

**K023098 ACCUFUSER, ACCUFUSER PLUS, STANDARD
PROCEDURE KIT, ACCUFUSER INFUSION KIT**Dec 9, 2002
82 days to decisionK023098 · Product code: **MEB** · General Hospital
Source: <https://www.510kdatabase.net/k023098/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Elastomeric (MEB)
Date received	Sep 18, 2002
Decision date	Dec 9, 2002
Days to decision	82 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mckinley Infuser
Location	San Francisco, CA, US
Contact	JOHN CHAPPELL
510(k) history	1 submissions · 1 cleared · 2002-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k023098/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 2, 2026