FDA 510(k) Premarket Notification

K132844 STRAUMANN CARES BONE LEVEL SCREW RETAINED BARS, STRAUMANN CARES BONE LEVEL SCREW RETAINED BRIDGES

CLEARED

Jan 15, 2014
126 days to decision

K132844 · Product code: **NHA** · Dental
Source: https://www.510kdatabase.net/k132844/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Sep 11, 2013
Decision date	Jan 15, 2014
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Straumann USA, LLC**
Location	Andover, MA, US
Contact	CHRISTOPHER KLACZYK
510(k) history	24 submissions · 24 cleared · 2012-2024

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k132844/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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