

K822596 VERTEX PERCUTANEOUS PUNCTURE NEEDLESep 21, 1982
25 days to decisionK822596 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k822596/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Aug 27, 1982
Decision date	Sep 21, 1982
Days to decision	25 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/k822596/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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