

**K831845 PER-Q-CATH TRAY**Aug 24, 1983  
77 days to decisionK831845 · Product code: **LJS** · General Hospital  
Source: <https://www.510kdatabase.net/k831845/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jun 8, 1983
Decision date	Aug 24, 1983
Days to decision	77 days
Third-party review	No

**APPLICANT**

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Company	<b>Gesco Intl., Inc.</b>
Location	Walker, MI, US
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/k831845/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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