FDA 510(k) Premarket Notification

K840646 CARDIOTRODE

CLEARED

Oct 26, 1984
254 days to decision

K840646 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k840646/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Feb 15, 1984
Decision date	Oct 26, 1984
Days to decision	254 days
Third-party review	No

APPLICANT

Company	**Kholber Corp.**
Location	Miami, FL, US
Contact	KEN M KOLBENSCHLAG
510(k) history	1 submissions · 1 cleared · 1984-1984

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k840646/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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