FDA 510(k) Premarket Notification

K865048 ECLIPSE+, MODEL NUMBER 7723

CLEARED

Feb 26, 1987
64 days to decision

K865048 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k865048/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 24, 1986
Decision date	Feb 26, 1987
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**Medtronic Vascular**
Location	Walker, MI, US
Contact	DAVID H MUELLER
510(k) history	475 submissions · 453 cleared · 1977-2023