

URGENT: SAFETY NOTICE - FSN02-2021

Biopsybell medical devices

Type of action: Immediate stop of product use

For the attention of: Distributors, Clinical Personnel, End Users

This letter contains important information that requires your immediate attention.

Dear Customer,

BIOPSYBELL SRL is conducting a Field Safety Corrective Action requesting to STOP with immediate effect using the Biopsybell devices with codes and batches listed in annex to this communication.

Description of the problem

BIOPSYBELL SRL has been informed by one of its sterilization service suppliers of a potential risk of incorrect sterilization of product batches sterilized by this supplier.

As part of the product release procedure, Biopsybell systematically checks all the technical information associated with sterilization of each single batch together with other product manufacturing and test information and formally releases the product on the market only when the outcome is positive.

Due to wilful falsification of the sterilization data and certificates by the supplier, it was not possible to intercept and identify any sterility problems.

This advisory warning applies to the product codes and batches listed in the annex.

Description of the devices involved

The devices involved are all Biopsybell devices with the codes and batches listed in the annex. Biopsybell also uses another supplier for the sterilization service, who is NOT involved in this problem. Therefore, batches not listed in the annex can be used as they are not impacted by the sterilization problem.

Clinical impact

The use of non-sterile devices in a clinical environment may lead to an increased risk of infection, which in extreme cases may cause severe harm or potentially lethal conditions.



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BIOPSYBELL SRL has to date not received any notification of adverse events or harm to patients, which might be associated with this field safety corrective action.

Request to clinical users:

- 1. Immediately stop using all products belonging to the batches listed in the annex still in your possession and contact Biopsybell/Biopsybell Representative for product return/disposal and related replacement.
- 2. Use products with codes and batches not listed in the annex available in your warehouse.
- 3. Contact your local BIOPSYBELL representative to discuss possible product alternatives.
- 4. Contact BIOPSYBELL or your Biopsybell representative in case of urgency or need for immediate replacement.

Action requested from the local distributor and customer

- 1. Circulate this safety notice to anyone in your organisation that needs to be aware of the problem.
 - If the product has been distributed further, identify the facilities/organisations involved and immediately forward this notice.
- 2. Place all products of the impacted batches in quarantine, awaiting the product return/disposal and related replacement, then complete the customer response form returning it to international1@biopsybell.it or international2@biopsybell.it, or to your local Biopsybell representative, as soon as possible.

BIOPSYBELL corrective actions

Biopsybell makes itself available to receive the involved parts still available in your warehouse and to replace them, after completing the form below and prior agreements with Biopsybell.



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Contact person

Should you have any questions on this letter, please contact:

Information	Contact
Technical and commercial information for abroad	e-mail: international1@biopsybell.it telephone: +39-0535-27850 e-mail: international2@biopsybell.it
	telephone: +39-0535-27850

We wish to underline that BIOPSYBELL's main objectives have always been the safety of patients and operators and the supply of quality products in full compliance with the regulations in force. We regret any inconvenience caused, but are also aware that we have done everything possible to prevent this situation. Nonetheless, as this was a case of wilful falsification by an external third party and we consider ourselves the injured party, we could not have prevented what occurred despite every possible control on our part and on the part of the Competent Bodies in charge of this control at this supplier. Thank you in advance for the support provided to BIOPSYBELL to solve this problem in as fast and effective a way as possible.

MIRANDOLA, 07/04/2021

BIOPSYBELL SRL

VICE PRESIDENT OF THE BOARD OF DIRECTORS

CARLO RICCA PRANDI BELLINI



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Customer response form – FSN02-2021- BIOPSYBELL SRL medical devices

Read together with the safety notice FSNo2-2021 and return the completed and signed form as soon as possible to:

international1@biopsybell.it; international2@biopsybell.it

• I confirm that this communication has been read and understood and that all the

recommended actions have be	en implemented as requested.		
Tick the appropriate box below:			
□ We have some of the impacto	ad davisas listad in the appey		
☐ We have <u>none</u> of the impacte	ed devices listed in the aimex		
OR			
☐ We have in our possession unit the following units have been p	ts of the impacted devices liste placed in quarantine and will NC	d in the annex and confirm that OT be used	
CODE	BATCH	QUANTITY IN QUARANTINE	



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Facility name:	
Department (if applicable):	
Address:	
Postal code:	City:
Contact name:	
Professional role within the	
organisation:	
Contact phone number:	Contact e-mail address:
Signature and stamp:	Date:

This form must be returned to BIOPSYBELL before this action can be considered closed for your facility.

PRODUCT CODE	PRODUCT BRAND NAME	PRODUCT DESCRIPTION	QUANTITY	PRODUCT BATCH
DKR1716CDFL10	DISKOM	DISKOM PERCUTANEOUS DISCECTOMY DEVICE 17G 16CM CENTIM. D. CONFEZIONAM. FORO LATERALE 10MM STERILE	10	200130
RESWB1112C	RENOVA SPINE KYPHOPLASTY WORKING CANNULA AND BEVEL TIP	RENOVA SPINE KYPHOPLASTY WORKING CANNULA AND BEVEL TIP 10G 12CM CENTIM. STERILE	1	2001105
KRVT1115200CBP-A	VERTEBROPLASTIC	VERTEBROPLASTIC 11G 15CM AGO VERTEBROPLASTICA P.TA BEVEL PLUS STERILE	1	2002542
GUN1820EC-OS	ESTER	AGO ESTER ONE SHOT 18G 20CM AUTOMATICOECOGENATO/CENTIMETRATO STERILE	7	2003656
DKR1716CDFL10	DISKOM	DISKOM PERCUTANEOUS DISCECTOMY DEVICE 17G 16CM CENTIM. D. CONFEZIONAM. FORO LATERALE 10MM STERILE	10	2003752
GUN1820EC-OS	ESTER ONE SHOT	AGO ESTER ONE SHOT 18G 20CM AUTOMATICOECOGENATO/CENTIMETRATO STERILE	5	20061503
GUN1825ECCH2-OS	ESTER 13 ONE SHOT	AGO ESTER 13 ONE SHOT 18G 25CM ECO/CENTIMET. CON AGO CHIBELL 22G 20CM STERILE	25	20061497
GUN1825EC-OS	ESTER ONE SHOT	AGO ESTER ONE SHOT 18G 25CM AUTOMATICOECOGENATO/CENTIMETRATO STERILE	25	20061496
GUN1820ECCH2-OS	ESTER 13 ONE SHOT	AGO ESTER 13 ONE SHOT 18G 20CM ECO/CENTIMET. CON AGO CHIBELL 22G 20CM STERILE	250	20071560
GUNTR1410EC	ESTER&INTRODUTTORE	AGO ESTER+INTRO 14G10CM AUTO ECO/CENT.STERILE	5	20071782
GUNTR1410EC-OS	ESTER ONE SHOT & INTRODUTTORE	ESTER ONE SHOT & INTRODUTTORE 14G 10CMSTERILE	5	20071778
BAN1112HCDTW	BONE ACCESS NEEDLE	BONE ACCESS NEEDLE 11G 12CM CANN CORONA E STILET TROCAR BIANCO DOPP CONF STERILE	1.000	20071820
BAN1112HCDBPW	BONE ACCESS NEEDLE	BONE ACCESS NEEDLE 11G 12CM CANN CORONA STIL.BEVEL PLUS BIANCO D.CONF. STERILE	1.000	20071852
03.51.10.0172	SCREWFIX	SCREW FIX 16G 35MM FENESTRATED OPEN CANNULA-2 CANNULE PER BUSTA,D.BUSTA ST	200	20081978
RESWD1115C	RENOVA SPINE KYPHOPLASTY WORKING CANNULA AND DIAMOND TIP	RENOVA SPINE KYPHOPLASTY WORKING CANNULA AND DIAMOND TIP 11G 15CM CENTIM. STERILE	500	2101323N