K131072 ORCHESTRA HYDROPHILIC GUIDEWIRE STANDARD ANGLED, ORCHESTRA HYDROPHILIC GUIDEWIRE STRAIGHT STIFF, ORCHESTRA HYDROPHILIC

CLEARED

Jul 9, 2013
83 days to decision

K131072 · Product code: **OCY** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k131072/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Guidewire, Gastroenterology-urology (OCY)
Date received	Apr 17, 2013
Decision date	Jul 9, 2013
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Coloplast A/S**
Location	Mchenry, IL, US
Contact	BRIAN SCHMIDT
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...