

**K833030 INTRAVASCULAR ADMINISTRATION SET**Nov 3, 1983  
57 days to decisionK833030 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k833030/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 7, 1983
Decision date	Nov 3, 1983
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Gish Biomedical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	75 submissions · 75 cleared · 1983-2009

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

Device record: <https://www.510kdatabase.net/k833030/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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