

**K770330 MONITOR, PULSE WAVE VELOCITY P607**Feb 24, 1977  
7 days to decisionK770330 · Product code: **DXG** · CardiovascularSource: <https://www.510kdatabase.net/k770330/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Feb 17, 1977
Decision date	Feb 24, 1977
Days to decision	7 days
Third-party review	No

**APPLICANT**

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Company	<b>Cyborg Corp.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k770330/> 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 July 10, 2026