

Rev 1: September 2018

**FSN Ref:** Manufacturer's ref number **FSCA Ref:** Manufacturer's ref number NNGYK90755/2025

**Date:** 02012026

**Urgent Field Safety Notice**  
**Device Commercial Name**

**For Attention of\*:** Identify either by name or role who needs to be aware of the hazard and/or take action. If this is multiple recipients then include full list.

<b>Contact details of local representative (name, e-mail, telephone, address etc.)*</b>
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**Urgent Field Safety Notice (FSN)**  
**Device Commercial Name**  
**Risk addressed by FSN**

<b>1. Information on Affected Devices*</b>	
1	1. Device Type(s)* Ultrasound gel for external use
1	2. Commercial name(s) Ultrasound gel 260 gr
1	3. Unique Device Identifier(s) (UDI-DI) 5999886483UHGEL4V
1	4. Primary clinical purpose of device(s)* External use for ultrasound examination
1	5. Device Model/Catalogue/part number(s)* Add as Appendix if necessary.
1	6. Software version Only where relevant.
1	7. Affected serial or lot number range 02.2027
1	8. Associated devices Within context of the FSCA eg for IVD reagents and platforms.

<b>2 Reason for Field Safety Corrective Action (FSCA)*</b>	
2	1. Description of the product problem* Laboratory control indicated bacteria infection
2	2. Hazard giving rise to the FSCA* The bacterium can cause problems in weakened human bodies.
2	3. Probability of problem arising Provide an indication (from incident data or prospective modelling) of the likelihood the problem will arise.
2	4. Predicted risk to patient/users From the output of the Health Hazard Evaluation indicate the anticipated risk (product of severity x probability) of patient/end user harm (direct or indirect).
2	5. Further information to help characterise the problem Include any further relevant statistics to help convey the seriousness of the issue.
2	6. Background on Issue Eg how the manufacturer became aware; brief details of relevant incidents; root cause if known; rationale for containment of problem to only affected devices; other risk mitigation or longer-term preventative action etc.
2	7. Other information relevant to FSCA This field may only contain additional information that is deemed necessary by the manufacturer to supplement information relevant to the FSCA.

<b>3. Type of Action to mitigate the risk*</b>	
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3.	<p><b>1. Action To Be Taken by the User*</b></p> <p> <input type="checkbox"/> Identify Device    <input type="checkbox"/> Quarantine Device    <input checked="" type="checkbox"/> Return Device    <input type="checkbox"/> Destroy Device  <input type="checkbox"/> On-site device modification/inspection  <input type="checkbox"/> Follow patient management recommendations  <input type="checkbox"/> Take note of amendment/reinforcement of Instructions For Use (IFU)  <input type="checkbox"/> Other    <input type="checkbox"/> None       </p> <p>Provide further details of the action(s) identified.</p>	
3.	2. By when should the action be completed?	Specify where critical to patient/end user safety
3.	<p>3. Particular considerations for: Choose an item.</p> <p>Is follow-up of patients or review of patients' previous results recommended? Choose an item.</p> <p>Provide further details of patient-level follow-up if required or a justification why none is required</p>	
3.	4. Is customer Reply Required? * (If yes, form attached specifying deadline for return)	Yes
3.	<p><b>5. Action Being Taken by the Manufacturer</b></p> <p> <input checked="" type="checkbox"/> Product Removal    <input type="checkbox"/> On-site device modification/inspection  <input type="checkbox"/> Software upgrade    <input type="checkbox"/> IFU or labelling change  <input type="checkbox"/> Other    <input type="checkbox"/> None       </p> <p>Provide further details of the action(s) identified.</p>	
3	6. By when should the action be completed?	Specify where critical to patient/end user safety
3.	7. Is the FSN required to be communicated to the patient /lay user?	Choose an item.
3	<p>8. If yes, has manufacturer provided additional information suitable for the patient/lay user in a patient/lay or non-professional user information letter/sheet?</p> <p>Choose an item.    Choose an item.</p>	

<b>4. General Information*</b>		
4.	1. FSN Type*	New
4.	2. For updated FSN, reference number and date of previous FSN	Provide reference and date of previous FSN if relevant
4.	<b>3. For Updated FSN, key new information as follows:</b>  Summarise any key difference in devices affected and/or action to be taken.	
4.	4. Further advice or information already expected in follow-up FSN? *	No
4	5. If follow-up FSN expected, what is the further advice expected to relate to:  Eg patient management, device modifications etc	
4	6. Anticipated timescale for follow-up FSN	For provision of updated advice.
4.	7. Manufacturer information (For contact details of local representative refer to page 1 of this FSN)	
	a. Company Name	<b>Rextra Ltd</b>
	b. Address	<b>1033 Budapest, Szentendrei ut 87.</b>
	c. Website address	<b>rextra.hu</b>
4.	8. The Competent (Regulatory) Authority of your country has been informed about this communication to customers. *Yes, reference NNGYK90755/2025	
4.	9. List of attachments/appendices:	If extensive consider providing web-link instead.
4.	10. Name/Signature	<b>Reichardt Geza</b>  

<b>Transmission of this Field Safety Notice</b>	
	<p>This notice needs to be passed on all those who need to be aware within your organisation or to any organisation where the potentially affected devices have been transferred. (As appropriate)</p> <p>Please transfer this notice to other organisations on which this action has an impact. (As appropriate)</p> <p>Please maintain awareness on this notice and resulting action for an appropriate period to ensure effectiveness of the corrective action.</p> <p>Please report all device-related incidents to the manufacturer, distributor or local representative, and the national Competent Authority if appropriate, as this provides important feedback.*</p>

Note: Fields indicated by \* are considered necessary for all FSNs. Others are optional.