FDA 510(k) Premarket Notification

K844684 COLOPLAST URO 2002 NIGHT DRAINAGE SYSTEM

CLEARED

Jan 8, 1985
39 days to decision

K844684 · Product code: **KNX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k844684/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Nov 30, 1984
Decision date	Jan 8, 1985
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**Coloplast A/S**
Location	Mchenry, IL, US
Contact	DAVE HEFFNER
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...