

**K951722 PERCEPTOR COMPENSATOR MORSE MANIFOLD**Mar 25, 1996  
346 days to decisionK951722 · Product code: **DRS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k951722/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Apr 14, 1995
Decision date	Mar 25, 1996
Days to decision	346 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Namic</b>
Location	Walker, MI, US
Contact	MARY M RUBIN
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/k951722/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 28, 2026