

**K120053 IPS E.MAX PRESS - ABUTMENT SOLUTIONS**Oct 18, 2012  
286 days to decisionK120053 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k120053/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Jan 6, 2012
Decision date	Oct 18, 2012
Days to decision	286 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ivoclar Vivadent, AG</b>
Location	Amherst, NY, US
Contact	DONNA MARIE HARTNETT
510(k) history	31 submissions · 31 cleared · 2004-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120053/> 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 June 15, 2026