

K823256 ULTRA-ASEPT ANGIOGRAPHIC SYRINGE/SPAREDec 30, 1982
59 days to decisionK823256 · Product code: **DXT** · Cardiovascular
Source: <https://www.510kdatabase.net/k823256/>**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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k823256/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 3, 2026