

K894648 ARBO TAB DISPOSABLE DIAGNOSTIC RESTING EKG ELECTROSep 5, 1989
43 days to decisionK894648 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k894648/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 2, 2026