

**K821032 MODEL SR-1020LF**May 3, 1982  
20 days to decisionK821032 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k821032/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 13, 1982
Decision date	May 3, 1982
Days to decision	20 days
Third-party review	No

**APPLICANT**

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Company	<b>Stewart-Riess Labs</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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

Device record: <https://www.510kdatabase.net/k821032/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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