

**K904747 PEI PLAQUE ELECTRODE ARRAY**Oct 9, 1991  
355 days to decisionK904747 · Product code: **DRF** · CardiovascularSource: <https://www.510kdatabase.net/k904747/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Oct 19, 1990
Decision date	Oct 9, 1991
Days to decision	355 days
Third-party review	No

**APPLICANT**

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Company	<b>Prucka Engineering, Inc.</b>
Location	Bellaire, TX, US
Contact	STEVEN KIM
510(k) history	6 submissions · 6 cleared · 1989-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904747/>; 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 July 4, 2026