

**K901151 I.V. ADMINISTRATION SET CONNECTING DEVICE**Jul 2, 1990  
112 days to decisionK901151 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k901151/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 12, 1990
Decision date	Jul 2, 1990
Days to decision	112 days
Third-party review	No

**APPLICANT**

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Company	<b>Dmd, Div. of Casta Pacific Co.</b>
Location	Nevada City, CA, US
Contact	JOHN T WRIGHT
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901151/>; 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 July 7, 2026