

K242322 Bone Marrow Biopsy NeedleJan 24, 2025
171 days to decisionK242322 · Product code: **KNW** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k242322/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Aug 6, 2024
Decision date	Jan 24, 2025
Days to decision	171 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Zhejiang Curaway Medical Technology Co., Ltd.
Location	Hangzhou, CN
Contact	Caili Cao
510(k) history	6 submissions · 6 cleared · 2023-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242322/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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