

K062592 CADD-SENTRY PRO MEDICATION SAFETY SOFTWARENov 29, 2006
89 days to decisionK062592 · Product code: **FRN** · General Hospital
Source: <https://www.510kdatabase.net/k062592/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Sep 1, 2006
Decision date	Nov 29, 2006
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Smiths Medical MD, Inc.
Location	St. Paul, MN, US
Contact	MELANIE HESS
510(k) history	20 submissions · 20 cleared · 2004-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062592/> 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 June 15, 2026