FDA 510(k) Premarket Notification

K102106 NIHON KHDEN PREFENSE EDNS-9000 SERIES CENTRAL NURSE STATION

CLEARED

Dec 17, 2010
143 days to decision

K102106 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k102106/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jul 27, 2010
Decision date	Dec 17, 2010
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nihon Kohden Corp.**
Location	Foothill, CA, US
Contact	STEVE GEERDES
510(k) history	9 submissions · 9 cleared · 2010-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k102106/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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