

**K900682 PHANTOM RADIOTRANSLUCENT ECG MONITORING  
ELECTRODE**Jul 31, 1990  
168 days to decisionK900682 · Product code: **DRY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k900682/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	Feb 13, 1990
Decision date	Jul 31, 1990
Days to decision	168 days
Third-party review	No

**APPLICANT**

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Company	<b>Contour Electrodes, Inc.</b>
Location	Herrin, IL, US
Contact	PARIS WALKER
510(k) history	4 submissions · 4 cleared · 1987-1990

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900682/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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