

**K250394 Fine TTO™**Oct 21, 2025  
251 days to decisionK250394 · Product code: **PBF** · Orthopedic  
Source: <https://www.510kdatabase.net/k250394/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopaedic Surgical Planning And Instrument Guides (PBF)
Date received	Feb 12, 2025
Decision date	Oct 21, 2025
Days to decision	251 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bodycad Laboratories, Inc.</b>
Location	Quebec City, CA
Contact	Nadine Adia
510(k) history	16 submissions · 16 cleared · 2017-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k250394/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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