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	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:	2020-07-01
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	CE
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	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	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	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