



MIRROR TRIAL
PARTICIPANT CONSENT FORM

Full title of Project: Efficacy of the Telescopic Mirror Implant for Age-related Macular Degeneration: The MIRROR Trial (acronym MIRROR). A Multicentre Randomised Controlled Clinical Trial

**Please
Initial
Box**

1. I confirm that I have read, or have read to me, and understand the Participant Information Sheet, v4.0 dated *day month* 2016 for the above study. I have had the opportunity to ask questions and these have been fully answered.

2. I confirm that I have had sufficient time to consider whether or not to participate in the study.

3. I understand that my participation is voluntary and that I am free to withdraw at any time, without giving any reason and without my legal rights or medical care being affected.

4. I understand the study is being conducted by the Belfast Health and Social Care Trust and that my personal information will be held securely on NHS/HSC Trust premises and handled in accordance with the provisions of the Data Protection Act 1998.

Copies: 1 original in Investigator Site File.1 original copy for patient. 1 copy in Patient's Medical Records

5. I understand that data collected as part of this study and relevant sections in my medical records may be looked at by authorised individuals from regulatory authorities or NHS/HSC Trusts and representatives from the study Sponsor where it is relevant to my taking part in this research. I give permission for these individuals to have access to this information.
6. I understand that images captured of my eyes will be anonymised and stored electronically on secure servers held in Queen's University Belfast.
7. I understand that the information I provide may be published as a report. Confidentiality and anonymity will be maintained and it will not be possible to identify me from any publications.
8. I agree to my GP being informed of my participation in the study.

I agree to take part in the above study.

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| | | |
| Name of Participant (Please print) | Date | Signature |

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| | | |
| Name of Person taking consent (Please print) | Date | Signature |

Copies: 1 original in Investigator Site File.1 original copy for patient. 1 copy in Patient's Medical Records