

**K220823 Preat Abutments**Oct 26, 2022  
219 days to decisionK220823 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k220823/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Mar 21, 2022
Decision date	Oct 26, 2022
Days to decision	219 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Preat Corporation</b>
Location	Grover Beach, CA, US
Contact	Chris Bormes
510(k) history	3 submissions · 3 cleared · 2019-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Aclivi, LLC</b>
Contact	Chris Brown

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220823/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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