

**K882655 PROCEDURE PACKS PRESS.  
MONITOR/ADMINISTRATION LINE**Sep 16, 1988  
79 days to decisionK882655 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k882655/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jun 29, 1988
Decision date	Sep 16, 1988
Days to decision	79 days
Third-party review	No

**APPLICANT**

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Company	<b>Procedure Packs, Inc.</b>
Location	Scottsdale, AZ, US
Contact	ROBERT S FUGATE
510(k) history	4 submissions · 4 cleared · 1988-1989

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882655/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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