

**K896892 KAWASUMI Y-TYPE BLOOD ADMINISTRATION SET**Sep 12, 1990  
278 days to decisionK896892 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k896892/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 8, 1989
Decision date	Sep 12, 1990
Days to decision	278 days
Third-party review	No

**APPLICANT**

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Company	<b>Kawasumi Laboratories Co., Ltd.</b>
Location	Canoga Park, CA, US
Contact	KENJIRO TANI
510(k) history	18 submissions · 18 cleared · 1987-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896892/>; 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 June 30, 2026