

**K881690 PORT-A-CATH PORTAL INTRODUCER FORCEPS**May 20, 1988  
31 days to decisionK881690 · Product code: **LKG** · General Hospital  
Source: <https://www.510kdatabase.net/k881690/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881690/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 2, 2026