FDA 510(k) Premarket Notification

K921401 BIOPSY FORCEPS

CLEARED

Oct 21, 1992
211 days to decision

K921401 · Product code: **DWZ** · Cardiovascular
Source: https://www.510kdatabase.net/k921401/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biopsy, Endomyocardial (DWZ)
Date received	Mar 24, 1992
Decision date	Oct 21, 1992
Days to decision	211 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Annex Medical, Inc.**
Location	Eden Prairie, MN, US
Contact	STUART J LIND
510(k) history	23 submissions · 23 cleared · 1989-1995