

**K801885 HANCOCK \* OBTURATORS**Sep 9, 1980  
33 days to decisionK801885 · Product code: **DTI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k801885/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sizer, Heart-valve, Prosthesis (DTI)
Date received	Aug 7, 1980
Decision date	Sep 9, 1980
Days to decision	33 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/k801885/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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