FDA 510(k) Premarket Notification

K090237 HG II FIXTURE SYSTEM

CLEARED

Apr 17, 2009
74 days to decision

K090237 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k090237/

SUBMISSION DETAILS

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

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