FDA 510(k) Premarket Notification

K982324 VECTRA PRO MODELS 2 AND 4

CLEARED

Feb 1, 1999
214 days to decision

K982324 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k982324/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 2, 1998
Decision date	Feb 1, 1999
Days to decision	214 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Chattanooga Group, Inc.**
Location	Mchenry, IL, US
Contact	CHERYL G DYER
510(k) history	70 submissions · 68 cleared · 1980-2001