

K010824 PERSONAL INFUSOR LOCAL PAIN MANAGEMENT PROCEDURAL KIT

Jun 5, 2001
78 days to decision

K010824 · Product code: **MEB** · General Hospital
Source: <https://www.510kdatabase.net/k010824/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Elastomeric (MEB)
Date received	Mar 19, 2001
Decision date	Jun 5, 2001
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k010824/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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