

**K900031 MODIFIED PEDIATRIC SILICONE FOLEY CATHETERS**Mar 6, 1990  
61 days to decisionK900031 · Product code: **KNY** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k900031/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter, G-u (KNY)
Date received	Jan 4, 1990
Decision date	Mar 6, 1990
Days to decision	61 days
Third-party review	No

**APPLICANT**

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Company	<b>Porges Corp.</b>
Location	Stamford, CT, US
Contact	HARRY M KAUFMAN
510(k) history	18 submissions · 18 cleared · 1988-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900031/>, 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