

**K852259 IVENT INTRAVENOUS ADMINISTRATION SET**Jun 7, 1985  
14 days to decisionK852259 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k852259/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 24, 1985
Decision date	Jun 7, 1985
Days to decision	14 days
Third-party review	No

**APPLICANT**

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Company	<b>Ivent Corp.</b>
Location	Walker, MI, US
Contact	CHRIS OZIMEK
510(k) history	3 submissions · 3 cleared · 1983-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k852259/>; 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 9, 2026