

**K090237 HG II FIXTURE SYSTEM**Apr 17, 2009  
74 days to decisionK090237 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k090237/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Feb 2, 2009
Decision date	Apr 17, 2009
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hiossen, Inc.</b>
Location	Fariless Hills, PA, US
Contact	MINJOO KIM
510(k) history	26 submissions · 26 cleared · 2009-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090237/> 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