FDA 510(k) Premarket Notification

K881690 PORT-A-CATH PORTAL INTRODUCER FORCEPS

CLEARED

May 20, 1988
31 days to decision

K881690 · Product code: **LKG** · General Hospital
Source: https://www.510kdatabase.net/k881690/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intraventricular (LKG)
Date received	Apr 19, 1988
Decision date	May 20, 1988
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**Pharmacia Deltec, Inc.**
Location	St. Paul, MN, US
Contact	EDWARD NUMAINVILLE
510(k) history	43 submissions · 35 cleared · 1987-1995