

**K813549 CONTINUOUS FLUSH DEVICE**Jan 12, 1982  
22 days to decisionK813549 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k813549/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)              |
| Submission type       | Traditional                                     |
| Device classification | Transducer, Blood-pressure, Extravascular (DRS) |
| Date received         | Dec 21, 1981                                    |
| Decision date         | Jan 12, 1982                                    |
| Days to decision      | 22 days   |
| Third-party review    | No  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Namic</b>                          |
| Location       | Walker, MI, US                        |
| 510(k) history | 8 submissions · 8 cleared · 1979-1996 |

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

Device record: <https://www.510kdatabase.net/k813549/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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