

**K792646 AMP DIRECT VASCULAR MONITORING SYSTEM**Feb 11, 1980  
52 days to decisionK792646 · Product code: **KRH** · CardiovascularSource: <https://www.510kdatabase.net/k792646/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ring, Annuloplasty (KRH)
Date received	Dec 21, 1979
Decision date	Feb 11, 1980
Days to decision	52 days
Third-party review	No

**APPLICANT**

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Company	<b>American Medical Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1978-1992

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

Device record: <https://www.510kdatabase.net/k792646/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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