


Manufacturer's declaration of conformity

Name and address	Care of Sweden AB Visit: Fabriksgatan 5A, SE-514 33 Tranemo, Mail: Box 146, SE-514 23 Tranemo Phone: +46 (0)771 106 600, Fax +46 (0)325 128 40 info@careofsweden.se
SRN	Not available yet.
Product group	CuroCell CLP reactive air mattresses
Basic UDI-DI	7331345A0020B0021G2
EMDN code	V080702 – NON ACTIVE ANTI-DECUBITUS MEDICAL MATTRESSES
Classification	Class I, according to Annex VIII (MDR (EU) 2017/745, rule 1
Product/Device name	CuroCell® S.A.M. PRO mattress
Article/Part number	List of Sales configuration CuroCell® S.A.M. PRO mattress (SAMP-CE-010)
Common Specifications	There are no applicable Common Specifications.
Mark of compliance	CE
Declaration	<p>We declare under our sole responsibility as Manufacturer that the product(s) listed above conform to the requirements of the MDR (EU) 2017/745. The product(s) meet(s) the relevant General Safety and Performance Requirements of Annex I.</p> <p>The conformity assessment procedure was performed following Annex II to III of MDR (EU) 2017/745. Any modification to the device, not authorized by us, will invalidate this declaration.</p>
Valid from:	2020-07-01
Updated:	2021-09-29
Manufacturer's signature	 Magnus Högberg, CEO
Date	2021-09-29



Manufacturer's declaration of conformity

Name and address	Care of Sweden AB Visit: Fabriksgatan 5A, SE-514 33 Tranemo, Mail: Box 146, SE-514 23 Tranemo Phone: +46 (0)771 106 600, Fax +46 (0)325 128 40 info@careofsweden.se
SRN	Not available yet.
Product group	CuroCell CLP reactive air mattress systems
Basic UDI-DI	7331345A0020B0020FY
EMDN code	V080701 - ACTIVE ANTI-DECUBITUS MEDICAL MATTRESSES
Classification	Class I, according to Annex VIII (MDR (EU) 2017/745, rule 13
Product/Device name	CuroCell® S.A.M. PRO
Article/Part number	List of Sales configuration CuroCell® S.A.M. PRO (SAMP-CE-010)
Common Specifications	There are no applicable Common Specifications.

Mark of compliance**Declaration**

We declare under our sole responsibility as Manufacturer that the product(s) listed above conform to the requirements of the MDR (EU) 2017/745. The product(s) meet(s) the relevant General Safety and Performance Requirements of Annex I.

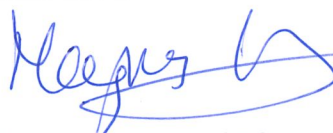
The conformity assessment procedure was performed following Annex II to III of MDR (EU) 2017/745. Any modification to the device, not authorized by us, will invalidate this declaration.

Valid from:

2020-09-25

Updated:

2021-09-29

Manufacturer's signature

Magnus Högberg, CEO

Date

2021-09-29



Manufacturer's declaration of conformity

Name and address	Care of Sweden AB Visit: Fabriksgatan 5A, SE-514 33 Tranemo, Mail: Box 146, SE-514 23 Tranemo Phone: +46 (0)771 106 600, Fax +46 (0)325 128 40 info@careofsweden.se
SRN	Not available yet.
Product group	CuroCell CLP reactive air mattress systems
Basic UDI-DI	7331345A0020B0020FY
EMDN code	V080701 - ACTIVE ANTI-DECUBITUS MEDICAL MATTRESSES
Classification	Class I, according to Annex VIII (MDR (EU) 2017/745, rule 13
Product/Device name	CuroCell® S.A.M. Switch
Article/Part number	List of Sales configuration CuroCell® S.A.M. Switch (SAMSW-CE-010)
Common Specifications	There are no applicable Common Specifications.
Mark of compliance	CE
Declaration	<p>We declare under our sole responsibility as Manufacturer that the product(s) listed above conform to the requirements of the MDR (EU) 2017/745. The product(s) meet(s) the relevant General Safety and Performance Requirements of Annex I.</p> <p>The conformity assessment procedure was performed following Annex II to III of MDR (EU) 2017/745. Any modification to the device, not authorized by us, will invalidate this declaration.</p>
Valid from:	2020-09-25
Updated:	2021-09-29
Manufacturer's signature	 Magnus Högberg, CEO
Date	2021-09-29



Manufacturer's declaration of conformity

Name and address	Care of Sweden AB Visit: Fabriksgatan 5A, SE-514 33 Tranemo, Mail: Box 146, SE-514 23 Tranemo Phone: +46 (0)771 106 600, Fax +46 (0)325 128 40 info@careofsweden.se
SRN	Not available yet.
Product group	CuroCell CLP reactive air mattress systems
Basic UDI-DI	7331345A0020B0020FY
EMDN code	V080701 - ACTIVE ANTI-DECUBITUS MEDICAL MATTRESSES
Classification	Class I, according to Annex VIII (MDR (EU) 2017/745, rule 13
Product/Device name	Pump CuroCell® S.A.M. PRO
Article/Part number	List of Sales configuration Pump CuroCell® S.A.M. PRO (SAMP-CE-010)
Common Specifications	There are no applicable Common Specifications.
Mark of compliance	CE
Declaration	<p>We declare under our sole responsibility as Manufacturer that the product(s) listed above conform to the requirements of the MDR (EU) 2017/745. The product(s) meet(s) the relevant General Safety and Performance Requirements of Annex I.</p> <p>The conformity assessment procedure was performed following Annex II to III of MDR (EU) 2017/745. Any modification to the device, not authorized by us, will invalidate this declaration.</p>
Valid from:	2020-09-25
Updated:	2021-09-29
Manufacturer's signature	 Magnus Högberg, CEO
Date	2021-09-29

