



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

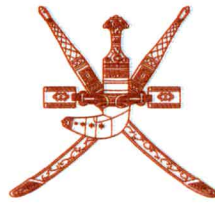
After Compliments,

Please find attached our Circular No 12 dated 22/1/2023 Regarding NCMDR recall of BIOHIT ColonView quick test from Ethicon Inc from (mfr: BIOHIT OYJ).

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. 12 / 2023

29 -06-1444 H

22 -01-2023

نتقدم بثقة
Moving Forward
with Confidence



Recall of BIOHIT ColonView quick test from BIOHIT OYJ

Source	NCMDR- National Center for Medical Devices Reporting- SFDA https://ncmdr.sfda.gov.sa/Secure/CA/CaViewRecall.aspx?caid=4&rid=18411
Product	BIOHIT ColonView quick test.
Description	IVD.
Manufacturer	BIOHIT OYJ.
Local Agent	HEALTH 4 ALL (LLC).
The affected products	REF 602250.02 Lot number: 22062004 exp. 2023-12
Reason	The test line intensities of the specified kit lot are lower than normal so that the results do not fulfill the acceptance criteria. Therefore, there is a risk for misdiagnosis, since false negative results are possible in case of weak positive samples.
Action	1. Discontinue distribution and dispose of all kits of the above-mentioned lot. 2. Review the previously generated results. 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

Dr. Mohammed Hamdan Al Rubaie

Director General

