

K820168 MENTOR ACTIVATOR NEUROMUSCULAR STIMUFeb 24, 1982
34 days to decisionK820168 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k820168/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jan 21, 1982
Decision date	Feb 24, 1982
Days to decision	34 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...

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