K060222 LOW BACK PAIN RELIEF SYSTEM /MODELS GM310PP,GM320PP,GM321PP, AND GM330PP

CLEARED

Apr 28, 2006
88 days to decision

K060222 · Product code: **NUH** · Neurology
Source: https://www.510kdatabase.net/k060222/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH)
Date received	Jan 30, 2006
Decision date	Apr 28, 2006
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Gemore Technology Co, Ltd.**
Location	Tan Shui, Taipei Hsien, TW
Contact	BODEN S.P. LAI
510(k) history	14 submissions · 14 cleared · 2003-2019