FDA 510(k) Premarket Notification

K991653 INVISION-PLUS INJECTION PORT CONNECTOR SYSTEM

CLEARED

Jun 24, 1999
42 days to decision

K991653 · 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