

K803064 ZIMMER ELECTRICAL MUSCLE STIMULATORDec 31, 1980
27 days to decisionK803064 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k803064/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Dec 4, 1980
Decision date	Dec 31, 1980
Days to decision	27 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	Zimmer, Inc.
Location	Warsaw, IN, US
Website	https://www.zimmerbiomet.com
510(k) history	374 submissions · 353 cleared · 1976-2026

Zimmer, Inc. is a leading orthopedic medical device manufacturer based in Warsaw, US. The company specializes in innovative surgical implants and trauma solutions. Zimmer, Inc. maintains a strong FDA 510(k) regulatory record with cleared devices from total submissions since 1976. Orthopedic devices represent approximately 90% of the company's submission portfolio. The company remains actively engaged in product development, with the latest FDA 510(k) clearance in 2026. Recent cleared devices reflect the company's focus on joint reconstruction and trauma fixation. Notable ...
