FDA 510(k) Premarket Notification

K882383 MAGGI SERIES ULTRA. NEEDLE, BIOPSY/CATHETER GUIDES

CLEARED

Nov 28, 1988
172 days to decision

K882383 · Product code: **DWZ** · Cardiovascular
Source: https://www.510kdatabase.net/k882383/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biopsy, Endomyocardial (DWZ)
Date received	Jun 9, 1988
Decision date	Nov 28, 1988
Days to decision	172 days
Third-party review	No

APPLICANT

Company	**CIVCO Medical Instruments Co., Inc.**
Location	Walker, MI, US
Contact	VICTOR J WEDEL
510(k) history	29 submissions · 29 cleared · 1982-2023

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k882383/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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