

Conveen Contour Leg bag, unsterile

27. May 2013
Version 1.0

Material Safety Data Sheet

Based on template version 3.0

Identification of the substance/mixture and of the company/undertaking

Product name: Conveen Contour Leg bag, unsterile
Product code: 05170, 05174, 05176
Product information: Continence

Manufacturer: Coloplast A/S
Holtedam 1
DK-3050 Humlebaek
Denmark
Telephone +45 49111111
msds@coloplast.com

USA

Coloplast Corp.
1601 West River Road N
Minneapolis, MN 55411
Telephone: +1-800-533-0464
www.us.coloplast.com

Canada

Coloplast Canada Corporation
3300 Ridgeway Drive, Unit 12
Mississauga, Ont. L5L 5Z9
Telephone: +1-877-820-7008
www.coloplast.ca

Europe

Coloplast A/S
Holtedam 1
DK-3050 Humlebaek
Telephone: +45 49 11 11 11
www.coloplast.com

Hazards identification

This product consists primarily of polymer materials. The products pose no immediate hazard.

Composition/information on ingredients

This product contains DEHP (di (2-ethylhexyl) phthalate), CAS 117-81-7, which is classified as hazardous under EU and US OSHA regulations. DEHP is identified as a Substance of Very High Concern (SVHC) by the EU (REACH). Due to its containment in a polymer matrix in the product, the substance poses no immediate hazard. Main ingredients and packaging materials are listed below.

Chemical name	CAS no.
PVC (polyvinylchloride)	93050-82-9
DEHP (di (2-ethylhexyl) phthalate)	117-87-1
Polypropylene	9003-07-0
Polyethylene	9002-88-4
Styrene-ethylene/butylene	66070-58-4
Acrylonitrile butadiene styrene	26657-42-1

Conveen Contour Leg bag, unsterile

Ethyl vinyl acetate	24937-78-8
Polyethylene terephthalate	25038-59-9
Silicone	-
Stabilizer	-
Softener	-
Pigments	-
Packaging:	
Polyamide (PA)/Polyethylene (PE) film	-9002-88-4
Corrugated cardboard	-
Coated paper	-

Disposal considerations

Dispose the device according to the recommended disposal technology at any approved facility. Under normal private use the product may be disposed of together with other household waste unless local regulations set special requirements. The disposal should always be in compliance with national, federal, state and local regulations. The product should not be discharged to the environment.

US

This product does not meet the criteria for hazardous waste as defined under the Resource Conservation and Recovery Act (RCRA) 40 CFR 261. Under normal private use the product may be disposed of together with other household waste per RCRA 40 RFT 261.4.B1.

European Union

Per The European Waste Catalogue (EWC), in accordance with EC Directive 75/422/ECC, the following Waste Code can be used: 18 01 04 00 wastes whose collection and disposal is not subject to special requirements in view of the prevention of infection (e.g. dressings, plaster casts, linen, disposable clothing, diapers). However, if the waste in view of the prevention of infection needs special requirements, other Waste Codes should be used. Under normal private use the product may be disposed of together with other household waste unless local regulations set special requirements.

Handling and storage

Handling: See instruction for use
Storage: Store until use as supplied and at room temperature unless other information is stated on the packaging or on the leaflet.

Other information

This MSDS is supplied as an additional service to the customer. The product is a medical device, which is regulated under the Council Directive 93/42/ECC, Medical Device Directive. The product has been evaluated according to the requirements of medical devices. According to current knowledge this product is considered non-toxic. For further information please contact Coloplast A/S.