Certificate of CE-Registration

mdi Europa

This is to certify that, in accordance with either medical device Directive 93/42/EEC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Hiermit wird bestätigt, daß mdi Europa GmbH als Bevollmächtigter gemäß § 7 Medizinproduktegesetz (MPG/nationale Umsetzung der Richtlinie für Medizinprodukte 93/42/EWG bzw. 98/79/EG) für den Hersteller

Composite Resources, Inc.
C-A-T Rersources, LLC
485 Lakeshore Parkway
Rock Hill, SC 29730
USA

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

die Anzeigepflicht gemäß § 25 MPG für die nachfolgend aufgeführten Medizinprodukte erfüllt hat. Den angezeigten Medizinprodukten sind die folgenden Registrierdaten zugeordnet worden:

UMDNS Code – Class	Medical Device	Registration-No.
14072 – Class I	Tourniquet	DE/CA09/0760/613-Ä1
14075 – Class I	Tourniquets, Cardiovascular	DE/CA09/0760/614-Ä1
16907 – Class I	Cuffs, Tourniquet, Disposable	DE/CA09/0760/615-Ä1
16632 – Class I	Tourniquets, Strap	DE/CA09/0760/616-Ä1

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Der Hersteller hat mdi Europa alle für das erstmalige Inverkehrbringen von Medizinprodukten erforderlichen Dokumente vorgelegt. Dazu gehört die Konformitätserklärung, die bestätigt, daß die Produkte die grundlegenden Anforderungen der Richt-linie 93/42/EWG bzw. 98/79/EG erfüllen. Ein Sicherheitsbeauftragter gemäß § 31 MPG wurde bestellt.

January 2016

Werner Sander
President & CEO

Declaration of Conformity Certificate

We **Composite Resources, Inc. (Parent Co.)**
C-A-T Resources, LLC (Manufacturer)
Derek G Thompson, (CFO)
485 Lakeshore Parkway
Rock Hill, SC 29730 USA
803-366-9700

Declare with sole responsibility, that our product/s:

UMDNS Code	UMDNS Description	Classification

The Combat Application Tourniquet® (C-A-T®). UMDNS codes 14072, 14075, 16907, and 16632, a Class 1 device.

Meet, the essential requirements of EITHER Council Directive 93/42/EEC OR Council Directive 98/79/EEC pertaining to medical devices.

We hereby appoint mdi Europa GmbH, Langenhagener Str. 71, D – 30855 Hannover-Langenhagen, Germany to act as European Authorized Representative as explicitly defined in Article 1,
§ 2(g) of Directive 98/79/EEC.

Signed this day 2nd of January 2016 ..... Derek G Thompson

mdi Europa use only!

The necessary pre-requisites for placing the CE mark on the above mentioned products and marketing them in all Member States of the European Union, have thus been fulfilled.

Signed this day 14 of Jan 20 16

mdi Europa GmbH
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Fax +49-511. 39 08 99 39

mdiEuropa
THE MEDICAL DEVICE SERVICE-MANAGEMENT