FDA 510(k) Premarket Notification

K854511 ELECATH PERCUTANEOUS LEFT ATRIAL CANNULATION SET

CLEARED

Feb 10, 1986
90 days to decision

K854511 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k854511/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Nov 12, 1985
Decision date	Feb 10, 1986
Days to decision	90 days
Third-party review	No

APPLICANT

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