FDA 510(k) Premarket Notification

K880571 CATH-TECH PORT IMPLANTABLE VASCULAR ACCESS SYSTEM

CLEARED

Mar 4, 1988
23 days to decision

K880571 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k880571/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Feb 10, 1988
Decision date	Mar 4, 1988
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Catheter Technology Corp.**
Location	Walker, MI, US
Contact	JACK SPEER
510(k) history	13 submissions · 13 cleared · 1983-1988