FDA 510(k) Premarket Notification

K032858 THE PHILIPS INTELLIVUE MP40, MP50, MP60, MP70,AND MP90 PATIENT MONITORS, RELEASE B.O.

CLEARED

Oct 10, 2003
28 days to decision

K032858 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k032858/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 12, 2003
Decision date	Oct 10, 2003
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Philips Medizin Systeme Boblingen GmbH**
Location	Andover, MA, US
Contact	HAUKE SCHIK
510(k) history	12 submissions · 12 cleared · 2001-2015