

# Comfeel Plus

19. April 2016  
Version 1.0

## Material Safety Data Sheet

Based on template version 5.0

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

Product name: Comfeel Plus (C+)  
Sacral  
Product code: 33285  
Product information: Wound care product  
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](http://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](http://www.coloplast.ca)

### Europe

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

### Hazards identification

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

### Composition/information on ingredients

This product is regulated as a medical device in European Economic Area (EEA). In other regions it may be regulated as a medical device, a cosmetic or not regulated.

This product does not contain substances classified as hazardous under EC Regulation No. 1272/2008/EC, Annex VI (EU) in concentrations above 0.1 % (w/w). Main ingredients and packaging materials are listed below.

Chemical name	CAS no.
Polyurethane film	9009-54-5
Adhesive	-
PETP film with and without blue pigment	25038-59-9
Blue pigment mix	147-14-8 and 1328-53-6

## Comfeel Plus

Silicone coating on PETP film  
PET/PE packaging film:

9016-00-6  
25038-59-9 / 9002-88-4

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### 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.

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### 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.

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### 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.

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Approved by	DKLILYC Lise Lyck Senior Biosafety Specialist Issuer 19-Apr-2016 10:57:15 GMT+0000
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Document Owner: DKLILYC Lise Lyck