

K823340 CLEAR VINYL MONITORING CONNECTORDec 30, 1982
52 days to decisionK823340 · Product code: **DXT** · Cardiovascular
Source: <https://www.510kdatabase.net/k823340/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Nov 8, 1982
Decision date	Dec 30, 1982
Days to decision	52 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/k823340/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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