

K831977 C TEC SERIES 9000 E.C.G. ELECTRODESep 9, 1983
81 days to decisionK831977 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k831977/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jun 20, 1983
Decision date	Sep 9, 1983
Days to decision	81 days
Third-party review	No

APPLICANT

Company	Speywood Medical, Ltd.
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k831977/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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