

# 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:**

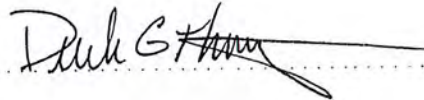
| <u>UMDNS Code</u> | <u>UMDNS Description</u> | <u>Classification</u> |
|-------------------|--------------------------|-----------------------|
|-------------------|--------------------------|-----------------------|

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

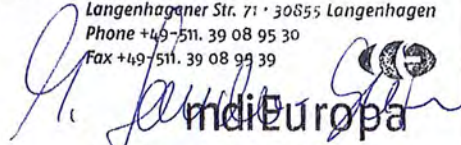


mdi Europa use only!

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 2016

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**mdiEuropa**

THE MEDICAL DEVICE SERVICE-MANAGEMENT