

**K173381 Nexxis OR**Feb 22, 2018  
115 days to decisionK173381 · Product code: **DXJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k173381/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Oct 30, 2017
Decision date	Feb 22, 2018
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Barco N.V.</b>
Location	La Jolla, CA, US
Contact	Eric Caus
Website	<a href="http://www.barco.com">http://www.barco.com</a>
510(k) history	60 submissions · 60 cleared · 1998-2024

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173381/> 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 May 31, 2026