FDA 510(k) Premarket Notification

# K901722 CHEMOSITE(TM) IMPLANTABLE DRUG DELIVERY SYSTEM

**CLEARED**

Jul 3, 1990
78 days to decision

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

## SUBMISSION DETAILS

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|---|---|
| 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

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