FDA 510(k) Premarket Notification

K243094 Quickdent Dental Implant System

CLEARED

Dec 19, 2024
80 days to decision

K243094 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k243094/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Sep 30, 2024
Decision date	Dec 19, 2024
Days to decision	80 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Quickdent Devices Private , Ltd.**
Location	West, Mumbai, Mumbai Surburban, IN
Contact	Mayur Khairnar
510(k) history	1 submissions · 1 cleared · 2024-2024

REGULATORY CONSULTANT

Consulting firm	**Blackwell Device Consulting**
Contact	Angela Blackwell

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