

**K961583 A PORT DETACHED CATHETER SYSTEM CATALOG
#1002**

Jul 18, 1996
85 days to decision

K961583 · Product code: **LJT** · General Hospital
Source: <https://www.510kdatabase.net/k961583/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Apr 24, 1996
Decision date	Jul 18, 1996
Days to decision	85 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Arrow Intl., Inc.
Location	Mchenry, IL, US
Contact	BRADLEY J ENEGREN
510(k) history	110 submissions · 105 cleared · 1976-2010

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Device record: <https://www.510kdatabase.net/k961583/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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