FDA 510(k) Premarket Notification

K880230 SIXALYZER

CLEARED

Mar 16, 1988
56 days to decision

K880230 · Product code: **IKK** · Physical Medicine
Source: https://www.510kdatabase.net/k880230/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Jan 20, 1988
Decision date	Mar 16, 1988
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Lukens Corp.**
Location	Mchenry, IL, US
Contact	LUTZ KAUFFMANN
510(k) history	11 submissions · 11 cleared · 1977-1988