

**K792430 CARDIASSIST EXTERNAL COUNTERPULSATION**Jan 23, 1980  
57 days to decisionK792430 · Product code: **DRN** · CardiovascularSource: <https://www.510kdatabase.net/k792430/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Counter-pulsating, External (DRN)
Date received	Nov 27, 1979
Decision date	Jan 23, 1980
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardiassist Corp.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k792430/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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