FDA 510(k) Premarket Notification

K053207 ARGUS, MODEL LCM

CLEARED

May 19, 2006
184 days to decision

K053207 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k053207/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Nov 16, 2005
Decision date	May 19, 2006
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Schiller AG**
Location	Baar, CH
Contact	MARKUS BUETLER
510(k) history	16 submissions · 16 cleared · 1985-2023