FDA 510(k) Premarket Notification

K813108 MODEL 500 VERI/500

CLEARED

Feb 19, 1982
108 days to decision

K813108 · Product code: **IPF** · Physical Medicine
Source: 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