

**K893660 ECG SIMULATOR MODEL ECGPLUS**Dec 5, 1989  
204 days to decisionK893660 · Product code: **DPS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k893660/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	May 15, 1989
Decision date	Dec 5, 1989
Days to decision	204 days
Third-party review	No

**APPLICANT**

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Company	<b>Bio-Tek Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	DAN CONTOIS
510(k) history	37 submissions · 35 cleared · 1980-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893660/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 21, 2026