

**K992104 PARSET, MODEL A10001E**Aug 23, 1999  
62 days to decisionK992104 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k992104/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jun 22, 1999
Decision date	Aug 23, 1999
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Rd Medical Mfg., Inc.</b>
Location	Culebra, PR, US
Contact	CARLOS A RODRIGUEZ-GARCIA
510(k) history	4 submissions · 4 cleared · 1999-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992104/> 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 4, 2026