

**K953300 V-CATH DUAL LUMEN E.S.P. PERIPHERAL INSERTED
CENTRAL CATHETER**Apr 30, 1996
291 days to decisionK953300 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k953300/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jul 14, 1995
Decision date	Apr 30, 1996
Days to decision	291 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Hdc Corp.
Location	Walker, MI, US
Contact	BRUCE FIELDS
510(k) history	30 submissions · 29 cleared · 1983-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k953300/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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