

K911769 RATEMINDER(TM) V INFUSION PUMP, MODIFICATIONJun 3, 1991
45 days to decisionK911769 · Product code: **FRN** · General Hospital
Source: <https://www.510kdatabase.net/k911769/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Apr 19, 1991
Decision date	Jun 3, 1991
Days to decision	45 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Critikon Company, LLC
Location	Mchenry, IL, US
Contact	JOHN MATSON
510(k) history	51 submissions · 51 cleared · 1979-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k911769/> 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 May 26, 2026