

**K800122 HANCOCK AORTIC PUNCH**Feb 1, 1980  
14 days to decisionK800122 · Product code: **DWS** · CardiovascularSource: <https://www.510kdatabase.net/k800122/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instruments, Surgical, Cardiovascular (DWS)
Date received	Jan 18, 1980
Decision date	Feb 1, 1980
Days to decision	14 days
Third-party review	No

**APPLICANT**

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Company	<b>Hancock Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1978-1980

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

Device record: <https://www.510kdatabase.net/k800122/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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