

K823253 VERTEX MONITORING CATHETERDec 30, 1982
59 days to decisionK823253 · Product code: **DXT** · CardiovascularSource: <https://www.510kdatabase.net/k823253/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Nov 1, 1982
Decision date	Dec 30, 1982
Days to decision	59 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k823253/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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