

**K891611 MODELS 6526,6527,6528 PERCU. TRANS. ANGIO.  
CATH.**Jun 12, 1989  
84 days to decisionK891611 · Product code: **LJT** · General Hospital  
Source: <https://www.510kdatabase.net/k891611/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Mar 20, 1989
Decision date	Jun 12, 1989
Days to decision	84 days
Third-party review	No

**APPLICANT**

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Company	<b>Medtronic Versaflex, Inc.</b>
Location	San Diego, CA, US
Contact	TIMOTHY J JOHNSON
510(k) history	2 submissions · 2 cleared · 1989-1989

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k891611/> 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 May 14, 2026