FDA 510(k) Premarket Notification

K903429 LIDO LINEA

CLEARED

Aug 14, 1990
14 days to decision

K903429 · Product code: **IKK** · Physical Medicine
Source: https://www.510kdatabase.net/k903429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Jul 31, 1990
Decision date	Aug 14, 1990
Days to decision	14 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