

Urgent Field Safety Notice  
Several Reference and batches of SAFIL & NOVOSYN  
Return of the Medical Device to the manufacturer  
Att. Users of above product

June 6<sup>th</sup>, 2018

Dear Sir or Madam,

B. Braun Surgical, S.A. is voluntarily recalling several reference/batches of Safil® and Novosyn®, a sterile, absorbable surgical multifilament sutures.

#### **Description of the medical device deficiency**

From an internal non conformity, the company detected that some units of the mentioned products have the pack damaged, as a consequence the product sterility is compromised.

#### **Potential harms associated are**

Wound infection (e.g. endomyometritis, localized/generalized peritonitis), abscess formation, adhesions, risk of wound dehiscence, sepsis that could lead to life-threatening injury.  
Treatment or reoperation might be necessary.

We would not recommend any specific monitoring of the patients that have been treated with the involved products since there is a risk of infection innate in any type of surgery. The hospital should act according to their established protocol for such complications.

#### **Identification of affected medical devices**

Reference name: **SAFIL®** (Several references affected, see attachment)  
Reference and batch number: Detailed list in the attachment

Reference name: **NOVOSYN®** (Several references affected see attachment)  
Reference and batch number: Detailed list in the attachment

## B. Braun Surgical, S.A.

### Actions to be taken

Please identify and quarantine if you still have the listed product in your warehouse.

Please check with your customers if they still have the listed product in their warehouse. If yes, ask them to send the product back to you immediately.

Once you have all affected units for return contact us for the management of the material.

Please, fill out the attached "Recall Confirmation Form" and send the completed form to us by July 6<sup>th</sup>, 2018.

This notice needs to be passed on all those who need to be aware within your organization and to any organization where the potentially affected devices have been transferred.

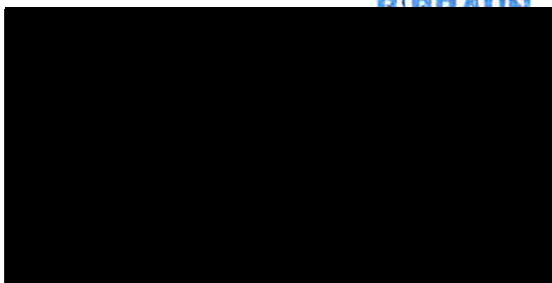
If you have any questions regarding this voluntary product recall, please contact us at the e-mail: [vigilance\\_CT@bbraun.com](mailto:vigilance_CT@bbraun.com).

We inform you that in accordance with the European Guidelines this recall has to be reported to the Competent Authority. Please check your national regulations and proceed accordingly.

We apologize the inconveniences we might have caused.

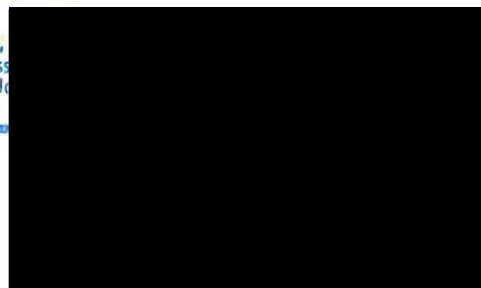
Thank you for your cooperation.

Yours faithfully,



B. Braun Surgical, S.A.

B. Braun Surgical,  
Carretera de Terrassa  
0191 Rubí (Barcelona)  
T: 93 586 62 00



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