

K203360 EK Implants and Abutments SystemJan 7, 2022
417 days to decisionK203360 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k203360/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Nov 16, 2020
Decision date	Jan 7, 2022
Days to decision	417 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Hiossen, Inc.
Location	Fariless Hills, PA, US
Contact	Peter Lee
510(k) history	25 submissions · 25 cleared · 2009-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k203360/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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