FDA 510(k) Premarket Notification

CLEARED

K020180 RELIEFBAND DEVICE, MODELS RB-DL, RB-EL, RB-RL

Mar 21, 2002
62 days to decision

K020180 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k020180/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jan 18, 2002
Decision date	Mar 21, 2002
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Woodside Biomedical, Inc.**
Location	Lake Forest, CA, US
Contact	CAROL PATTERSON
510(k) history	6 submissions · 6 cleared · 1998-2002