FDA 510(k) Premarket Notification

K142624 Neuromaster G1 MEE200

CLEARED

Apr 24, 2015
220 days to decision

K142624 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k142624/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Sep 16, 2014
Decision date	Apr 24, 2015
Days to decision	220 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nihon Kohden Corporation**
Location	Tokyo, JP
Contact	Natalie Kennel
510(k) history	18 submissions · 18 cleared · 2015-2025