FDA 510(k) Premarket Notification

K920377 MORTARA INSTRUMENT MODEL ECG PREVU

CLEARED

Apr 28, 1992
90 days to decision

K920377 · Product code: **LOS** · Cardiovascular
Source: https://www.510kdatabase.net/k920377/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jan 29, 1992
Decision date	Apr 28, 1992
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mortara Instrument, Inc.**
Location	Walker, MI, US
Contact	JOHN PRIMOZICH
510(k) history	51 submissions · 51 cleared · 1983-2019