

**K895942 GESCO DUAL LUMEN PER-Q-CATH-DL**Jul 2, 1990  
264 days to decisionK895942 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k895942/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Oct 11, 1989
Decision date	Jul 2, 1990
Days to decision	264 days
Third-party review	No

**APPLICANT**

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Company	<b>Gesco Intl., Inc.</b>
Location	Walker, MI, US
Contact	GEORGE E SINKO
510(k) history	26 submissions · 22 cleared · 1978-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895942/>; 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 26, 2026