

## Purilon Gel



29. May 2019  
Version 2.0

### Safety Data Sheet

Based on template version 7.0

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

Product name:	Purilon Gel
Product code:	03900, 03903, 03906
Product information:	Wound care product
Manufacturer:	Coloplast A/S Holtedam 1 DK-3050 Humlebaek Denmark Telephone +45 49111111 msds@coloplast.com

#### USA

Coloplast Corp.  
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Minneapolis, MN 55411  
Telephone: +1-800-533-0464  
[www.us.coloplast.com](http://www.us.coloplast.com)

#### Canada

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Concord, ON, L4K 0J1  
Telephone: +1-888-880-8605  
[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)

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#### Hazards identification

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

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#### 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.

The safety of this medical device has been evaluated according to the requirements in the Medical Device Directive (Council Directive 93/42/EEC of 14 June 1993).

This product does not contain Substances of Very High Concern (SVHC) in concentrations above 0.1 %w/w according to the candidate list, article 59 (10) European REACH regulation (EC) No. 1907/2006.

## Purilon Gel

Main ingredients and packaging materials are listed below.

<b>Chemical name</b>	<b>CAS no.</b>
Hydrocolloid	-
Water	7732-18-5

<b>Packaging</b>	
PE (Polyethylene)	9002-88-4
Pigments	-

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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 SDS 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.