

**K071851 HDC STERILE SPIDER SCREW**Oct 18, 2007  
105 days to decisionK071851 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k071851/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jul 5, 2007
Decision date	Oct 18, 2007
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hdc S.R.L.</b>
Location	Bethesda, MD, US
Contact	GUIDO BONAPACE
510(k) history	3 submissions · 3 cleared · 2005-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071851/> 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 23, 2026