

K823920 COEURLOCK STERILE DISPOS. ANGIOGRAPHICFeb 18, 1983
52 days to decisionK823920 · Product code: **DXT** · CardiovascularSource: <https://www.510kdatabase.net/k823920/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Dec 28, 1982
Decision date	Feb 18, 1983
Days to decision	52 days
Third-party review	No

APPLICANT

Company	Coeur Laboratories, Inc.
Location	NC, US
510(k) history	10 submissions · 10 cleared · 1983-1997

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Device record: <https://www.510kdatabase.net/k823920/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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