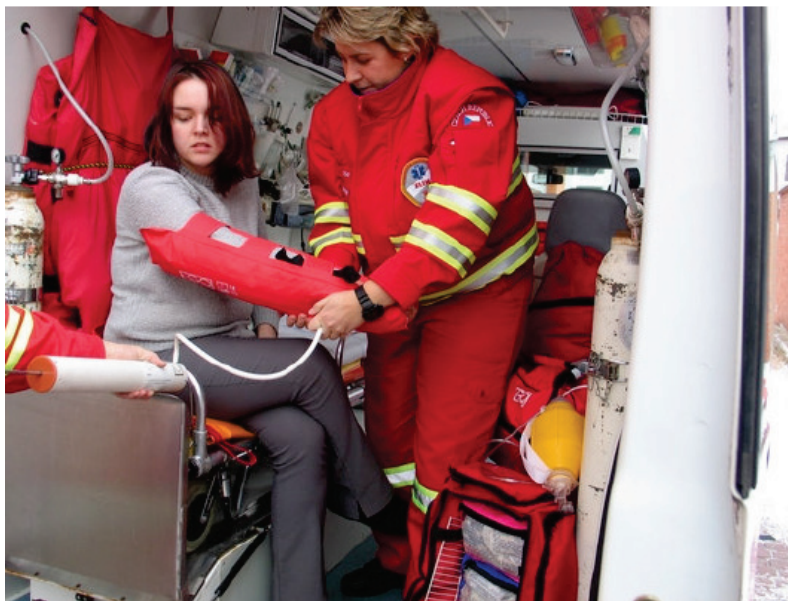


HANGING GRIPSACK UNDER THE HELICOPTERE

MADE BY COMPANY EGO ZLIN SPOL. S R. O.



BASIC INFORMATION ABOUT OUR COMPANY:

Company name: EGO Zlín, spol. s r.o. (Ltd.)
Address: U pekárny 438, 763 14 Zlín - Štípa,
the Czech Republic
Company ID: 469 024 73
VAT code: CZ 469 024 73
Director: Ing.Pavel Kostka
Sales manager: Ing. Marie Eibelová
Tel: + 420 577 100 038
Fax: + 420 577 914 363
E-mail: export@egozlin.cz



The producer company EGO Zlin has established and certified system of quality management ISO 9001:2009

Hanging gripsack under the helicoptere is made from polyamide textile materials, straps and ropes, using stitch technology. Fixing and suspending metal parts are used from the assortment of mountaineering and aeronautics. The bag serves for patient's protection against weather effects as well as prevention from catching cold.

The vacuum mattress EM – 10 is put at the bottom of the suspension bag. After laying and fixing of the patient over chest and hips, the vacuum pump is used for shaping and ensures such a manner the stiffness needed of the whole system.

The suspension bag is used for fast transport of injured person by aerial or mountain services, or for saving the persons at fires, mining or other disasters.

It consists of:

- ZV – 10 - Suspension fixation bag
- EM – 10 - Vacuum mattress
- EM – 20 - Vacuum pump with pedal



TYPE CERTIFICATE



Registration Number 058/05/07/02/0

issued for the producer:
EGO Zlín, spol. s r.o.
 Štípa 438
 CZ - 763 14 Zlín – Štípa
 Ident.-no.: 46902473
 for the product.

Name: **Device for transport of patient**
 Type Designation: **VP-10, VP-20, EZD-10, ET-10, ZV-10**
 Options: **-**
 Point of Production: **EGO Zlín spol. s r.o.**
Štípa 438, CZ - 763 14 Zlín - Štípa

at which the evaluation, certification and audit of the quality management system were carried in accordance with the certification system of the TÜV CZ according to the ČSN EN 45011:1999, whose results are given in the Report no. 0424/90/01/BT/IZ/S from April 18th, 2001, and the annual supervisory audit of the quality system, which results are given in the Report no. 0133/90/05/BT/IZ/S.

The mentioned product type fulfils the applicable requirements of following regulations/ normative documents that have underlain for its evaluation:
Decree of the government no. 338/2004 Coll. (Directive of the Council 93/42/EEC as amended by the 98/79/EC) setting the technical requirements for the medical devices.
ČSN EN 1441:1998 (EN 1441:1997)

This certificate is valid till: **April 30th, 2011**

Details and validity conditions are given in the annex to this certificate that is its integral part and contains 1 page.

Prague, February 24th, 2005



Ing. Ivo Dršťák
 Manager of the Certification
 Body

TÜV CZ s.r.o., Novoborská 894, 142 21 Prague 4 • CR Ident.-No: 63987121
 Municipal Court in Prague, volume C, insert no. 3443, entry date: July 20th, 1999
 Product Certification Body

F.0-028295 (08.05.04)

Annex to the certificate no. 058/05/07/02/0

1. The product sample was applied for the evaluation and certification on March 12th, 2001.
2. The certificate was issued on the technical basis of the manufacturer. See the Evaluation Report no 0424/90/01/BT/IZ/S from April 18th, 2001 and the Evaluation Report no 0133/90/05/BT/IZ/S from February 5th, 2005.
3. Detailed technical data characterizing the product type/sample: See the Evaluation Report no 0424/90/01/BT/IZ/S from April 18th, 2001
4. The list of important parts of the technical documentation: See the Evaluation Report no 0424/90/01/BT/IZ/S from April 18th, 2001
5. Validity conditions
 - The certificate shall only promote its holder, the said product and production places.
 - The transmission of this certificate to third parties is inadmissible as well as its using by them.
 - Any modifications of the product shall be announced to TÜV CZ to enable to decide about an additional conformity evaluation.
 - TÜV CZ performs the supervision over the regular functioning of the quality system on the basis of the concluded contract of inspection activity on schedule once yearly
 - This certificates can be renewed on demand
 - This certificate shall only be reproduced in full including all annexes.
 - The right to use the trademark TÜV CZ was established to this certificate.
 - The certificate holder undertakes to keep files about all eventually complaints concerned the conformity of the product with the requirements of regulations and directives and to release this files to the certification body TÜV CZ.
 - The not mentioned details (advertising, using of trademark and certificates) are regulated by the General conditions for product certification in the actual version.

2.2.2010