

K250216 OsteoProbe (OP-100)Sep 11, 2025
230 days to decisionK250216 · Product code: **QGQ** · Orthopedic
Source: <https://www.510kdatabase.net/k250216/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bone Indentation Device (QGQ)
Date received	Jan 24, 2025
Decision date	Sep 11, 2025
Days to decision	230 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Active Life Scientific, Inc.
Location	Santa Barbara, CA, US
Contact	Alexander Proctor
510(k) history	3 submissions · 2 cleared · 2021-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k250216/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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