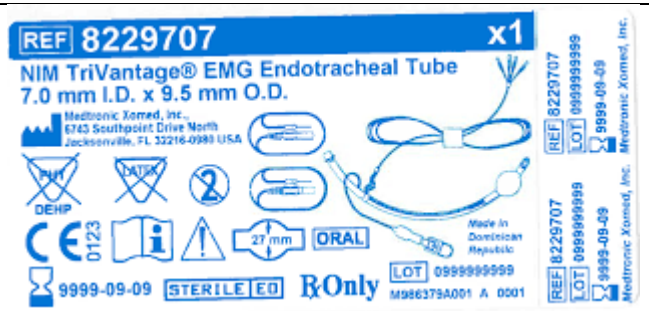
Medical Devices Safety Notice

The National Health Regulatory Authority would like to alert all governmental and private healthcare facilities, local agents and distributors that the below medical device:

Device Details	
Device Name	NIM Trivantage EMG Endotracheal Tube
Lot No.	8229707
Manufacturer	Medtronic SA
Country of Origin	USA
Reference	https://recalls-rappels.canada.ca/en/alert-recall/nim-trivantage-emg-endotracheal-tube
Device picture	
Reason of Recall	NHRA initiates this FSN due to problems with the Trivantage EMG tubes such as having difficulty passing the electrode check screen on the NIM mainframe immediately after intubating the patient, losing the connection between the EMG tube and the mainframe during a procedure resulting in them getting an "electrode off" or "channel not reading message" on the monitor and being unable to proceed with nerve monitoring, or getting excess noise from the tube that would sometimes also result in "lead-off" issues.
Action should be taken	Please stop using the above mentioned medical device and contact the authorized representative **Behzad** at **info@behzadmedical.com.bh** to take the necessary action for recall.

Your cooperation is highly appreciated in improving health services in the Kingdom of Bahrain.

For more information please contact Medical_Devices@nhra.bh

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