FDA 510(k) Premarket Notification

K823249 LIFESCOPE II/FOUR

CLEARED

Nov 24, 1982
23 days to decision

K823249 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k823249/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Nov 1, 1982
Decision date	Nov 24, 1982
Days to decision	23 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