INTCO MEDICAL(HK) CO., LTD.



FLAT/RM 19C, LOCKHART CENTRE, 301-307 LOCKHART ROAD, WAN CHAI. HONG KONG

Tel:+86 511 83174088 Fax: +86 511 83174188

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Document Number: CE-DC-001 Version: A/5

EC Declaration of Conformity

Manufacturer: whose single Authorized Representative:

INTCO Medical (HK) Co., Limited FLAT/RM 19C, LOCKHART CENTRE, 301-307 LOCKHART ROAD, Borkstrasse 10, 48163 Muenster, WAN CHAI, HONG KONG

> Tel:+86 511 83174088 Fax: +86 511 83174188

We, the manufacturer, herewith declare that the products

MedNet EC-REP GmbH

Germany

Tel: +49-251-32266-61 Fax: +49-251-32266-22

Cold Packs

(HypaCool Instant Cold Pack, Instant Ice Pack Mini, Instant Plus Ice Pack, Instant Ice Pack, INSTANT COLD COMPRESS, HOT & COLD COMPRESS, Instant Soothe COLD pack, Cold Pack, Instant Freeze, Instant cold pack, Instant Cold Compress, cold pad, Instant Cool Pack, Instant Perineal Cold Pack, Cool Power Compress, Cool Pack Mini, Cool Pack Midi, Cool Pack Maxi, Cold Compress)

Model codes

(T129525, Q2281, 7710, 713, 710, ICE1069, 151168, 122864, INSD, N14843, N14844, 09135, 09136, 007909, 100196.0,99228,99242,ICE0609,ICE0506,ICE1125,ICE0505,751520-Q,751315-Q,KID2,122865,132935,HY-303,913-001 .30649,193349.2,1130109,I114-15*23,I1114-13*15,04-018M,25.00200)

UMDNS Code: 10932

meet the provisions of the Council Directive 93/42/EEC which apply to them.

The medical device has been assigned to class II a according to Annex IX, Rule 9 of the Directive 93/42/EEC. It bears the mark



The product concerned has been manufactured under a quality management system according to Annex V and Annex VII of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

> **TÜV Rheinland LGA Products GmbH** Tillystraße 2, 90431, Nürnberg, Germany Certificate No.: DD 2068388-1

> > Issue date: 11.09.2020 Expiry date: 26.05.2024

following the procedure relating to the EC Declaration of Conformity set out in Annex V and Annex VII of Directive 93/42/EEC.

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This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

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