YOUR DISPOSABLE EXCELLENCE

DICHIARAZIONE DI CONFORMITA'

CE

Il fabbricante:
Ragione sociale: DISPOTECH SRL
Sede: GORDONA (SO)- VIA AL PIANO, 29
Cod. fiscale/Part.IVA: 00672170149
R.E.A.: 47213

DICHIARA che il prodotto: GHIACCIO SPRAY

è conforme ai requisiti essenziali del Dgs.46/1997 che recepisce la Direttiva Dispositivi Medici 93/42/CEE e s.m.i. compresa la direttiva 2007/47/CE recepita con D.lgs. 37/2010

Classe del dispositivo: IIa allegato IX regola 9

In rispondenza a quanto stabilito dalle norme armonizzate:
Decreto Legislativo 24.02.1997, n.46 - Attuazione della direttiva 93/42/CEE, concernente i dispositivi medici
MDD 93/42/CEE e successive modifiche integrative (es.: Direttiva Europea 2007/47/CE) - Direttiva dispositivi medici
REGOLAMENTO (UE) 2023/607 DEL PARLAMENTO EUROPEO E DEL CONSIGLIO del 15 marzo 2023 che modifica i regolamenti (UE) 2017/745 e (UE) 2017/746 per quanto riguarda le disposizioni transitorie per determinati dispositivi medici e dispositivi medico-diagnostici in vitro
MEDDEV 2.12-1 rev.8 Linee guida sul sistema di vigilanza dei dispositivi medici
UNI EN ISO 13485 :2016 Dispositivi medici – Sistemi di gestione della qualità – Requisiti per scopi regolamentari
UNI CEI EN ISO 15223-1:2017 Simboli da utilizzare nelle etichette del dispositivo medico, nell' etichettatura e nelle informazioni che devono essere fornite – parte I requisiti generali
UNI CEI EN ISO 14971:2020 Applicazione della gestione dei rischi ai dispositivi medici
UNI CEI EN 1041:2013 Informazioni fornite dal fabbricante di dispositivi medici
ISO 10993-1:2018 Valutazione biologica dei dispositivi medici
MEDDEV 2.7.1 Linee guida sulla valutazione clinica dei dispositivi medici
DECRETO LEGISLATIVO 14 MARZO 2003 n°65 Attuazione delle direttive 1999/45/CE e 2001/60/CE relative alla classificazione, all'imballaggio e all'etichettatura dei preparati pericolosi
DIRETTIVA 2013/10/UE DELLA COMMISSIONE del 19 marzo 2013 che modifica la direttiva 75/324/CEE del Consiglio per il ravvicinamento delle legislazioni degli Stati membri relative agli aerosol al fine di adattare le sue disposizioni concernenti l'etichettatura al regolamento (CE) n. 1272/2008 del Parlamento europeo e del Consiglio relativo alla classificazione, all'etichettatura e all'imballaggio delle sostanze e delle miscele
DIRETTIVA (UE) 2016/2037 della Commissione, del 21 novembre 2016, che modifica la direttiva 75/324/CEE del Consiglio per quanto riguarda la pressione massima ammissibile dei generatori aerosol e adegua le sue disposizioni concernenti l'etichettatura al regolamento (CE) n. 1272/2008 del Parlamento europeo e del Consiglio relativo alla classificazione, all'etichettatura e all'imballaggio delle sostanze e delle miscele
DIRETTIVA 75/324/CEE direttiva madre aerosol attuazione DPR 21/07/1982 n°741
NORME AEROSOL AIA

Riferimento certificato di conformità CE n°: 26556
Emesso dall'Ente Notificato n°: 0546 CERTIQUALITY SRL- Via Gaetano Giordano, 4- 20123 MILANO- ITALY
Prima emissione: 20/02/2004
Emissione corrente: 22/03/2021

Luogo: Gordona

Data emissione documento_24/01/2024_

Il legale rappresentante
(responsabile rilascio del prodotto)

DISPOTECH S.R.L.
Via Al Piano, 29
23020 GORDONA (SO)
Tel. 0039 0343 36711 Fax 0039 0343 36567
Codice Fiscale e Partita Iva 00672170149

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Manufacturer's Declaration

in relation to Regulation (EU) 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices, in particular with respect to

- the validity of certificates issued under Council Directive 90/385/EEC on Active Implantable Medical Devices (AIMDD) or Council Directive 93/42/EEC on Medical Devices (MDD) (Directive Certificates) *and/or*[1]
- the compliance of the devices and us as their manufacturer with the conditions for the continued placing on the market and putting into service

Manufacturer name	Dispotech srl
Manufacturer address and contact details	Via al piano, 29 23020 GORDONA (SO) Italia Tel. +39 0343 36711 Fax. +39 0343 36567 e-mail: info@dispotech.it
Single Registration Number (SRN) (if available)	IT-MF-000010735

Authorised Representative name (if applicable)	N.A.
Authorised Representative address and contact details	N.A.
Single Registration Number (SRN) (if available)	N.A.

Notified body name (if applicable)	Certiquality srl □ See attached schedule
Notified body number (if applicable)	0546

[1] The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body.

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Tel. +39 0343 36711 - Fax. +39 0343 36567 - e-mail: info@dispotech.it - www.dispotech.com
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	□ See attached schedule
Directive Certificate number(s) to which this confirmation is made (if applicable)	26556 □ See attached schedule
Original expiry date as indicated on the Directive Certificate prior to the extension of the validity (if applicable)	28/01/2024 □ See attached schedule
End date of extended validity/transition period	31/12/2028 □ See attached schedule

We, as the manufacturer declare under our sole responsibility:

- for the above listed **Directive Certificate** (or see attached schedule, if multiple certificates) the conditions for the legal extension of validity as required in Article 120.2 of the MDR are met *and/or*[2]
- the listed **device(s)** in the attached schedule and we as their manufacturer are in compliance with the conditions listed in Article 120.3c of the MDR for continued placing on the market and putting into service,

namely by fulfilling the following conditions:

➢ **Directive Certificate(s)** as listed above or in the attached schedule

- Directive Certificate(s) covering the listed device(s) was/were issued after 25 May 2017, was/were valid on 26 May 2021 and have not been withdrawn afterwards.

  Choose applicable statements:

  ☐ Expired *before* 20 March 2023:

  ☐ Before the original date of expiry as indicated on the Directive Certificate(s), we and the notified body have signed written agreement(s) in accordance with Section 4.3, second subparagraph of Annex VII to this Regulation for the conformity assessment(s) in respect of the device(s) covered by the expired certificate(s) or in respect of a device(s) intended to substitute that/those device(s), or

  ☐ A Competent Authority has granted a derogation from the applicable conformity assessment procedure in accordance with Article 59(1) MDR (may be provided upon request), or

[2] The first condition is not applicable in case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body

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☐ A Competent Authority has required the manufacturer, in accordance with Article 97(1) MDR, to carry out the applicable conformity assessment procedure (may be provided upon request)

Choose one of the following statements only if a derogation per Article 59(1) or a requirement per Article 97(1) has been granted by a Competent Authority:

☐ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

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☒Expired/expires *after* 20 March 2023:

Choose one applicable statement:

☒Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitute(s) and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

- **Upclassified devices**

In case of devices for which the conformity assessment procedure pursuant to MDD did not require the involvement of a notified body, for which the declaration of conformity was drawn up prior to 26 May 2021 and for which the conformity assessment procedure pursuant to this Regulation requires the involvement of a notified body:

Choose one applicable statement:

☐ Formal application(s) to the notified body in accordance with Section 4.3, first subparagraph of Annex VII MDR for conformity assessment has/have been made or will be made/submitted by us to a notified body no later than 26 May 2024 for the device(s) listed in the attached schedule or its/their substitutes and signed written agreement(s) is/will be in place in accordance with Section 4.3, second subparagraph of Annex VII MDR before 26 September 2024.

☐ We do not intent to lodge an application for conformity assessment by 26 May 2024, therefore the transition period will end on 26 May 2024.

- **Quality Management System (QMS)**

Choose one applicable statement:

☐ A QMS in accordance with Article 10(9) MDR will be put in place by no later than 26 May 2024.

☒A QMS in accordance with Article 10(9) MDR is in place.

☐ A notified body has issued the attached certificate for the MDR-compliant QMS.

- **Device(s) as listed in the attached schedule**

- The device(s) continue to comply with the AIMDD or MDD.
- There are no significant changes in the design and intended purpose.
- The device(s) do not present an unacceptable risk to health or safety of patients, users or other persons, or to other aspects of the protection of public health.

Signed for and on behalf of the manufacturer:

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Full Company Name DISPOTECH SRL

Location & Date GORDONA, 27/09/2023

Signature, Print Name, Title Mr MASSIMO MORTAROTTI, LEGAL REPRESENTATIVE

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Schedule of Devices

The above Manufacturer's Declaration is valid for the following devices:

Identification of the device(s)[3] (e.g., device name, family/group name device model or catalogue number)	**Directive Certificate number(s) to which this confirmation is made** (if applicable)	**Original expiry date as indicated on the Directive Certificate (s) prior to the extension of the validity** (if applicable)	**Notified Body name and number that issued the Directive Certificate** (if applicable)	**Notified Body name and number where the MDR application was lodged/contract signed** (if applicable)	**End date of extended validity / transition period**	**Substitute Device(s)** (if applicable)
INSTANT COLD PACK	26556	28/01/2024	0546	1936	31/12/2028	n.a.
ICE SPRAY	26556	28/01/2024	0546	1936	31/12/2028	n.a.

[3] for devices with AIMDD/MDD certificate(s) the identification should be as in the certificate, and only if the certificate has a generic scope it should be as defined above)

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