FDA 510(k) Premarket Notification

K935732 CARDIO PERFECT ST 2001

CLEARED

Apr 4, 1996
855 days to decision

K935732 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k935732/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Dec 1, 1993
Decision date	Apr 4, 1996
Days to decision	855 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardio Control BV**
Location	The Netherlands, NL
Contact	DICK F VAN LUIJK
510(k) history	4 submissions · 4 cleared · 1992-1997