



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 43 dated 01/03/2023 Regarding NCMDR recall of Rescue Drill Guide Conical Connection RP from (mfr: Nobel Biocare Services 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. 43/2023

08-08-1444 H

01-03-2023

نتقدم بثقة
Moving Forward
with Confidence

رؤية عمان
2040
Oman Vision

Recall of Rescue Drill Guide Conical Connection RP from Nobel Biocare Services AB.

Source	NCMDR- National Center for Medical Devices Reporting- SFDA https://ncmdr.sfda.gov.sa/Secure/CA/CAViewRecall.aspx?caid=4&rid=18460
Product	Rescue Drill Guide Conical Connection RP.
Description	Dental Retrieval Instrument.
Manufacturer	Nobel Biocare Services AB.
The affected products	Catalog number: 37487 Lot number: 164473
Reason	Batch 164473 has been manufactured in an incorrect raw material, Titanium Grade 5 (Ti6Al4V ELI) instead of Stainless Steel (W.1.4305).
Action	1. Inspect your stock and quarantine affected devices. 2. Return all affected stock. 3. 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

Ph. Ahmed Al harbi

Acting Director Genera

