

K901722 CHEMOSITE(TM) IMPLANTABLE DRUG DELIVERY SYSTEM

Jul 3, 1990
78 days to decision

K901722 · Product code: **LJT** · General Hospital
Source: <https://www.510kdatabase.net/k901722/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Apr 16, 1990
Decision date	Jul 3, 1990
Days to decision	78 days
Third-party review	No

APPLICANT

Company	Device Labs, Inc.
Location	Medway, MA, US
Contact	ELTON M TUCKER
510(k) history	6 submissions · 5 cleared · 1988-1993

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k901722/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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