FDA 510(k) Premarket Notification

K962455 FOUR-CHANNEL PREAMPLIFIER

CLEARED

Jan 13, 1997
202 days to decision

K962455 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k962455/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Jun 25, 1996
Decision date	Jan 13, 1997
Days to decision	202 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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