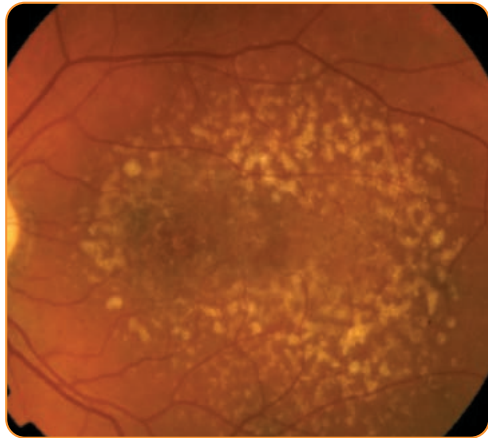


# Identification of patients suitable for EyeMax Mono™ implantation



## Early/intermediate AMD

- EyeMax Mono used as 'insurance' against progression to geographic atrophy (GA)
- No initial benefit vs standard monofocal IOL; benefit with EyeMax Mono is conferred with the onset of central GA
- Caution with rapidly progressing phenotypes (e.g. diffuse trickling)



## GA with foveal island

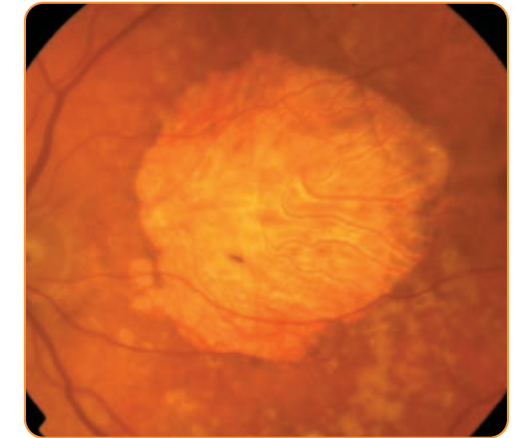
- Proceed with caution
- No initial benefit over standard monofocal IOL; benefit with EyeMax Mono is conferred when the foveal island is lost
- Counsel the patient to expect significant loss of function when the central foveal island is lost



## Centre-involving GA with functioning retina within 2 disc diameters of anatomical centre<sup>1,2</sup>

### • Ideal initial patient for EyeMax Mono surgery

- Ensure glasses are prescribed post-operatively (avoid varifocals)
- Allow time for neuroadaptation
- Likely no value in using eccentric fixation as criterion for implantation, the patient may adopt this post-operatively even if not present before



## Centre-involving GA with no functioning retina within 2 disc diameters of anatomical centre

- Likely no optical benefit of EyeMax Mono vs a standard monofocal IOL

**Dry AMD is a progressive condition and patients lose visual acuity and function over time**

**But EyeMax Mono maximises the capacity of surviving retina to support activities of daily living**

EyeMax Mono should be used in patients with:

Cataract requiring surgery<sup>1,2</sup>

Centre-involving dry AMD<sup>1,2</sup>

Sufficient remaining macular function within 10° of the foveal centre<sup>1,2</sup>

EyeMax Mono may be implanted binocularly or monocularly, with the crystalline lens retained or a standard monofocal lens implanted in the fellow eye.<sup>1-3</sup> If there is significant interocular difference in visual acuity then consider operating on the better-seeing eye first

- Counsel the patient that gains in visual acuity may not necessarily translate to functional improvement
- Approximately 5% of patients may experience no gain in visual acuity<sup>1</sup>

EyeMax Mono is designed to optimise visual outcomes in patients with dry AMD. Please refer to the Instructions For Use. Adverse events should be reported as soon as possible to [eventreporting@invua.com](mailto:eventreporting@invua.com).

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**Post-operative target refraction for EyeMax Mono should be between emmetropia and +3 D<sup>1-3</sup>**

**Avoid anisometropia of >2 D<sup>1-4</sup>**

- By refining the optic of EyeMax Mono for a 0 to +3 D outcome it gives the option of leaving the patient with +3 D correction with glasses, providing 10–20% ‘magnification’ if their vision is very poor

**Target refraction will depend on:<sup>1-3</sup>**

- Existing refraction
- Patient preference
- Severity of the maculopathy

Patient's original vision:	Target for:
Better than 0.1 decimalised	Emmetropia (not myopic side of emmetropia)
Worse than 0.1 decimalised	+3 D
Already hypermetropic	Consider leaving hypermetropic

**Determine the IOL power using the correct biometry formula for the patient's axial length<sup>1,2,4</sup>**

**Select the appropriate EyeMax Mono lens using the A-constant of 119.2<sup>1,2,4</sup>**

- The need for continued use of glasses should be explained upfront to manage patient expectations<sup>1-3</sup>
- Separate glasses for near and distance are probably better than varifocals for this patient group
- Low vision aids should be used/prescribed as normal

**No toric version of EyeMax Mono available. Astigmatism should be managed as for implantation of a standard monofocal lens**

**References:**

1. Qureshi MA, et al. Eur J Ophthalmol 2018;28:198–203;
2. Robbie SJ, et al. J Refract Surg 2018;34:718–25;
3. EyeMax Mono, data on file 2019;
4. National Institute for Health and Clinical Excellence. 2018. Available at: <https://www.nice.org.uk/guidance/ng77/>. Accessed January 2020.