

**K832882 DIF-STET**Dec 6, 1983  
102 days to decisionK832882 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k832882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Aug 26, 1983
Decision date	Dec 6, 1983
Days to decision	102 days
Third-party review	No

**APPLICANT**

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Company	<b>Intersect Systems, Inc.</b>
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1983-2002

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

Device record: <https://www.510kdatabase.net/k832882/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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