

**K800015 AXIOM VESSEL LOOPS**Mar 25, 1980  
81 days to decisionK800015 · Product code: **KDC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k800015/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Jan 4, 1980
Decision date	Mar 25, 1980
Days to decision	81 days
Third-party review	No

**APPLICANT**

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Company	<b>Axiom Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-2000

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

Device record: <https://www.510kdatabase.net/k800015/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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