

**K890453 REUSABLE MLYOCARDIAL BIOPSY FORCEPS
(BIOPTOMES)**Mar 31, 1989
60 days to decisionK890453 · 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	Dip, Inc.
Location	Mchenry, IL, US
Contact	VAN HOF
510(k) history	56 submissions · 56 cleared · 1979-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890453/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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