



CERTIFICATE OF CE (MDD) NOTIFICATION

Ref. No.: CH 2702-2014

Date: 11/04/2014

Order No.: CH 2555-2014

THIS IS TO CERTIFY THAT, ACCORDING TO THE EUROPEAN COUNCIL DIRECTIVE 93/42/EEC WE, HERE AT OBELIS S.A. (O.E.A.R.C.) PERFORMED ALL NOTIFICATION DUTIES AND RESPONSIBILITIES AS THE EUROPEAN AUTHORIZED REPRESENTATIVE (EC REP) OF:

NAME: KT HEALTH, LLC

ADDRESS: D/B/A KT TAPE, 7 SOUTH 1550 WEST, SUITE #600 LINDON, UT 84042, U.S.A

AS STIPULATED AND DEMANDED BY THE AFOREMENTIONED DIRECTIVE.

The Manufacturer declares that the Class I * device complies with the Directive including all essential requirements.

The Manufacturer has provided Obelis s.a. (O.E.A.R.C.) with all the appropriate declarations as per the European Council Directive 93/42/EEC article 14 requirements, including the EC Declaration of Conformity (according to Annex VII) confirming that their Class I medical devices, as stipulated here above, are fulfilling the applicable requirements of the European Council Directive 93/42/EEC.

The notification of the following medical devices has been completed by Obelis s.a. (O.E.A.R.C.) on the 21/03/2014 in compliance with the European Council Directive 93/42/EEC and 2007/47/EC - article 14 requirements

CLASS I MEDICAL DEVICE: PLEASE SEE ANNEX A - LIST OF DEVICES (1 PAGE, 1 DEVICE)

As of the 22/03/2014, and as long as the Manufacturer will continue complying with the hereabove mentioned requirements**, he therefore:

- Is required to affix the CE marking on this device;
- May place this device in the European community Territory.

OBELIS s.a. - O.E.A.R.C

Registered address :
Bd Général Wahis 53
1030 Bruxelles

Tel. +32 2 732 59 64 - Fax +32 2 732 60 03

G. Elkayam
Mr. G. Elkayam CEO
Obelis sa

date & stamp

Evelien Jonckheere
Brussels Enterprise
Commerce & Industry

date & stamp



SEEN
by the Brussels Chamber of Commerce
Evelien Jonckheere
Brussels, the 15 AVR. 2014

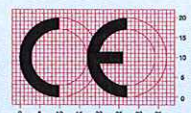


*also applicable to Class I s & m

** and provided that the product classification will not be rejected by the competent authorities

Obelis European Authorized Representative Center is a member of the European Association of Authorized Representatives (E.A.A.R.), ISO 9001 : 2008 and ISO 13485 : 2003 certified in accordance to the profession of a European Authorized Representative.

Registered Address: Bd. Général Wahis 53- 1030 Brussels | Registered Office Address: Av. de Tervueren 34 B44- 1040 Brussels - Belgium
T: + 32 (0) 2 732 5964 | F: + 32 (0) 2 732 6003 | Email: mail@obelis.net | Website: www.obelis.net



Annex A* – List of devices

(Article 9, section 1 of the Directive 93/42/EEC on medical devices)

No.	Generic Device Term	Commercial name	Class**	Rule	Catalogue reference number	Short description and intended use	GMDN code***
1	Elastic Sports Tape – Kinesiology Tape	KT TAPE	1	Rule 1	10000024	Kinesiology tape for pain relief and support	16866

* Annex A is part of the Agreement.

** The here above product list classification is based on the classification claim of the manufacturer and under its sole responsibility (MDD 93/42/EEC, article 2 & Annex IX; MEDDEV 2.4/1 Rev.9, chapter 3.1 & 3.3)

*** GMDN codes are mandatory information to perform the Notification

Manufacturer's Name

Obelis S.A.

BECI

SA

Signature: Ryan Dewey

Signature: [Signature]

Signature: _____

Date: 1-13-14

Date: 14/4/2014

Date: CHAMBRE DE COMMERCE ET D'INDUSTRIE DE BRUXELLES

Stamp:

Stamp: OBELIS s.a. - O.E.A.R.C
 Registered address :
 Bd Général Wahis 53
 1030 Bruxelles
 Tél. +32 2 732 59 54 - Fax +32 2 732 60 03

Stamp: Beci 15-04-2014
 KAMER VOOR HANDEL EN NIJVERHEID VAN BRUSSEL

SINCE 1988

OBELIS s.a. Anti-Counterfeiting Label