FDA 510(k) Premarket Notification

K030661 STRYKER PAINPUMP2

CLEARED

May 30, 2003
88 days to decision

K030661 · Product code: **MEA** · General Hospital
Source: https://www.510kdatabase.net/k030661/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion, Pca (MEA)
Date received	Mar 3, 2003
Decision date	May 30, 2003
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Instruments**
Location	Kalamazoo, MI, US
Contact	NICOLE PETTY
510(k) history	73 submissions · 73 cleared · 1994-2026