

K873442 ECG ELECTRODE SYSTEMNov 23, 1987
88 days to decisionK873442 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k873442/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Aug 27, 1987
Decision date	Nov 23, 1987
Days to decision	88 days
Third-party review	No

APPLICANT

Company	Hiltrode, Inc.(A/S) Subsidiary Munkenes A/S
Location	Cambridge, MA, US
Contact	RIKK I LARSEN
510(k) history	1 submissions · 1 cleared · 1987-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k873442/>; 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 7, 2026