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	PERFUSOR LINE, PE, LL, 200 CM
Device Model	8723060
Batch No.	22C11E8SC6 22D16E8SC6
Manufacturer	B. Braun Melsungen AG
Country of Origin	Germany
Reference	Link
Reason of Recall	NHRA initiates this FSN due to risk for holes and leakages in the combinations of Original Perfusor® Line. The deviation might harbour the risk for the patient of microbial contamination, under supply, open patient access, or air infusion.
Action should be taken	Please stop using the above mentioned medical device and contact the authorized representative Wael Pharmacy Co. W.L.L at vincent@waelpharmacy.com 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

Recall 2023 0008

04/May/2023