

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 Resources, LLC
485 Lakeshore Parkway
Rock Hill, SC 29730
USA**

as stipulated and demanded by the aforementioned 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 Richtlinie 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:

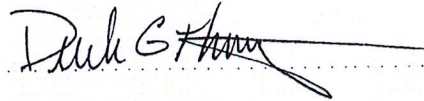
<u>UMDNS Code</u>	<u>UMDNS Description</u>	<u>Classification</u>
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The Combat Application Tourniquet® (C-A-T®). UMDNS codes 14072, 14075, 16907, and 16632, a Class 1 device.

**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

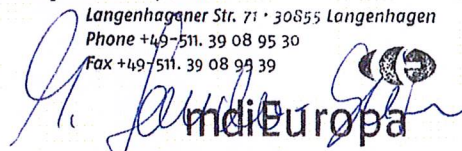


mdi Europa use only!

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

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mdiEuropa

THE MEDICAL DEVICE SERVICE-MANAGEMENT