



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

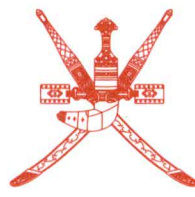
After Compliments,

Please find attached our Circular No 180 dated 02/10/2022 Regarding NCMDR Recall of BD Veritor™ Plus Analyzers from (mfr: BD).

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. 180/2022

06-03-1444 H

02-10-2022

بدمع بثقة
Moving Forward
with Confidence



Recall of BD Veritor™ Plus Analyzers from BD.

Source	GHC- Gulf Health Council.		
Product	BD Veritor™ Plus Analyzers.		
Manufacturer	BD.		
Local agent	Aston Medical Supplies LLC.		
The affected products	Product Code (REF)	Serial number	Expiration Date
	256066	All unexpired analyzers	Up to & including 31/08/2024
Reason	Addendum to the warning dated 24/1/2022 No. MOHAP/O/22/000563, BD has updated from an “advisory” to a “product removal” of unexpired analyzers within an expiry date		
Action	1. Quarantine all affected lots and return to the supplier for replacement. If you choose to continue use of the BD Veritor™ Plus Analyzer whilst waiting for your new device, BD recommends following the guidance from the company to reduce as much as possible, the risks associated with the described defect. 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

