

K103640 MIDMARK IQECGMar 22, 2011
99 days to decisionK103640 · Product code: **DPS** · Cardiovascular
Source: <https://www.510kdatabase.net/k103640/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Dec 13, 2010
Decision date	Mar 22, 2011
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103640/> 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 May 25, 2026