FDA 510(k) Premarket Notification

K800319 ALPHA DETECTOR

CLEARED

Mar 3, 1980
20 days to decision

K800319 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k800319/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Feb 12, 1980
Decision date	Mar 3, 1980
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**M.O.E Electronics Research & Prod.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1980-1985