

**K800179 INFRASONDE AUTO/INFLATOR ELEC. VIBRATOR**Feb 11, 1980  
13 days to decisionK800179 · Product code: **DXW** · CardiovascularSource: <https://www.510kdatabase.net/k800179/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Phonocatheter, Intracavitary (DXW)
Date received	Jan 29, 1980
Decision date	Feb 11, 1980
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Sphygmetrics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1980

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

Device record: <https://www.510kdatabase.net/k800179/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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