Sultanate of Oman
Ministry of Health
Directorate General of Pharmaceutical
Affairs and Drug Control
Muscat

سلطنة عُمان
وزارة الصحة
المديرية العامة للصيدلة
والرقابة الدوائية
مسقط

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. 18/14 **dated** 27/10/22 **Regarding GHC Recall of DLP® Left Heart Vent Catheters from (mrf: Medtronic).**

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**

PADC
المديرية العامة للصيدلة والرقابة الدوائية
Directorate General of Pharmaceutical Affairs & Drug Control

ص.ب: ٣٩٣ مسقط - الرمز البريدي: ١٠٠ - هاتف: ٢٢٣٥٧١١١ - فاكس: ٢٢٣٥٨٤٨٩
P.O. Box: 393 Muscat - Postal Code: 100 - Tel: 22357111 - Fax: 22358489
dgpa_dc Email: dg-padc@moh.gov.om

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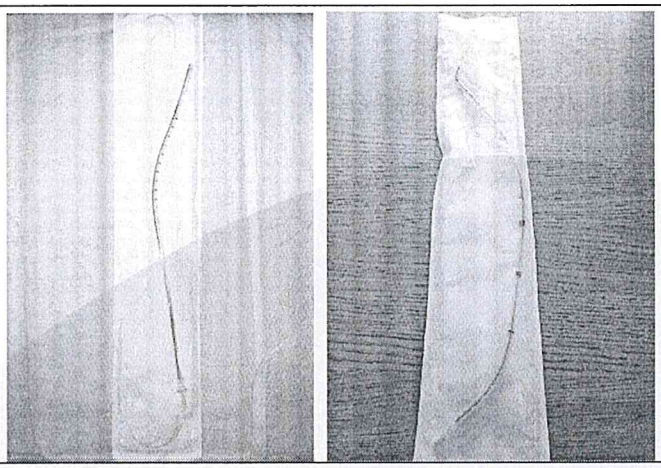
Circular No. 184 / 2021

20 -03-1443 H

27 -10-2021

Recall of DLP® Left Heart Vent Catheters from Medtronic.

Source	GHC- Gulf Health Council.
Product	DLP® Left Heart Vent Catheters.
Manufacturer	Medtronic.
Local Agent	Alzahrawi Medical Supplies LLC.
The affected products	Product Name: DLP® Left Heart Vent Catheters, Model Number: 16 Fr 12116,18 Fr 12118. Product Name: DLP® Left Heart Vent Catheters, Model Number: 18 Fr 12118.
Reason	Medtronic is voluntarily recalling specific models of unused above-mentioned products, due to the potential for a wire protrusion through the left heart vent catheter tip. If unnoticed prior to the procedure, this wire protrusion could lead to tissue damage (abrasion/perforation) and thereby to a longer duration of the procedure and/or surgical repair. In those procedures where a Model 12116 or Model 12118 was used, monitor patients in the acute post-op care setting for bleeding.
Action	1. Do not use any affected products, and return all unused affected product. 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 contact E-mail: Med-device@moh.gov.om

Dr. Mohammed Hamdan Al Rubaie

DIRECTOR GENERAL

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