

K932143 FLEXI-TRAK ANCHORING DEVICEDec 1, 1993
215 days to decisionK932143 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932143/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026