FDA 510(k) Premarket Notification

K881412 MYOCARD BIOPSY FORCEP

CLEARED

Sep 8, 1988
157 days to decision

K881412 · Product code: **DWZ** · Cardiovascular
Source: https://www.510kdatabase.net/k881412/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biopsy, Endomyocardial (DWZ)
Date received	Apr 4, 1988
Decision date	Sep 8, 1988
Days to decision	157 days
Third-party review	No

APPLICANT

Company	**Fehling Medizintechnik GmbH**
Location	West Germany, DE
Contact	GUIDO FEHLING
510(k) history	1 submissions · 1 cleared · 1988-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k881412/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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