

**K913311 SAFETY INJECTION PORT, MODIFICATION**Sep 13, 1991  
56 days to decisionK913311 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k913311/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 19, 1991
Decision date	Sep 13, 1991
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Riverside Technology, Inc.</b>
Location	Lisle, IL, US
Contact	W. C MILLER
510(k) history	1 submissions · 1 cleared · 1991-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913311/> 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 7, 2026