

**K905436 NOBELPHARMA DRILLING EQUIPMENT DEC 500**Mar 4, 1991  
90 days to decisionK905436 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k905436/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 4, 1990
Decision date	Mar 4, 1991
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Nobelpharma USA, Inc.</b>
Location	Chicago, IL, US
Contact	THOMAS MULDOON
510(k) history	64 submissions · 64 cleared · 1990-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905436/>; 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 July 3, 2026