FDA 510(k) Premarket Notification

K880018 LIDO(TM) WORKSET

CLEARED

Feb 26, 1988
52 days to decision

K880018 · Product code: **IKK** · Physical Medicine
Source: https://www.510kdatabase.net/k880018/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Jan 5, 1988
Decision date	Feb 26, 1988
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Loredan Biomedical, Inc.**
Location	Mchenry, IL, US
Contact	KAREN COMISSO
510(k) history	9 submissions · 9 cleared · 1983-1993