

**K060222 LOW BACK PAIN RELIEF SYSTEM /MODELS
GM310PP,GM320PP,GM321PP, AND GM330PP**Apr 28, 2006
88 days to decisionK060222 · 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 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060222/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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