

75 Corporate Drive Trumbull, CT, US 06611 T: +1 203 601 5200 www.coopersurgical.com

February 27, 2023

### URGENT: MEDIA FIELD SAFETY CORRECTIVE ACTION

SAGE™ Vitrification Media Kit

Dear Valued CooperSurgical Customer,

CooperSurgical, Inc. is issuing a Field Safety Corrective Action (FSCA) for one (1) lot number of its SAGE Vitrification Media Kit (Reference Number ART-8026, **LOT 211112-002333)** (referred to in this letter as the "Product").

Per the Product Instructions for Use, these Products are intended for the vitrification and containment of human oocytes and embryos (pronuclear zygotes through day three cleavage stage embryos and blastocyst stage embryos) in Assisted Reproductive Technology (ART) procedures. This Product is designed to be used in conjunction with the SAGE™ Warming Kit (ART-8031) for warming and recovery of specimens.

#### Reason for Recall:

It has come to CooperSurgical's attention that certain SAGE Vitrification Media Kits within **LOT 211112-002333** may contain mislabeled vials. Specifically, the impacted kits may contain vials labeled as Vitrification Solution (VS) but actually contain Equilibration Solution (ES).

#### Risk to Health:

Use of an impacted SAGE Vitrification Media Kit from **LOT 211112-002333** may impact the viability of oocytes/embryos.

#### **Actions to be Taken:**

### For CUSTOMERS and DISTRIBUTORS:

- Inspect your inventory for Product from this lot. Note the lot's labeled expiration date is November 12, 2022. Lot information can be found on the box and bottle labeling.
  - o If found, discontinue use / distribution of the Product immediately and quarantine.
- Please complete the appropriate included Acknowledgment Form and send it via email to <u>Recall@coopersurgical.com</u>.

#### For CUSTOMERS:

 Check your records to determine whether you utilized SAGE Vitrification Media Kits from LOT 211112-002333. If so, follow up with your local sales representative for more information regarding implications of its use.



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## For DISTRIBUTORS:

• If the Product has been distributed to your customers, send a copy of this letter and the Customer Acknowledgement Form to each of these customers.

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We regret any inconvenience caused by this FSCA. CooperSurgical is committed to high quality, safe and effective Products.

Where appropriate, Regulatory Agencies will be notified of this issue.

You may reach us at +1-203-601-5200 during normal operating hours of 09:00 - 17:00 M-FEST. Follow the phone prompt to enter extension 3300. Our email address is Recall@coopersurgical.com.

Sincerely,

•••

Sr. Post Market Surveillance Manager



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### **Customer Acknowledgement Form**

## IMMEDIATE RESPONSE REQUIRED - TIME-SENSITIVE ACTION NEEDED

Please complete this form and return it via email to <a href="Recall@coopersurgical.com">Recall@coopersurgical.com</a>.or via fax to +1.203.601.9870, ATTN: Product Surveillance.

Customer Account #:		Account Name:		
Street Address:		Country, Town, State, & Zip Code:		
Contact Name:	Phone Number:		Email address:	
I have read and understand the notice 27, 2023.	e instructions prov	ided in the letter	dated February □ Yes □ No	
AGE Vitrification Media Kit (part r		-	-	
Please check the appropriate boxe	-		f applicable.	
☐ We have NO record of use for		•		
			t / have contacted our local sales	
representative for more inform		•		
☐ We have the following affected affected Product for return to		acility and Will d	iscontinue use and quarantine the	
Part Number	Lot Nu	ımbers	Quantity of kits to be Returned	
ART-8026	211112	-002333		
Have any adverse events been associated with affected Product(s)?				
If yes, please explain:				
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If you have additional questions, please contact a CooperSurgical Product Surveillance representative at **+1.203.601.5200** Ext. **3300** or email us at <a href="mailto:Recall@coopersurgical.com">Recall@coopersurgical.com</a>.



Signature

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# **Distributor Acknowledgement Form**

# IMMEDIATE RESPONSE REQUIRED - TIME-SENSITIVE ACTION NEEDED

Please complete this form and return it via email to <a href="Recall@coopersurgical.com">Recall@coopersurgical.com</a> or via fax to +1.203.601.9870, ATTN: Product Surveillance.

Customer Account #:		Account Name:		
Street Address:		Country, Town, State, & Zip Code:		
Contact Name:	Phone Number:		Email address:	
have read and understand the notice 27, 2023.	instructions prov	ided in the letter d	ated February	□ Yes □ N
SAGE Vitrification Media Kit (part nu	mber: ART-8026,	LOT 211112-0023	33)	
	•	the table if applic	able.	
Please check the appropriate box be  ☐ We have NO kits of the affected I  ☐ We have the following affected P distribution and quarantine the a  Product/Part Number	ot at our facility. roduct at our facil ffected Product fo	ity and will discont	inue its	to be Returned
<ul> <li>We have NO kits of the affected I</li> <li>We have the following affected P distribution and quarantine the a</li> <li>Product/Part</li> </ul>	ot at our facility. roduct at our facil ffected Product fo	ity and will discont r return to Cooper	inue its Surgical.	to be Returned
<ul> <li>We have NO kits of the affected I</li> <li>We have the following affected P distribution and quarantine the a</li> <li>Product/Part Number</li> </ul>	ot at our facility. roduct at our facil ffected Product fo  Lot N  21111  mers:	ity and will discont r return to Cooper lumbers 2-002333	inue its Surgical.  Quantity of kits	to be Returned
We have <b>NO</b> kits of the affected I  We have the following affected P distribution and quarantine the a  Product/Part Number ART-8026  Quantity of devices shipped to custor	ot at our facility. roduct at our facil ffected Product fo  Lot N  21111  mers: ed to customers, p  Date and N	ity and will discont r return to Cooper lumbers 2-002333	inue its Surgical.  Quantity of kits  fthe following:	to be Returned

**Printed Name**