FDA 510(k) Premarket Notification

CLEARED

K083250 NEUROMED ELECTROANALGESIC DELIVERY SYSTEMS

Aug 18, 2009
287 days to decision

K083250 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k083250/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Nov 4, 2008
Decision date	Aug 18, 2009
Days to decision	287 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aa Neuromed Corp.**
Location	Port Coquitlam, Bc, CA
Contact	HERMANN DOLKER
510(k) history	1 submissions · 1 cleared · 2009-2009