FDA 510(k) Premarket Notification

K920099 GYMMAN

CLEARED

Mar 9, 1992
60 days to decision

K920099 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k920099/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jan 9, 1992
Decision date	Mar 9, 1992
Days to decision	60 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Lumiscope Co., Inc.**
Location	Edison, NJ, US
Contact	STEPHEN L WHITTENBU
510(k) history	13 submissions · 13 cleared · 1988-2003