

K170726 Biopsy ForcepsJun 2, 2017
85 days to decisionK170726 · Product code: **DWZ** · Cardiovascular
Source: <https://www.510kdatabase.net/k170726/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biopsy, Endomyocardial (DWZ)
Date received	Mar 9, 2017
Decision date	Jun 2, 2017
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Fehling Instruments GmbH & Co. KG
Location	Karlstein, DE
Contact	Ulrike Lindner
510(k) history	2 submissions · 2 cleared · 2016-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k170726/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 1, 2026