FDA 510(k) Premarket Notification

K250216 OsteoProbe (OP-100)

CLEARED

Sep 11, 2025
230 days to decision

K250216 · 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