

**K811110 MODELS: 52-01-0, 52-01-1**Jun 11, 1981  
49 days to decisionK811110 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k811110/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 23, 1981
Decision date	Jun 11, 1981
Days to decision	49 days
Third-party review	No

**APPLICANT**

---

Company	<b>Y</b>
Location	Mchenry, IL, US
510(k) history	78 submissions · 78 cleared · 1978-1996

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k811110/> 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 July 3, 2026