

K896703 TBD IONTOPHORESIS ELECTRODEFeb 26, 1990
90 days to decisionK896703 · Product code: **EGJ** · Physical Medicine
Source: <https://www.510kdatabase.net/k896703/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SD
Submission type	Traditional
Device classification	Device, Iontophoresis, Other Uses (EGJ)
Date received	Nov 28, 1989
Decision date	Feb 26, 1990
Days to decision	90 days
Third-party review	No

APPLICANT

Company	Empi
Location	Walker, MI, US
Contact	STACY MATTSON
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...

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k896703/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 21, 2026