

**K171622 Dentium Ti-Base**Jan 11, 2018  
223 days to decisionK171622 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k171622/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Jun 2, 2017
Decision date	Jan 11, 2018
Days to decision	223 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dentium Co., Ltd.</b>
Location	Dunwoody, GA, US
Contact	Sangpil Yoon
510(k) history	22 submissions · 22 cleared · 2004-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k171622/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 27, 2026