

**K833271 NAMIC CONTINUOUS FLUSH DEVICE**Dec 1, 1983  
71 days to decisionK833271 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k833271/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 21, 1983
Decision date	Dec 1, 1983
Days to decision	71 days
Third-party review	No

**APPLICANT**

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Company	<b>North American Instrument Corp.</b>
Location	Walker, MI, US
510(k) history	28 submissions · 28 cleared · 1983-1996

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Device record: <https://www.510kdatabase.net/k833271/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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