FDA 510(k) Premarket Notification

K020881 MODIFICATION TO PERSONAL INFUSOR LOCAL PAIN MANAGEMENT PROCEDURAL KIT

CLEARED

Mar 27, 2002
9 days to decision

K020881 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k020881/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Pump, Infusion (FRN)
Date received	Mar 18, 2002
Decision date	Mar 27, 2002
Days to decision	9 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Science Incorporated**
Location	Bloomington, MN, US
Contact	LYNN WEIST
510(k) history	4 submissions · 4 cleared · 1997-2002