FDA 510(k) Premarket Notification

K062592 CADD-SENTRY PRO MEDICATION SAFETY SOFTWARE

CLEARED

Nov 29, 2006
89 days to decision

K062592 · 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