FDA 510(k) Premarket Notification

K896500 CWD/PPG MODULE

CLEARED

Jul 18, 1990
246 days to decision

K896500 · Product code: **JOM** · Cardiovascular
Source: https://www.510kdatabase.net/k896500/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Photoelectric, Pneumatic Or Hydraulic (JOM)
Date received	Nov 14, 1989
Decision date	Jul 18, 1990
Days to decision	246 days
Third-party review	No

APPLICANT

Company	**Life Sciences Manufacturing, Inc.**
Location	Mchenry, IL, US
Contact	ROBERT S CUTLER
510(k) history	12 submissions · 12 cleared · 1979-1991