

**K010538 PARSET NEEDLELESS PRIMARY SET FOR BLUNT CANNULA, A10003E, PARSET NEEDLELESS SECONDARY SET WITH BLUNT CANNULA, A14003E**Mar 30, 2001  
35 days to decisionK010538 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k010538/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 23, 2001
Decision date	Mar 30, 2001
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010538/> 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