FDA 510(k) Premarket Notification

K014020 NT MEDITENS PLUS, TYPE 290

CLEARED

Jul 11, 2002
217 days to decision

K014020 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k014020/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 6, 2001
Decision date	Jul 11, 2002
Days to decision	217 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bio-Medical Research, Ltd.**
Location	Washington Dc, DC, US
Contact	MICHELLE SAWYER
510(k) history	32 submissions · 31 cleared · 1996-2021