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	Contact Lenses Solutions	
Device Model	**Model Name**	**Lot No.**
	RENU Multiplus Multi-purpose 360ml	ME2970, MF5135, MF1656, ME3081
	RENU Multiplus Multi-purpose 120ml	ME3124, ME4078
	Starter Kit RENU Multiplus Multi-purpose 60ml	ME3124, ME3848, MF1849
	BIOTRUE Multiplus Multi-purpose 300ml	ME3693
	BIOTRUE Multiplus Multi-purpose 120ml	ME2438
	Starter Kit BIOTRUE Multiplus Multi-purpose 60ml	MF1642
Manufacturer	Bausch and Lomb INC	
Country of Origin	Italy	
Reference	https://bausch.co.uk/news/bausch-lomb-announces-voluntary-recall#:~:text=LAVAL%2C%20Quebec%2C%20May%2027%2C,conditioning%20solution%2C%20Boston%C2%AE%20Simplus	

For more information please contact Medical_Devices@nhra.bh

Recall 2021 0060

Device picture	
Reason of Recall	NHRA initiates this FSN due to the sterilization process of the mentioned lots can not be confirmed and doesn't comply with the standards.
Action should be taken	Please stop using the above mentioned lots and contact the authorized representative **Optica** at elena.feleciano@optica.net 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