FDA 510(k) Premarket Notification

K950646 INTEGRATOR

CLEARED

Apr 30, 1996
442 days to decision

K950646 · Product code: **LXM** · Physical Medicine
Source: https://www.510kdatabase.net/k950646/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Feb 13, 1995
Decision date	Apr 30, 1996
Days to decision	442 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Moyco Union Broach, Div., Moyco Technologies, Inc.**
Location	York, PA, US
Contact	CHRIS M LEHR
510(k) history	1 submissions · 1 cleared · 1996-1996