

**K191919 Elos Accurate Hybrid Base**Feb 10, 2020  
207 days to decisionK191919 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k191919/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Jul 18, 2019
Decision date	Feb 10, 2020
Days to decision	207 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Elos Medtech Pinol A/S</b>
Location	Goerløse, DK
Contact	Tina Friis Poulsen
510(k) history	12 submissions · 12 cleared · 2018-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k191919/> 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 26, 2026