

**K823214 DOSIMETRIC RELEASE PROGRAM - ADMIN. SET**Jan 12, 1983  
76 days to decisionK823214 · Product code: **FPA** · Chemistry  
Source: <https://www.510kdatabase.net/k823214/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Oct 28, 1982
Decision date	Jan 12, 1983
Days to decision	76 days
Third-party review	No

**APPLICANT**

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Company	<b>Cutter Laboratories, Inc.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.bayer.com">https://www.bayer.com</a>
510(k) history	39 submissions · 39 cleared · 1976-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823214/>; 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 22, 2026