FDA 510(k) Premarket Notification

K802887 MODEL EEG 4221

CLEARED

Dec 16, 1980
29 days to decision

K802887 · Product code: **GWQ** · Neurology
Source: https://www.510kdatabase.net/k802887/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Full-montage Standard Electroencephalograph (GWQ)
Date received	Nov 17, 1980
Decision date	Dec 16, 1980
Days to decision	29 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