

**K232151 Dynamic TiBase**Oct 17, 2023  
90 days to decisionK232151 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k232151/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Jul 19, 2023
Decision date	Oct 17, 2023
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Talladium Espa?a, SL</b>
Location	Lleida, ES
Contact	Xavier Soca Filella
510(k) history	7 submissions · 7 cleared · 2021-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k232151/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 24, 2026