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	Welch Allyn Spot Vital Signs 4400 Device
Device Model	44WT-4
Lot No.	manufactured on or before November 9, 2024
Manufacturer	BAXTER
Country of Origin	USA
Reference	Link
Reason of Recall	NHRA initiates this FSN due to the discovery that certain Welch Allyn reusable blood pressure cuffs labeled “not made with natural rubber latex” contain a latex rubber band around the instructions for use (IFU), which may expose individuals with latex allergies to potential allergic reactions, including anaphylaxis.
Action should be taken	Please stop using the above mentioned medical device and contact the authorized representative Behzad Medical Establishment at imran@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

Recall 2025 0039

19/Oct/2025