FDA 510(k) Premarket Notification

# K183518 Preat Abutments

**CLEARED**

Mar 18, 2019
90 days to decision

K183518 · Product code: **NHA** · Dental
Source: https://www.510kdatabase.net/k183518/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Abutment, Implant, Dental, Endosseous (NHA) |
| Date received | Dec 18, 2018 |
| Decision date | Mar 18, 2019 |
| Days to decision | 90 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

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

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k183518/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026