

K223199 2.1 x 255mm Drill, 22mm StopNov 15, 2022
33 days to decisionK223199 · Product code: **HBE** · Neurology
Source: <https://www.510kdatabase.net/k223199/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Drills, Burrs, Trephines & Accessories (simple, Powered) (HBE)
Date received	Oct 13, 2022
Decision date	Nov 15, 2022
Days to decision	33 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Biomet Microfixation
Location	Jacksonville, FL, US
Contact	Jay Sharma
510(k) history	24 submissions · 24 cleared · 2012-2024

REGULATORY CONSULTANT

Consulting firm	MRC Global, LLC
Contact	Danielle Besal

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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