

K873669 COMFEEL ULCUS ULCER CARE SYSTEMNov 17, 1987
67 days to decisionK873669 · Product code: **FRO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k873669/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Dressing, Wound, Drug (FRO)
Date received	Sep 11, 1987
Decision date	Nov 17, 1987
Days to decision	67 days
Third-party review	No

APPLICANT

Company	Coloplast A/S
Location	Mchenry, IL, US
Contact	RICHARD A HAMER
Website	http://www.coloplast.com/
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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