

**K850112 OMNIMED IV SOLUTION ADMINISTRATION SET**Feb 22, 1985  
39 days to decisionK850112 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k850112/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 14, 1985
Decision date	Feb 22, 1985
Days to decision	39 days
Third-party review	No

**APPLICANT**

---

Company	<b>Omni Intl., Inc.</b>
Location	Walker, MI, US
Contact	RAMAN KADEVARI
510(k) history	12 submissions · 12 cleared · 1982-1999

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k850112/>; 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 27, 2026