FDA 510(k) Premarket Notification

K891753 DUAL LUMEN SUB-CLAVIAN CATHETER, DLSC 600 & 800

CLEARED

Jun 8, 1989
77 days to decision

K891753 · Product code: **LFJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k891753/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Mar 23, 1989
Decision date	Jun 8, 1989
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Akcess Medical Products, Inc.**
Location	New Brunswick, NJ, US
Contact	EDWARD J HOPKINS
510(k) history	36 submissions · 28 cleared · 1989-1994