FDA 510(k) Premarket Notification

K091386 MICROFUSE DUAL RATE INFUSER

CLEARED

Oct 2, 2009
144 days to decision

K091386 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k091386/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Pump, Infusion (FRN)
Date received	May 11, 2009
Decision date	Oct 2, 2009
Days to decision	144 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Numia Medical Technology, LLC**
Location	Lyndonville, VT, US
Contact	ERIC FLACHBART
510(k) history	1 submissions · 1 cleared · 2009-2009