FDA 510(k) Premarket Notification

K894648 ARBO TAB DISPOSABLE DIAGNOSTIC RESTING EKG ELECTRO

CLEARED

Sep 5, 1989
43 days to decision

K894648 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k894648/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jul 24, 1989
Decision date	Sep 5, 1989
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Arbo Medical, Inc.**
Location	Wilton, CT, US
Contact	WAYNE SHOCKLOSS
510(k) history	15 submissions · 15 cleared · 1989-1995