

**K821776 SURGITEK PERCUTANEOUS NEPHROST. STENT**Aug 10, 1982  
54 days to decisionK821776 · Product code: **LJE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k821776/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Nephrostomy (LJE)
Date received	Jun 17, 1982
Decision date	Aug 10, 1982
Days to decision	54 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Engineering Corp.</b>
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-1993

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k821776/> 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 28, 2026