

**K823565 MULTI-LUMEN INTRAVENOUS CATHETER**Dec 16, 1982  
13 days to decisionK823565 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k823565/>**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	Dec 3, 1982
Decision date	Dec 16, 1982
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Mcperson Enterprises, Inc.</b>
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823565/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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