

**K020881 MODIFICATION TO PERSONAL INFUSOR LOCAL PAIN
MANAGEMENT PROCEDURAL KIT**Mar 27, 2002
9 days to decisionK020881 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020881/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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