surgery.  |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Bilateral simultaneous (rapid sequential) cataract surgery  |
| Comparator   | Unilateral eye cataract surgery   |
| Outcomes   | <ul> <li>Visual acuity</li> <li>Visual function</li> <li>Complication rates (including refractive surprise)</li> <li>Health-related quality of life</li> <li>Patient satisfaction</li> <li>Resource use and cost</li> </ul>   |
| Other criteria for inclusion/exclusion of studies                      | Exclude: non-OECD countries   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | Medical comorbidities   |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the appropriate timing of second eye surgery, taking into account issues such as refractive power after first eye surgery?  |
| Objectives   | To determine the appropriate timing of second eye surgery, taking into account issues such as refractive power after first eye surgery.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation in the second eye   |
| Intervention   | Different timing of second eye surgery  |
| Comparator   | Different timings vs. each other No second eye surgery  |
| Outcomes   | <ul> <li>Visual acuity</li> <li>Visual function</li> <li>Complication rates (including refractive surprise)</li> <li>Falls</li> <li>Health-related quality of life</li> <li>Patient satisfaction</li> <li>Resource use and cost</li> </ul>  |
| Other criteria for inclusion/exclusion of studies                      | Exclude: non-OECD countries   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of capsular tension rings applied during phacoemulsification cataract surgery?  |
| Objectives   | To determine the effectiveness of capsular tension rings applied during phacoemulsification cataract surgery.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
| Status   | Articles published from 1990 onwards  |
|  | 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards   |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Capsular tension rings  |
| Comparator   | No capsular tension rings   |
| Outcomes   | <ul> <li>Post-operative complications (decentration)</li> <li>Visual acuity</li> <li>Post-operative refraction</li> <li>Resource use and cost</li> </ul>  |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | <ul><li>Pseudoexfoliation</li><li>Multifocal lenses</li><li>Toric lenses</li></ul>  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of interventions to increase pupil size to improve visual outcomes and reduce complications during phacoemulsification cataract surgery?  |
| Objectives   | To determine the effectiveness of interventions to increase pupil size to improve visual outcomes and reduce complications during phacoemulsification cataract surgery.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials then observational   |
| Status   | Articles published from 1990 onwards  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Interventions to increase pupil size  |
|  | <ul> <li>Intracameral mydriatics (with or without anaesthesia) – please provide list of<br/>mydriatics (tropicamide, phenylephrine)</li> </ul>  |
|  | <ul> <li>Viscomydriasis with a high-viscosity ophthalmic viscosurgical device (OVD) e.g.<br/>sodium hyaluronate</li> </ul>  |
|  | <ul> <li>Manual separation: synechiolysis and/or pupillary membranectomy with spatula and<br/>forceps</li> </ul>  |
|  | Mechanical pupillary stretching using iris hooks  |
|  | Sphincter cutting   |
|  | <ul> <li>Use of mechanical pupil dilation/expansion devices e.g. Graether pupil expander<br/>(Eagle Vision); Siepser Iris Protector ring; Perfect Pupil device (Milvella); Morcher<br/>Pupil Dilator (Morcher GmbH); Oasis Iris Expander (Oasis Medical, Inc.); Malyugin<br/>Ring (MicroSurgical Technology)</li> </ul> |
| Comparator   | <ul><li>No additional procedure</li><li>Each other</li></ul>  |
| Outcomes   | Complications (capsular rupture, haemorrhage)   |
|  | Post-operative complications (inflammation, distorted pupils)   |
|  | Visual acuity   |
|  | Visual function   |
| 011  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.                 |
| Subgroup analyses (treatment effect modifiers)                         | People with floppy iris syndrome  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of postoperative eye shields to prevent complications after cataract extraction?  |
| Objectives   | To determine the effectiveness of postoperative eye shields to prevent complications after cataract extraction.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | <ul><li>Postoperative eye shields</li><li>Length of time with eye shield</li></ul>  |
| Comparator   | <ul><li>No postoperative eye shields</li><li>Different lengths of time</li></ul>  |
| Outcomes   | <ul><li>Accidental trauma</li><li>Patient satisfaction</li><li>Resource use and cost</li></ul>  |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Deview question  | What is the effectiveness of prophylactic antiseptics (for example, topical iodine) and   |
| Review question  | antibiotics to prevent endophthalmitis after cataract surgery?  |
| Objectives   | To evaluate the effectiveness of the following interventions to prevent endophthalmitis after cataract surgery:   |
|  | Prophylactic antiseptics (for example, topical iodine)  |
|  | Prophylactic antibiotics  |
| Type of review   | Intervention  |
| Language   | Any   |
| Study design   | Randomised controlled trials (RCTs)   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing any cataract surgery  |
| Interventions  | • 1) Antiseptics (povidone iodine, chlorhexidine, tisept, presept) vs. no antiseptics   |
|  | <ul> <li>2a) Preoperative antibiotics (in theatre, several days before surgery) vs. no<br/>preoperative antibiotics</li> </ul>  |
|  | <ul> <li>2b) Timing of intraoperative antibiotics (i.e. administered up to the end of the<br/>operation e.g. with infusion in the middle of operation, at end of procedure)</li> </ul>  |
|  | <ul> <li>2c) Route of administration of intraoperative antibiotics (topical, parenteral,<br/>intravitreous, intracameral, subconjunctival, infusion during surgery) with or without<br/>postoperative antibiotics vs. no intraoperative antibiotics or different routes vs. each<br/>other</li> </ul>   |
|  | • 2d) Postoperative (early e.g. few days and longer term e.g. ≥1 week) topical and systemic antibiotics vs. no postoperative antibiotics  |
|  | • 2e) Different types of postoperative antibiotics vs. each other   |
|  | 2f) Duration and frequency of postoperative antibiotics   |
|  | <ul> <li>2g) Timing of antibiotics i.e. preoperative vs. intraoperative vs. postoperative vs.<br/>combinations of timing of administration</li> </ul>   |
| Comparators  | As above  |
| Outcomes   | <ul> <li>Endophthalmitis rates: verified/confirmed/culture positive (preferred), suspected, any</li> <li>Adverse effects of treatment</li> <li>Best corrected distance visual acuity</li> <li>Resource use and costs</li> </ul>   |
| Other criteria for inclusion/exclusion of studies                      | • Exclude: non-OECD (Organisation for Economic Co-operation and Development) countries as pathogens and care are likely to be different   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses  | People on warfarin  |
| (treatment effect modifiers)   | <ul> <li>People with an increased risk of infection (inflammatory blepharitis, tear duct<br/>obstruction)</li> </ul>  |
|  | People who are immunocompromised  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of prophylactic topical corticosteroids and/or NSAIDs to prevent inflammation and cystoid macular oedema after phacoemulsification cataract surgery?  |
| Objectives   | To evaluate the effectiveness of prophylactic topical corticosteroids and/or non-<br>steroidal anti-inflammatory drugs (NSAIDs) to prevent inflammation and cystoid<br>macular oedema following phacoemulsification cataract surgery  |
| Type of review   | Intervention  |
| Language   | Any   |
| Study design   | Randomised controlled trials (RCTs); single and both eyes   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | <ul> <li>Corticosteroid drops vs. no treatment</li> <li>NSAID drops vs. no treatment</li> <li>Combination of corticosteroid and NSAID drops vs. no treatment</li> <li>Combination of intraoperative/postoperative corticosteroid injection and postoperative drops vs. no treatment (steroid vs NSAID vs no treatment)</li> <li>Corticosteroid drops vs. NSAID drops</li> <li>Timing of postoperative treatment (e.g. 2 vs 4 vs 6 weeks)</li> <li>Different dosing (frequency and duration) of postoperative treatment</li> </ul> |
| Comparator   | As above  |
| Outcomes   | <ul> <li>Inflammation rates</li> <li>Cystoid macular oedema (clinically symptomatic, optical coherence tomography-verified)</li> <li>Best corrected distance visual acuity</li> <li>Adverse effects of treatment e.g. raised intraocular pressure (steroid-induced glaucoma), allergies (such as sensitivity to preservatives)</li> <li>Resource use and costs</li> </ul>   |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Include: incident pseudophakic cystoid macular oedema</li> <li>Exclude: non-OECD (Organisation for Economic Co-operation and Development) countries as pathogens and care are likely to be different</li> </ul>  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE.   |
| Subgroup analyses (treatment effect modifiers)                         | <ul> <li>Higher risk populations e.g. diabetic macular oedema</li> <li>Complicated vs. uncomplicated phacoemulsification cataract surgery</li> </ul>  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of interventions to reduce the impact of peroperative posterior capsule rupture?  |
| Objectives   | To determine the effectiveness of interventions to reduce the impact of perioperative posterior capsule rupture.  |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
| Status   | Articles published from 1990 onwards 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation who experience a peroperative posterior capsule rupture   |
| Intervention   | <ul> <li>Anterior vitrectomy + Triamcinolone</li> <li>Timing and type of lens insertion</li> <li>Early versus late lens removal when lens fallen into back of eye</li> </ul>  |
| Comparator   | <ul><li>Anterior vitrectomy</li><li>Different timings and types</li><li>Other timing</li></ul>  |
| Outcomes   | <ul> <li>Visual acuity</li> <li>Visual function</li> <li>Complications (inflammation and pressure)</li> <li>Quality of life</li> <li>Resource use and cost</li> </ul>   |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What is the effectiveness of interventions used to manage cystoid macular oedema following cataract surgery?  |
| Objectives   | To determine the effectiveness of interventions used to manage cystoid macular oedema following cataract surgery.   |
| Type of review   | Intervention  |
| Language   | English   |
| Study design   | Randomised controlled trials  |
| Status   | Articles published from 1990 onwards 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | <ul> <li>NSAIDs</li> <li>SAIDs</li> <li>Diamox</li> <li>Periocular and intraocular steroids</li> <li>Intraocular Anti-VEGF</li> <li>Vitrectomy</li> </ul>   |
| Comparator   | <ul><li>No intervention</li><li>Each other</li></ul>  |
| Outcomes   | <ul> <li>Visual acuity</li> <li>Further surgery (for non-vitrectomy interventions)</li> <li>Macular thickness</li> <li>Time to resolution</li> <li>Adverse events</li> <li>Quality of life</li> <li>Resource use and cost</li> </ul>  |
| Other criteria for inclusion/exclusion of studies                      | None specified  |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included RCTs will be assessed using the Cochrane's risk of bias tool. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE. |
| Subgroup analyses (treatment effect modifiers)                         | <ul><li>Retinal/vascular disease</li><li>Diabetic macular oedema</li></ul>  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details   |
|--|---|
| Review question  | What are the early and late complications of cataract surgery?  |
| Objectives   | To determine the early and late complications of phacoemulsification cataract surgery.  |
| Type of review   | Epidemiological   |
| Language   | English   |
| Study design   | Randomised controlled trials  |
|  | Observational studies   |
| Status   | No date restrictions  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation   |
| Intervention   | Not relevant  |
| Comparator   | Not relevant  |
| Outcomes   | All complications   |
|  | Loss of visual acuity   |
|  | Loss of visual function   |
|  | Health-related quality of life  |
|  | Resource use and cost   |
| Other criteria for inclusion/exclusion of studies                      | Exclude: non-OECD countries   |
| Review strategies (data extraction, quality assessment, data analysis) | Citation appraisal against specified criteria to identify relevant studies. Data extracted and included studies will be assessed using relevant risk of bias tools. Where appropriate, data will be meta-analysed. The overall quality of the evidence for each outcome will be assessed using GRADE and/or CERQual as appropriate. |
| Subgroup analyses (treatment effect modifiers)                         | None specified  |
| Baseline characteristics to be extracted in evidence tables            | None specified  |

|  | Details  |
|--|--|
| Review question  | What should the postoperative assessment include?  |
| Objectives   | To determine the content of the postoperative assessment following phacoemulsification cataract surgery.   |
| Type of review   | Qualitative  |
| Language   | English  |
| Study design   | Interviews, surveys, mixed-methods designs   |
| Status   | Articles published from 1990 onwards   |
|  | 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards  |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation  |
| Intervention   | Not relevant   |
| Comparator   | Not relevant   |
| Outcomes   | Content in postoperative assessment  |
|  | Resource use and cost  |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Include studies that used qualitative methods for data collection (including focus<br/>group and individual interviews, observation, and document analysis) and qualitative<br/>methods for data analysis (including thematic analysis or any other appropriate<br/>qualitative analysis method that enabled analysis of text and observations and<br/>narrative presentation of findings)</li> </ul> |
|  | <ul> <li>Exclude non-qualitative research and studies for example, narrative reviews,<br/>commentaries, editorials/letters, opinion pieces, case studies/reports.</li> </ul>   |
|  | <ul> <li>Exclude studies that collected data using qualitative methods but<br/>analysed/presented the data using only quantitative methods.</li> </ul>   |
| Review strategies (data extraction, quality assessment, data analysis) | The quality of included papers will be assessed using appropriate study design specific checklists.  |
|  | Data will be analysed qualitatively (thematic synthesis methods) and overall confidence in the evidence for each outcome will be assessed using the CERQual tool.  |
| Subgroup analyses (treatment effect modifiers)                         | None specified   |
| Baseline characteristics to be extracted in evidence tables            | None specified   |

|  | Details  |
|--|--|
| Review question  | Who and in what setting should carry out the postoperative assessment?   |
| Objectives   | To determine who and in what setting should carry out the postoperative assessment.  |
| Type of review   | Qualitative  |
| Language   | English  |
| Study design   | Interviews, surveys, mixed-methods designs   |
| Status   | Articles published from 1990 onwards 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards   |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation  |
| Intervention   | Not relevant   |
| Comparator   | Not relevant   |
| Outcomes   | <ul> <li>Investigations performed</li> <li>Further interventions - re referral rates</li> <li>Additional medications prescribed</li> <li>Delays in diagnosis and treatment</li> <li>Resource use and costs</li> </ul>  |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Include studies that used qualitative methods for data collection (including focus group and individual interviews, observation, and document analysis) and qualitative methods for data analysis (including thematic analysis or any other appropriate qualitative analysis method that enabled analysis of text and observations and narrative presentation of findings)</li> <li>Exclude non-qualitative research and studies for example, narrative reviews, commentaries, editorials/letters, opinion pieces, case studies/reports.</li> <li>Exclude studies that collected data using qualitative methods but analysed/presented the data using only quantitative methods.</li> </ul> |
| Review strategies (data extraction, quality assessment, data analysis) | The quality of included papers will be assessed using appropriate study design specific checklists.  Data will be analysed qualitatively (thematic synthesis methods) and overall confidence in the evidence for each outcome will be assessed using the CERQual tool.   |
| Subgroup analyses (treatment effect modifiers)                         | None specified   |
| Baseline characteristics to be extracted in evidence tables            | None specified   |

|   | Details  |
|---|--|
| Review question   | What issues should be considered when organising postoperative care?   |
| Objectives  | To determine what issues should be considered when organising postoperative care.  |
| Type of review  | Qualitative  |
| Language  | English  |
| Study design  | Interviews, surveys, mixed-methods designs   |
| Status  | Articles published from 1990 onwards   |
| Ciaido  | 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards  |
| Population  | <ul> <li>Adults (18 years and over) undergoing phacoemulsification cataract surgery with<br/>intraocular lens implantation</li> </ul>  |
|   | People at high risk of non-adherence to standard postoperative care  |
| Intervention  | Not relevant   |
| Comparator  | Not relevant   |
| Outcomes  | Planned preoperatively at pre assessment   |
|   | Resource use and cost  |
| Other criteria for inclusion/exclusion of studies           | <ul> <li>Include studies that used qualitative methods for data collection (including focus<br/>group and individual interviews, observation, and document analysis) and qualitative<br/>methods for data analysis (including thematic analysis or any other appropriate<br/>qualitative analysis method that enabled analysis of text and observations and<br/>narrative presentation of findings)</li> </ul> |
|   | • Exclude non-qualitative research and studies for example, narrative reviews, commentaries, editorials/letters, opinion pieces, case studies/reports.   |
|   | <ul> <li>Exclude studies that collected data using qualitative methods but<br/>analysed/presented the data using only quantitative methods.</li> </ul>   |
| Review strategies (data extraction, quality                 | The quality of included papers will be assessed using appropriate study design specific checklists.  |
| assessment, data analysis)                                  | Data will be analysed qualitatively (thematic synthesis methods) and overall confidence in the evidence for each outcome will be assessed using the CERQual tool.  |
| Subgroup analyses (treatment effect modifiers)              | None specified   |
| Baseline characteristics to be extracted in evidence tables | None specified   |

|  | Details  |
|--|--|
| Review question  | What is the appropriate time to assess outcomes in the postoperative period?   |
| Objectives   | To determine the appropriate time to assess outcomes in the postoperative period.  |
| Type of review   | Qualitative  |
| Language   | English  |
| Study design   | Interviews, surveys, mixed-methods designs   |
| Status   | Articles published from 1990 onwards 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards   |
| Population   | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation  |
| Intervention   | Not relevant   |
| Comparator   | Not relevant   |
| Outcomes   | Resource use and cost     Stable visual outcome  |
| Other criteria for inclusion/exclusion of studies                      | <ul> <li>Include studies that used qualitative methods for data collection (including focus<br/>group and individual interviews, observation, and document analysis) and qualitative<br/>methods for data analysis (including thematic analysis or any other appropriate<br/>qualitative analysis method that enabled analysis of text and observations and<br/>narrative presentation of findings)</li> </ul> |
|  | • Exclude non-qualitative research and studies for example, narrative reviews, commentaries, editorials/letters, opinion pieces, case studies/reports.   |
|  | <ul> <li>Exclude studies that collected data using qualitative methods but<br/>analysed/presented the data using only quantitative methods.</li> </ul>   |
| Review strategies (data extraction, quality assessment, data analysis) | The quality of included papers will be assessed using appropriate study design specific checklists.  |
|  | Data will be analysed qualitatively (thematic synthesis methods) and overall confidence in the evidence for each outcome will be assessed using the CERQual tool.  |
| Subgroup analyses (treatment effect modifiers)                         | None specified   |
| Baseline characteristics to be extracted in evidence tables            | None specified   |

|   | Details  |
|---|--|
| Review question   | If the postoperative assessment and care are undertaken outside of the hospital, how should outcomes between surgical units and these providers be effectively communicated?   |
| Objectives  | To determine how outcomes between surgical units and postoperative care providers should be effectively communicated, if the postoperative assessment and care are undertaken outside of the hospital.   |
| Type of review  | Qualitative  |
| Language  | English  |
| Study design  | Interviews, surveys, mixed-methods designs   |
| Status  | Articles published from 1990 onwards 1990 chosen as a date to correspond with the widespread adoption of phacoemulsification in the UK onwards   |
| Population  | Adults (18 years and over) undergoing phacoemulsification cataract surgery with intraocular lens implantation  |
| Intervention  | Not relevant   |
| Comparator  | Not relevant   |
| Outcomes  | Not relevant   |
| Other criteria for inclusion/exclusion of studies           | <ul> <li>Include studies that used qualitative methods for data collection (including focus<br/>group and individual interviews, observation, and document analysis) and qualitative<br/>methods for data analysis (including thematic analysis or any other appropriate<br/>qualitative analysis method that enabled analysis of text and observations and<br/>narrative presentation of findings)</li> </ul> |
|   | <ul> <li>Exclude non-qualitative research and studies for example, narrative reviews, commentaries, editorials/letters, opinion pieces, case studies/reports.</li> <li>Exclude studies that collected data using qualitative methods but</li> </ul>  |
|   | analysed/presented the data using only quantitative methods.   |
| Review strategies (data extraction, quality                 | The quality of included papers will be assessed using appropriate study design specific checklists.  |
| assessment, data analysis)                                  | Data will be analysed qualitatively (thematic synthesis methods) and overall confidence in the evidence for each outcome will be assessed using the CERQual tool.  |
| Subgroup analyses (treatment effect modifiers)              | None specified   |
| Baseline characteristics to be extracted in evidence tables | None specified   |