FDA 510(k) Premarket Notification

K832050 MORTARA INSTRUMENT #101

CLEARED

Dec 13, 1983
172 days to decision

K832050 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k832050/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Jun 24, 1983
Decision date	Dec 13, 1983
Days to decision	172 days
Third-party review	No

APPLICANT

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