

**K823191 DISPOSABLE PLASTIC STOPCOCK**Dec 30, 1982  
65 days to decisionK823191 · Product code: **DXT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k823191/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Oct 26, 1982
Decision date	Dec 30, 1982
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Vertex Medical Corp.</b>
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1982-1984

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Device record: <https://www.510kdatabase.net/k823191/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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