



L'Union, 19<sup>TH</sup> of October 2016

**For the attention of the General Manager and/or of the Medical Device Vigilance Manager**

Re: Product recall

Dear Sir or Madam,

Following a validation verification of our sterilization procedures on the premises of one of our providers, we wish to inform you that as a precautionary measure we are making a voluntary product recall concerning the following batches:

ARTICLE	REFERENCE	BATCH	QUANTITY

We identified a risk in connection with sterilization conformity which could lead to infection in the days following surgery.

The cause of the incident is diagnosed and we have taken the necessary corrective steps to prevent recurrence.

In the context of this recall procedure, we are requesting a return of products from the batches quoted above to be sent to us at the following address:

**KASIOS SAS**  
18 chemin de la Violette  
31240 L'Union - FRANCE

As soon as we receive these products, they will be replaced free of charge.

Please address any queries concerning the recall to the KASIOS Medical Device Vigilance Unit:

Mr. Alain LERCH - Medical Device Vigilance Manager: [alain.lerch@kasios.com](mailto:alain.lerch@kasios.com) – +33(0)6.80.98.65.44

Mrs Adeline FRANCOIS - Medical Device Vigilance Deputy: [adeline.francois@kasios.com](mailto:adeline.francois@kasios.com) – +33(0)5 34.27.33.25

We much regret the inconvenience caused by this recall procedure. We wish to express our sincere apologies and the hope that we may look forward to enjoying your continued trust.

Yours sincerely,

Alain LERCH  
Medical Device Vigilance Manager