

K242432 MT-BoneAug 19, 2024
3 days to decisionK242432 · Product code: **DZI** · Dental
Source: <https://www.510kdatabase.net/k242432/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Bone, Powered (DZI)
Date received	Aug 16, 2024
Decision date	Aug 19, 2024
Days to decision	3 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Mectron S.P.A.
Location	Carasco, IT
Contact	Chiara Acunzo
510(k) history	13 submissions · 13 cleared · 2009-2025

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

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