FDA 510(k) Premarket Notification

K101956 STRAUMANN MEMBRAGEL

CLEARED

Dec 16, 2010
157 days to decision

K101956 · Product code: **NPK** · Dental
Source: https://www.510kdatabase.net/k101956/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Barrier, Synthetic, Intraoral (NPK)
Date received	Jul 12, 2010
Decision date	Dec 16, 2010
Days to decision	157 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Straumann USA (On Behalf of Institut Straumann Ag)**
Location	Andover, MA, US
Contact	JANET C KAY
510(k) history	2 submissions · 2 cleared · 2007-2010

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k101956/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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