

**K894430 MODIFIED LABEL TO PERCU. INTRO. W/SIDEPORT & 3-WAY**Aug 9, 1989  
23 days to decisionK894430 · Product code: **DYB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k894430/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jul 17, 1989
Decision date	Aug 9, 1989
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Daig Corp.</b>
Location	Mchenry, IL, US
Contact	J FLEISCHHACKER
510(k) history	63 submissions · 63 cleared · 1977-2000

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894430/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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