

K991653 INVISION-PLUS INJECTION PORT CONNECTOR SYSTEMJun 24, 1999
42 days to decisionK991653 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k991653/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 13, 1999
Decision date	Jun 24, 1999
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Rymed Technologies, Inc.
Location	Shelton, CT, US
Contact	AL WEISENBORN
510(k) history	3 submissions · 3 cleared · 1996-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k991653/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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