

**K810755 SQUINN VITATEK 500 SERIES PHYSIOL. MONI**Apr 3, 1981  
14 days to decisionK810755 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k810755/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 20, 1981
Decision date	Apr 3, 1981
Days to decision	14 days
Third-party review	No

**APPLICANT**

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Company	<b>Vitatek</b>
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1982

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

Device record: <https://www.510kdatabase.net/k810755/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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