

**K955839 MULTI-MED MULTI-LUMEN CENTRAL VENOUS  
CATHETER**Mar 25, 1996  
90 days to decisionK955839 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k955839/>**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 26, 1995
Decision date	Mar 25, 1996
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Baxter Edwards</b>
Location	Irvine, CA, US
Contact	IRENE PARKER
510(k) history	11 submissions · 10 cleared · 1993-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955839/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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