FDA 510(k) Premarket Notification

K890453 REUSABLE MLYOCARDIAL BIOPSY FORCEPS (BIOPTOMES

CLEARED

Mar 31, 1989
60 days to decision

K890453 · Product code: **DWZ** · Cardiovascular
Source: https://www.510kdatabase.net/k890453/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biopsy, Endomyocardial (DWZ)
Date received	Jan 30, 1989
Decision date	Mar 31, 1989
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Dlp, Inc.**
Location	Mchenry, IL, US
Contact	VAN HOF
510(k) history	56 submissions · 56 cleared · 1979-1997