FDA 510(k) Premarket Notification

CLEARED

K113496 VIRTURE MALE SLING SYSTEM WITH ALEXIS WOUND RETRACTOR CONVENIENCE KIT

Feb 14, 2012
81 days to decision

K113496 · Product code: **OTM** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k113496/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, For Stress Urinary Incontinence, Male (OTM)
Date received	Nov 25, 2011
Decision date	Feb 14, 2012
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Coloplast A/S**
Location	Mchenry, IL, US
Contact	JANELL COLLEY
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...