FDA 510(k) Premarket Notification

K032871 PLANET MONITORING SYSTEM

CLEARED

Nov 10, 2003
56 days to decision

K032871 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k032871/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Sep 15, 2003
Decision date	Nov 10, 2003
Days to decision	56 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Larsen & Toubro Limited**
Location	Crofton, MD, US
Contact	E.J. Smith
510(k) history	18 submissions · 18 cleared · 2003-2011