FDA 510(k) Premarket Notification

K071869 TENS/IF 14

CLEARED

Mar 7, 2008
245 days to decision

K071869 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k071869/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 6, 2007
Decision date	Mar 7, 2008
Days to decision	245 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Emsi**
Location	Alexandria, VA, US
Contact	CHERITA JAMES
510(k) history	6 submissions · 6 cleared · 2008-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k071869/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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