

K842812 MENTOR 150 PLUS TRANSCUTANEOUS ELEC.Aug 8, 1984
21 days to decisionK842812 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k842812/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 18, 1984
Decision date	Aug 8, 1984
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Mentor Corp.
Location	Mchenry, IL, US
510(k) history	61 submissions · 61 cleared · 1977-2013

Mentor Corp. is a surgical aesthetics and medical device company based in McHenry, US. Now part of Johnson & Johnson MedTech, the brand supplies products to plastic surgeons and specialists worldwide. Mentor has received FDA 510(k) clearances from total submissions since its first clearance in 1977. The company's regulatory record spans General & Plastic Surgery, Gastroenterology & Urology, Obstetrics & Gynecology, and Radiology device categories. The latest clearance was recorded in 2013, reflecting the company's historical significance in surgical device innovation. Men...