FDA 510(k) Premarket Notification

K932143 FLEXI-TRAK ANCHORING DEVICE

CLEARED

Dec 1, 1993
215 days to decision

K932143 · Product code: **KNY** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k932143/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter, G-u (KNY)
Date received	Apr 30, 1993
Decision date	Dec 1, 1993
Days to decision	215 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Convatec, A Division of E.R. Squibb & Sons**
Location	Walker, MI, US
Contact	MARILYN DREYLING
510(k) history	81 submissions · 68 cleared · 1982-2002