FDA 510(k) Premarket Notification

K903945 EXEL A-V FISTULA SET

CLEARED

Nov 20, 1990
85 days to decision

K903945 · Product code: **FIE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k903945/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Aug 27, 1990
Decision date	Nov 20, 1990
Days to decision	85 days
Third-party review	No

APPLICANT

Company	**Exel Intl.**
Location	Culver City, CA, US
Contact	ESHAGH HAMID
510(k) history	18 submissions · 18 cleared · 1986-2000