

K813108 MODEL 500 VERI/500Feb 19, 1982
108 days to decisionK813108 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k813108/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Nov 3, 1981
Decision date	Feb 19, 1982
Days to decision	108 days
Third-party review	No

APPLICANT

Company	Verite
Location	Mchenry, IL, US
510(k) history	10 submissions · 9 cleared · 1979-1985

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Device record: <https://www.510kdatabase.net/k813108/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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