FDA 510(k) Premarket Notification

K843503 AEGIS AMBULATORY MONITORING SYSTEM

CLEARED

Oct 26, 1984
50 days to decision

K843503 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k843503/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 6, 1984
Decision date	Oct 26, 1984
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Aegis Medical Systems, Inc.**
Location	Marlton, NJ, US
Contact	JOHN R MORENO
510(k) history	1 submissions · 1 cleared · 1984-1984