

**K830666 APNEA/BRADYCARDIA MONITOR HB 95**Jul 18, 1983  
136 days to decisionK830666 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k830666/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Mar 4, 1983
Decision date	Jul 18, 1983
Days to decision	136 days
Third-party review	No

**APPLICANT**

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Company	<b>Harvey-Bostwick, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983

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

Device record: <https://www.510kdatabase.net/k830666/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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