

K810895 ESOPHAGEAL PILL ELECTRODEMay 13, 1981
40 days to decisionK810895 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k810895/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Apr 3, 1981
Decision date	May 13, 1981
Days to decision	40 days
Third-party review	No

APPLICANT

Company	Con Med Equipment, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-1983

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Device record: <https://www.510kdatabase.net/k810895/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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