FDA 510(k) Premarket Notification

K170421 ET Hybrid Abutment

CLEARED

Jan 11, 2018
332 days to decision

K170421 · Product code: **NHA** · Dental
Source: https://www.510kdatabase.net/k170421/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Feb 13, 2017
Decision date	Jan 11, 2018
Days to decision	332 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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