

**K833569 COMFEEL BARRIER CREAM**Dec 27, 1983  
76 days to decisionK833569 · Product code: **EXB** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k833569/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Ostomy (EXB)
Date received	Oct 12, 1983
Decision date	Dec 27, 1983
Days to decision	76 days
Third-party review	No

**APPLICANT**

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Company	<b>Coloplast A/S</b>
Location	Mchenry, IL, US
Website	<a href="http://www.coloplast.com/">http://www.coloplast.com/</a>
510(k) history	71 submissions · 68 cleared · 1983-2023

Coloplast A/S is a Danish multinational medical device manufacturer based in McHenry, US. The company develops and markets devices for ostomy, urology, continence, and wound care. Coloplast has received FDA 510(k) clearances from total submissions since its first clearance in 1983. The company's regulatory portfolio is dominated by Gastroenterology & Urology devices, including catheter systems, guidewires, and access sheaths. The latest clearance on record dates to 2023, reflecting the company's historical engagement with FDA regulatory pathways. Notable cleared devices i...

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k833569/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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