

K190552 JJ Implant SystemDec 6, 2019
276 days to decisionK190552 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k190552/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Mar 5, 2019
Decision date	Dec 6, 2019
Days to decision	276 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Jj Implants
Location	Chalakudy, IN
Contact	Jiju John
510(k) history	1 submissions · 1 cleared · 2019-2019

REGULATORY CONSULTANT

Consulting firm	PaxMed International, LLC
Contact	Kevin A. Thomas

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k190552/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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