

**K834106 INTRAVENOUS INFUSION CONTROLLER**May 30, 1984  
183 days to decisionK834106 · Product code: **LDR** · General Hospital  
Source: <https://www.510kdatabase.net/k834106/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Nov 29, 1983
Decision date	May 30, 1984
Days to decision	183 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/k834106/>; 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