FDA 510(k) Premarket Notification

K130222 STRAUMANN DENTAL IMPLANT SYSTEM

CLEARED

Apr 29, 2013
90 days to decision

K130222 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k130222/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 29, 2013
Decision date	Apr 29, 2013
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Straumann USA, LLC**
Location	Andover, MA, US
Contact	JENNIFER M JACKSON, MS
510(k) history	24 submissions · 24 cleared · 2012-2024

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k130222/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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