Unimax Medical Systems Inc.
8F-2, No. 127, Lane 235, Pao Chiao Road, Xindian Dist,
New Taipei City 231, Taiwan
Tel: +886-2-8919-1698
www.unimaxmeds.com

Rev 1: September 2018
FSN Ref: FSN-EP24001 FSCA Ref: FCA-EP24001

Date: 02/02/2024

Urgent Field Safety Notice
Detachable Endo Retrieval Pouch

For Attention of*:Theatre Manager

Contact details of local representative (name, e-mail, telephone, address etc.)*
Mölnlycke Health Care AB Contact's Name: Geetha Sundaram Iranaveeran Email: vigilance@molnlycke.com Phone: +46 31 722 30 00 Country: Sweden Address: Gamlestadsvägen 3C – Gothenburg – Sweden Post Code: SE 402 52 Box 13080

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Detachable Endo Retrieval Pouch

1. **Information on Affected Devices***	
1.	1. Device Type(s)*
	Detachable Endo Retrieval Pouch Small (250-300ml) / 10mm introducer diameter Medium /Large (500-700ml)/ 10mm introducer diameter Extra Large (1150-1500ml)/12mm and 15 mm introducer diameter
1.	2. Commercial name(s)
	Detachable Endo Retrieval Pouch
1.	3. Unique Device Identifier(s) (UDI-DI)
	07323190272792 (model 899102) 07323190272808 (model 899103) 07323190272815 (model 899104) 07323190272907 (model 899112)
1.	4. Primary clinical purpose of device(s)*
	The detachable endo pocket is a device that is used to collect and extract specimens during laparoscopic surgery.
1.	5. Device Model/Catalogue/part number(s)*
	899102; 899103; 899104; 899112
1.	6. Software version
	n/a
1.	7. Affected serial or lot number range
	Please refer to the web-link for look up: https://reurl.cc/374Xe8
1.	8. Associated devices
	n/a

2 **Reason for Field Safety Corrective Action (FSCA)***	
2.	1. Description of the product problem*
	The mechanism of the listed article number operates in a way that the tube within detaches during the removal process. If the tube is not precisely fixed, part of the tube may stretch out from the opening after detachment and fall into the abdomen of the patient.
2.	2. Hazard giving rise to the FSCA*
	The reported incidence is potentially serious to patients as the extending part may fall into the cavity.
2.	3. Probability of problem arising
	Overall occurrence rate: within 0.0001
2.	4. Predicted risk to patient/users
	Prolonged surgery or surgical intervention
2.	5. Further information to help characterise the problem
	n/a
2.	6. Background on Issue
	The device is used to contain and remove specimen removed during laparoscopic surgery. The mechanism of the listed article number operates in a way that the tube within detaches during the removal process. If the tube is not precisely fixed, part of the tube may stretch out from the opening after detachment and fall into the abdomen of the patient. It was thus decided to proceed with a

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	field safety corrective action to replace the current version with an improved design variant thus reducing the potential for the tube stretching out / falling into the patient's abdomen.
2.	7. Other information relevant to FSCA
	n/a

	3. Type of Action to mitigate the risk*	
3.	**1. Action To Be Taken by the User***	
	☒ Identify Device ☒ Quarantine Device ☐ Return Device ☒ Destroy Device ☐ On-site device modification/inspection ☐ Follow patient management recommendations ☐ Take note of amendment/reinforcement of Instructions For Use (IFU) ☐ Other ☐ None Provide further details of the action(s) identified.	
3.	2. By when should the action be completed?	Estimated within 6 months
3.	3. Particular considerations for: n/a Is follow-up of patients or review of patients' previous results recommended? n/a Provide further details of patient-level follow-up if required or a justification why none is required	
3.	4. Is customer Reply Required? * (If yes, form attached specifying deadline for return)	Yes
3.	**5. Action Being Taken by the Manufacturer** ☒ Product Removal ☐ On-site device modification/inspection ☐ Software upgrade ☐ IFU or labelling change ☐ Other ☐ None Provide further details of the action(s) identified.	
3	6. By when should the action be completed?	• Distribution of the FSN scheduled for 1-3 weeks. • Expected return of customer reply form in 1-2 months. • Replacement of affected devices scheduled for 3-6 months depends on the quantity.
3.	7. Is the FSN required to be communicated to the patient /lay user?	No
3	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?	
	n/a n/a	

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	4. General Information*	
4.	1. FSN Type*	New
4.	2. For updated FSN, reference number and date of previous FSN	n/a
4.	**3.** For Updated FSN, key new information as follows:	
	n/a	
4.	4. Further advice or information already expected in follow-up FSN? *	Not planned yet
4	5. If follow-up FSN expected, what is the further advice expected to relate to:	
	n/a	
4	6. Anticipated timescale for follow-up FSN	n/a
4.	7. Manufacturer information (For contact details of local representative refer to page 1 of this FSN)	
	a. Company Name	Unimax Medical Systems Inc.
	b. Address	8F-2, No. 127, Lane 235, Pao Chiao Road, Xindian Dist., New Taipei City, Taiwan
	c. Website address	http://www.unimaxmeds.com
4.	8. The Competent (Regulatory) Authority of your country has been informed about this communication to customers. * Yes	
4.	9. List of attachments/appendices:	Please refer to the web-link for look up: https://reurl.cc/374Xe8
4.	10. Name/Signature	**Partheeban Chinnamuthu / Regulatory Specialist**
		

	Transmission of this Field Safety Notice
	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) Please transfer this notice to other organisations on which this action has an impact. (As appropriate) Please maintain awareness on this notice and resulting action for an appropriate period to ensure effectiveness of the corrective action. 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. *

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