

K770521 ELECTROCARDIOGRAPHIC ELECTRODES, DISP.Mar 29, 1977
11 days to decisionK770521 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k770521/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Mar 18, 1977
Decision date	Mar 29, 1977
Days to decision	11 days
Third-party review	No

APPLICANT

Company	Dobre, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1977-1993

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Device record: <https://www.510kdatabase.net/k770521/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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