

**K790666 VITALITH S 3000 SERIES**Jun 1, 1979  
56 days to decisionK790666 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k790666/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)          |
| Submission type       | Traditional                                 |
| Device classification | Implantable Pacemaker Pulse-generator (DXY) |
| Date received         | Apr 6, 1979                                 |
| Decision date         | Jun 1, 1979                                 |
| Days to decision      | 56 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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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k790666/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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