

K792522 MODEL 980D TENS NEUROPACERDec 13, 1979
7 days to decisionK792522 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k792522/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 6, 1979
Decision date	Dec 13, 1979
Days to decision	7 days
Third-party review	No

APPLICANT

Company	Empi
Location	Walker, MI, US
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...
