

K161987 UF(II) Narrow Implant System - Fixture, UF(II) Narrow Implant System - SuprastructureFeb 21, 2017
217 days to decisionK161987 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k161987/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jul 19, 2016
Decision date	Feb 21, 2017
Days to decision	217 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Dio Corporation
Location	Los Angeles, CA, US
Contact	SANGIN PARK
510(k) history	14 submissions · 14 cleared · 2010-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k161987/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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