

K090922 EMPI ACTIVE TRANSCUTANEOUS NERVE STIMULATORJun 5, 2009
65 days to decisionK090922 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k090922/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Apr 1, 2009
Decision date	Jun 5, 2009
Days to decision	65 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Empi
Location	Walker, MI, US
Contact	SANDRA WALROD
Website	http://www.empik.com
510(k) history	60 submissions · 56 cleared · 1977-2010

Empi is a historical medical device manufacturer based in Walker, US. The company specialized in electrotherapy and neurostimulation devices. Empi received FDA 510(k) clearances from total submissions between 1977 and 2010. The company's primary focus was physical medicine and neurology devices, including transcutaneous nerve stimulators, iontophoresis systems, and electrotherapy equipment. This regulatory record spans over three decades of device development and market clearance. Empi is now a historical record only, with no FDA 510(k) clearances issued in more than 13 y...

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Device record: <https://www.510kdatabase.net/k090922/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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