

CarboFlex™



PRODUCT DESCRIPTION¹:

CarboFlex® Odour Control Dressing is a sterile five-layer non-adhesive dressing with an absorbent wound contact layer (containing Alginate and Hydrofiber™) an odour absorbing charcoal pad, an absorbent padding layer and soft flocking outer contained within a peel pouch.

CarboFlex™ products

Dressing Size cm	CVT number	Pieces/package
8x15cm	403203	5
10x10cm	403202	10
15x20cm	403204	5



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INTENDED USE¹:

CarboFlex[®] is intended for the management of malodorous wounds. It may be used as a primary or secondary dressing

INTENDED USER¹:

CarboFlex[®] is intended to be used by healthcare professionals, patient caregivers and patients.

DIRECTIONS FOR USE¹:

Always check instructions for use. Please see IFU in product box before clinical use

ADDITIONAL INFORMATION¹:

Can be used with compression therapy	YES
Waterresistant	NO
MRI safe	NO

INDICATIONS FOR USE¹:

CarboFlex[®] is Indicated for the management of malodorous wounds. CarboFlex[®] may be used on infected malodorous wounds under medical supervisions with appropriate antibiotic therapy if indicated, together with frequent monitoring of the wound.

PRECAUTIONS AND OBSERVATIONS¹:

- CarboFlex[®] dressing may be used under compression therapy.
- This product is supplied sterile. If the immediate product packaging is damaged, do not use.
- If required, the wound should be debrided and necrotic tissue removed.
- Store at room temperature; avoid refrigeration and exposure to high humidity.
- The CarboFlex[®] dressing is for single-patient use only and should not be re-used. Re-use may lead to increased risk of infection or cross contamination. Physical properties of the device may no longer be optimal for intended use.

Due to an increased level of manganese content in the dressing (<20ppm), the use of CarboFlex[®] wound dressing is not recommended for infants less than 12 months of age.

DURATION OF USE¹:





With noninfected malodorous wounds, CarboFlex[®] may be left undisturbed for up to three days. If the wound is infected, then CarboFlex[®] should be changed more frequently.



Do not re-use

CONTRAINDICATIONS¹:

Do not use on individuals with a known sensitivity to the dressing or its components.

STERILIZATION ¹ :	
Sterile	YES
Method	CarboFlex is sterilised using gamma irradiation and is not intended to be resterilised by the user
CERTIFICATIONS ¹ :	
CE mark	CE 2797
MDR 2017/745	Class IIb
MATERIAL SPECIFICATIONS ¹ :	
List of ingredients	 This dressing has an absorbent wound contact layer (containing alginate and Hydrofiber®), an odour absorbing activated charcoal pad, an absorbent padding layer, and a soft flocking outer contained within a peel pouch.
STORAGE AND HANDLING ¹ :	
Before Use	Keep away from sunlight
  	Keep dry.
	Store at room temperature (10°C - 25°C/50°F - 77°F).
After use	After use, this product may be a potential biohazard. Handle and dispose of in accordance with accepted medical practice and applicable local laws and regulations.

References:
1. CarboFlex IFU Revised 2020-02
1260676V1

Convatec Limited
First Avenue,
Deeside Industrial Park Deeside,
Flintshire CH5 2NU,
UK



UNOMEDICAL A/S
Aaholmvej 1-3, Osted
4320 LEJRE
DENMARK



Convatec Denmark A/S
CVR 31477093,
Transformervej 14, 1. , 2860 Soborg,
Denmark
Tlf.: +45 48 16 74 74
www.convatec.dk

Convatec Norway AS
Org. nummer: MVA 992 780 118
Visiting address:
Nils Hansens Vei 8, 0667 Oslo
Postadresse:
Postboks 6464 Etterstad, 0605 Oslo
Tlf.: +47 21 09 67 90
www.convatec.no

Convatec (Sweden) AB
Organisationsnummer: 556754-8473,
Box 3096, 16973 Solna, Sweden
Visiting address:
Gustav III:s Boulevard 42, plan 9,
16973 Solna, Sweden
Tlf.: +46 020-21 22 22
www.convatec.se

Convatec Oy
Y-tunnus 1644589-9,
Aviapolis, Karhumäentie 3, 01530 VANTAA,
Finland
P: +358 020 7659 600
www.convatec.fi

For full description and instructions for use please see IFU in product box
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