

K892438 KENDALL CURITY INTRAVASCULAR CUT DOWN TUBESJul 24, 1989
105 days to decisionK892438 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k892438/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 10, 1989
Decision date	Jul 24, 1989
Days to decision	105 days
Third-party review	No

APPLICANT

Company	The Kendal Co.
Location	Mchenry, IL, US
Contact	WILLIAM D CURTIN
510(k) history	63 submissions · 60 cleared · 1980-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k892438/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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