FDA 510(k) Premarket Notification

K800648 CADWELL 7200

CLEARED

Apr 21, 1980
28 days to decision

K800648 · Product code: **IKN** · Neurology
Source: https://www.510kdatabase.net/k800648/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electromyograph, Diagnostic (IKN)
Date received	Mar 24, 1980
Decision date	Apr 21, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Cadwell Laboratories, Inc.**
Location	Walker, MI, US
510(k) history	46 submissions · 46 cleared · 1979-2007