FDA 510(k) Premarket Notification

K791234 MODEL EEG-4217 ELECTROENCEPHALOGRAPH

CLEARED

Sep 12, 1979
75 days to decision

K791234 · Product code: **GWQ** · Neurology
Source: https://www.510kdatabase.net/k791234/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Full-montage Standard Electroencephalograph (GWQ)
Date received	Jun 29, 1979
Decision date	Sep 12, 1979
Days to decision	75 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