



To:

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. 129 dated 28/7/22 Regarding NCMDR recall of Microvisc 1%, Microvisc Plus 1,4%, Microvisc Phaco 2,5% from (mfr: Bohus BioTech AB).

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. 129 / 2022

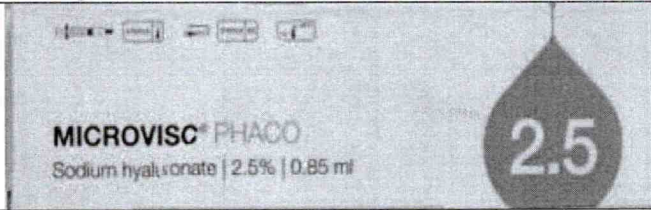
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28 -07-2022

بنقدم بثقة
Moving Forward
with Confidence



Recall of Microvisc 1%, Microvisc Plus 1,4%, Microvisc Phaco 2,5% from Bohus BioTech AB.

Source	NCMDR- National Center for Medical Devices Reporting- SFDA https://ncmdr.sfda.gov.sa/Secure/CA/CAViewRecall.aspx?caid=6&rid=16187
Product	Microvisc 1%, Microvisc Plus 1,4%, Microvisc Phaco 2,5%.
Description	Ophthalmological technology - liquid media for ophthalmological purposes.
Manufacturer	Bohus BioTech AB.
The affected products	Include all syringe sizes, 0,55 ml, 0,85 ml and 1,0 ml. This also include all products being under own label/private label.
Reason	A deviation in sterilization process.
Action	1. Recall all products within expiry date. 2. Contact the local agent for remedial action.
Product image	
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

