FDA 510(k) Premarket Notification

K904113 OPERATIVE RECORDING CAMERA

CLEARED

Oct 30, 1990
54 days to decision

K904113 · Product code: **IKO** · Physical Medicine
Source: https://www.510kdatabase.net/k904113/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hammer, Reflex, Powered (IKO)
Date received	Sep 6, 1990
Decision date	Oct 30, 1990
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Medical Dynamics, Inc.**
Location	Englewood, CO, US
Contact	JO BREHM
510(k) history	18 submissions · 17 cleared · 1988-1998