

**K791380 VA 1000**Sep 19, 1979  
57 days to decisionK791380 · Product code: **DTA** · Cardiovascular  
Source: <https://www.510kdatabase.net/k791380/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Pacemaker Electrode Function (DTA)
Date received	Jul 24, 1979
Decision date	Sep 19, 1979
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Vitatron Medical BV</b>
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1976-1986

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Device record: <https://www.510kdatabase.net/k791380/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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