FDA 510(k) Premarket Notification

K090172 STAR 55 MODEL 100

CLEARED

Jun 5, 2009
133 days to decision

K090172 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k090172/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jan 23, 2009
Decision date	Jun 5, 2009
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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