

K013717 BRENTWOOD REAL TIME ST AND ARRHYTHMIA ANALYSIS SOFTWARE LIBRARY

Feb 6, 2002
90 days to decision

K013717 · Product code: **DPS** · Cardiovascular
Source: <https://www.510kdatabase.net/k013717/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Nov 8, 2001
Decision date	Feb 6, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Brentwood Medical Technology Corp.
Location	Torrance, CA, US
Contact	RUOMEI ZHANG
510(k) history	7 submissions · 7 cleared · 2000-2011

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k013717/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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