

**EC Certificate**  
**Full Quality Assurance System according to**  
**Medical Devices Directive 93/42/EEC Annex-II Section 3**

**Certificate Number: 1984-MDD-21-818**

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the national legislation to which the undersigned is subjected, transposing annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive.

**Organization:**

**Rumex International Ltd.**

311 Shoreham Street Sheffield, South Yorkshire, S2 4FA, UK

**Product:** Hydrophobic Intraocular lenses

**Types:** AquaFree Yellow Aspheric Hydrophobic, AquaFree Yellow Preloaded

**Product:** Hydrophilic Intraocular lenses

**Types:** Hydro-Sense Aspheric, Hydro-Sense Aspheric Yellow, Hydro-4 Aspheric

The certificate is valid till expiration date, subject to successful completion of periodical surveillance audits. Please contact Kiwa for details.

**Report Number:** M.6028.01

**Expiry Date:** 27 May 2024

Kiwa Belgelendirme Hizmetleri A.Ş. is Notified Body under Council Directive 93/42/EEC concerning medical devices with identification number: 1984



**Muhtesem Gökhan Yücel**  
Head of Notified Body

20 May 2021, Istanbul, Turkey