FDA 510(k) Premarket Notification

K100100 DIO STEADY EXTERNAL IMPLANT SYSTEM

CLEARED

Nov 10, 2010
302 days to decision

K100100 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k100100/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 12, 2010
Decision date	Nov 10, 2010
Days to decision	302 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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