



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 232 dated 29/10/2023 Regarding NCMDR Field Safety Notice of CARDIOHELP-i from (mfr: MAQUET Cardiopulmonary GmbH).

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





نتقدم بثقة  
Moving Forward  
With Confidence

رؤية عمان  
2040  
Oman Vision

Circular No. 232 / 2023

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29 -10-2023

Field Safety Notice of CARDIOHELP-i from MAQUET Cardiopulmonary GmbH.

Source	NCMDR - National Center Medical Device Reporting- SFDA. <a href="https://ncmdr.sfda.gov.sa/Secure/CA/CAViewRecall.aspx?caid=4&amp;rid=19742">https://ncmdr.sfda.gov.sa/Secure/CA/CAViewRecall.aspx?caid=4&amp;rid=19742</a>
Product	CARDIOHELP-i.
Description	Heart-lung bypass system.
Manufacturer	MAQUET Cardiopulmonary GmbH.
Local agent	Mustafa Sultan Science & Industry Co.LLC.
The affected products	Article number: 701048012 Unique Device Identifier: 04037691658384 Serial numbers: 90414549, 90414727 & 90414729
Reason	It was found that the fixation (weld) of the nut bolt to the slide rail plate was insufficient. The deficient weld allows the nut bolt to break off from the slide rail plate with slight manual force.
Action	1. Affected CARDIOHELP-i are not requested to be returned and can be used for intra-hospital support as usual. Inter-hospital transport is not permitted until the replacement is performed. 2. Please do not to fixate any accessory at the side rail until the replacement is performed. 3. You will be contacted by Getinge representative to arrange the replacement of the slide rail holders. 4. Contact the local agent for remedial action.
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: <a href="mailto:Med-device@moh.gov.om">Med-device@moh.gov.om</a>

Dr. Mohammed Hamdan Al Rubaie  
Director General

