

**K810297 DATASCOPE PERCOR=DL DUAL-LUMEN INTRA**May 6, 1981  
90 days to decisionK810297 · Product code: **DSP** · CardiovascularSource: <https://www.510kdatabase.net/k810297/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Balloon, Intra-aortic And Control (DSP)
Date received	Feb 5, 1981
Decision date	May 6, 1981
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Datascope Corp.</b>
Location	Mchenry, IL, US
510(k) history	136 submissions · 135 cleared · 1976-2019

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k810297/> 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 20, 2026