FDA 510(k) Premarket Notification

K831274 WLECTROCARDIOGRAPH ECG-7204

CLEARED

Oct 27, 1983
191 days to decision

K831274 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k831274/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Apr 19, 1983
Decision date	Oct 27, 1983
Days to decision	191 days
Third-party review	No

APPLICANT

Company	**Nihon Kohden America, Inc.**
Location	Foothill Ranch, CA, US
510(k) history	166 submissions · 163 cleared · 1979-2012