FDA 510(k) Premarket Notification

K990809 CADISCOPE ELECTRONIC STETHOSCOPE

CLEARED

Jun 9, 1999
90 days to decision

K990809 · Product code: **DQD** · Cardiovascular
Source: https://www.510kdatabase.net/k990809/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Mar 11, 1999
Decision date	Jun 9, 1999
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Caditec AG**
Location	Mundelein, IL, US
Contact	BETTY LOCK
510(k) history	1 submissions · 1 cleared · 1999-1999