FDA 510(k) Premarket Notification

K761031 ELECATH ARTERIAL EMBOLECTOMY CATHETER

CLEARED

Dec 13, 1976
28 days to decision

K761031 · Product code: **DXE** · Cardiovascular
Source: https://www.510kdatabase.net/k761031/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Nov 15, 1976
Decision date	Dec 13, 1976
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Electro-Catheter Corp.**
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995