

**K895932 PSG(TM) EDM(TM) INFUSION CATHETER**Jan 4, 1990  
86 days to decisionK895932 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k895932/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 10, 1989
Decision date	Jan 4, 1990
Days to decision	86 days
Third-party review	No

**APPLICANT**

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Company	<b>Peripheral Systems Group</b>
Location	Mountain View, CA, US
Contact	JESSICA AYRES
510(k) history	17 submissions · 17 cleared · 1990-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895932/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 20, 2026