FDA 510(k) Premarket Notification

K060058 STELLAR 404T

CLEARED

Jan 20, 2006
11 days to decision

K060058 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k060058/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jan 9, 2006
Decision date	Jan 20, 2006
Days to decision	11 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

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