



To: **Muscat**
THE DIRECTOR GENERAL OF HEALTH SERVICES IN ALL GOVERNORATES
Commanding Officer, Armed Forces Hospital (Al Khoudh & Salalah)
Director General of Engineering Affairs, MOH
Director General of Royal Hospital
Director General of Khoula Hospital
Director General of Medical Supplies (MOH)
Director General of Pvt. Health Est. Affairs (to kindly arrange distribution to all Pvt. Hospitals)
Hospital Director (Al Nahda Hospital)
Hospital Director (Al Massara Hospital)
The Head of Medical Services in SQU Hospital
The Head of Medical Services in Royal Oman Police
The Head of Medical Services in Ministry of Defence
The Head of Medical Services in The Diwan
The Head of Medical Services in The Sultan's Special Force
The Head of Medical Services in Internal Security Services
The Head of Medical Services in Petroleum Development of Oman
The Head of Medical Services in LNG Oman
ALL PRIVATE PHARMACIES & DRUG STORES

After Compliments,

Please find attached our Circular No. 267 dated 21/12/23 Regarding NCMDR recall of Hydrophobic acrylic pre-loaded intraocular lens from (mfr: Cristalens Industrie).

Copy to:

- Director, Office of H.E. The Undersecretary for Health Affairs
- Director of Medical Device Control, DGPA&DC
- Director of Pharmacovigilance & Drug Information Dept, DGPA&DC
- Director of Drug Control Department, DGPA&DC
- Director of Pharmaceutical Licensing Department, DGPA&DC
- Director of Central Quality Control Lab., DGPA&DC
- Supdt. of Central Drug Information





Circular No. 267/2023

08-06-1445 H

29-12-2023

نتقدم بثقة
Moving Forward
with Confidence



Recall of Hydrophobic acrylic pre-loaded intraocular lens from Cristalens Industrie

Source	NCMDR- National Center for Medical Devices Reporting- SFDA https://ncmdr.sfda.gov.sa/Secure/CA/CaViewRecall.aspx?caid=4&rid=19817
Product	Hydrophobic acrylic pre-loaded intraocular lens.
Description	Sterile Preloaded Hydrophilic Intraocular Lenses.
Manufacturer	Cristalens Industrie.
The affected products	Intraocular lens -Model: ARTIS PLE Dioptry 12.5D to 14.5D; ARTIS T PL E diopter 14D. See Appendix 1 in the attachment for a list of serial numbers.
Reason	The power of the implanted lens does not correspond to the power desired by the practitioner.
Action	1. Please do not use the affected products. 2. Return the affected products. 3. Contact the local agent for remedial action.
comments	Healthcare professionals are encouraged to report any adverse events Suspected to be associated with the above device or any other medical device to Department of Medical Device Control through the E-mail: Med-device@moh.gov.om

Dr. Mohammed Hamdan Al Rubaie
Director General

