FDA 510(k) Premarket Notification

K875188 LUER LOCK CONNECTORS / RIDIG P.V.C.

CLEARED

Mar 2, 1988
76 days to decision

K875188 · Product code: **LFJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k875188/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Dec 17, 1987
Decision date	Mar 2, 1988
Days to decision	76 days
Third-party review	No

APPLICANT

Company	**Vas-Cath of Canada , Ltd.**
Location	Mchenry, IL, US
Contact	KATHERINE CREWE
510(k) history	19 submissions · 19 cleared · 1979-1988