

**K820453 THE INT-STOPPER**Mar 8, 1982  
18 days to decisionK820453 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k820453/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 18, 1982
Decision date	Mar 8, 1982
Days to decision	18 days
Third-party review	No

**APPLICANT**

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Company	<b>Burrton Medical Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1979-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k820453/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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