

**K791140 PREGELLED DISPOSABLE EELCTRO. SERIES 600**Jul 10, 1979  
22 days to decisionK791140 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k791140/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jun 18, 1979
Decision date	Jul 10, 1979
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Hayes Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1979

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k791140/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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