FDA 510(k) Premarket Notification

K955470 ACTION POTENTIAL SIMULATION THERAPY DEVICE

CLEARED

Oct 22, 1996
327 days to decision

K955470 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k955470/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Nov 30, 1995
Decision date	Oct 22, 1996
Days to decision	327 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Neurotech, Inc.**
Location	Washington, DC, US
Contact	J J M WILKINS
510(k) history	3 submissions · 3 cleared · 1986-1996