FDA 510(k) Premarket Notification

K833848 DUAL LUMEN SUBCLAVIAN CANNULA

CLEARED

Dec 8, 1983
34 days to decision

K833848 · Product code: **LFJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k833848/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Nov 4, 1983
Decision date	Dec 8, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Shiley, Inc.**
Location	Mchenry, IL, US
510(k) history	174 submissions · 174 cleared · 1976-1993