



# Declaration of Conformity for Medline US products

**Declaration of conformity n°** DC\_Sensicare PI  
**Revision n°** 01  
**Technical file #** CE2009005 (MDD) and CEP2009005 (PPE)

<b>Legal Manufacturer</b>	<b>Medline Industries, Inc.</b> Three Lake Drive, Northfield, Illinois 60093 USA
<b>EC Representative</b>	<b>Medline International France SAS</b> 5 rue Charles Lindbergh, 44110 Châteaubriant - France
<b>Product range</b>	Surgical Gloves – Sensicare PI
<b>Product codes</b>	See attached list
<b>Classification</b>	Medical Device class IIa Sterile - rule 6 Personal Protective Equipment Category III
<b>GMDN Codes</b>	56293 Polyisoprene surgical glove, non-powdered

## European Union Regulations:

<b>European Representative</b> We herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the following EU Council Directive(s) as transposed into national laws.	
<b>Applicable directive:</b>	Medical Devices Directive: Council Directive 93/42/EEC of 14 June, 1993 as amended.
<b>Annex 93/42/EEC</b>	Annex II
<b>Notified Body</b>	BSI The united Kingdom – CE 0086
<b>Certificate n°</b>	CE 557201
<b>First Issued (Place/Date)</b>	The United Kingdom, 20 <sup>th</sup> January, 2010
<b>Applicable standards</b>	See Technical File CE2009005
<b>Applicable directive:</b> Personal Protective Equipment: Council Directive 89/686/EEC of 21 December 1989.	
<b>EC-type examination per Article 10</b>	
Certificate n°	CE 664227
Notified Body (name/number/address)	BSI The united Kingdom – CE 0086
First Issued (Place/Date)	The United Kingdom, 8 <sup>th</sup> February 2017
Applicable standards	See Technical File CEP2009005
<b>Conformity assessment procedure per Article 11</b>	
Article 11 A or B	11A
Certificate n°	CE 651845
Notified Body (name/number/address)	BSI The united Kingdom – CE 0086
First Issued (Place/Date)	The United Kingdom, 8 <sup>th</sup> June 2016

## Australian Regulations:

This is a declaration of conformity made under clause 1.8 of Schedule 3 to the Therapeutic Goods (Medical Devices) Regulations 2002.  
 Each kind of medical device to which the system has been applied complies with the applicable provisions of the essential principles, the classification rules, and the full quality assurance procedures, at each stage, from the design of the device until its final inspection before being supplied.

All supporting documentation is retained at the premises of the manufacturer.

### **Authorised Signatory:**

**Kenneth Smith**  
 International Quality Operations Manager

44110 Châteaubriant - France  
 Place

20/11/2017  
 Date

05.INT.02

Duration of archiving: End of life and support period

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List of Medline US finished products for declaration of conformity

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Revision n°00  
Effective date: 21 Oct 16  
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Surgical gloves - Sensicare PI

DoC nr: DC Sensicar Rev: 01

Product number	Description	GMDN Code	Class	PPE Class
MSG9055	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III
MSG9060	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III
MSG9065	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III
MSG9070	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III
MSG9075	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III
MSG9080	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III
MSG9085	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III
MSG9090	Sensicare PI, Surgical glove, Sterile, Synthetic, Polyisoprene, Powder-Free	56293	Ila	III

\* GMDN code with description

GMDN Code

56293 Polyisoprene surgical glove, non-powdered