FDA 510(k) Premarket Notification

K011698 STRAUMANN GBR SYSTEM

CLEARED

Aug 23, 2001
83 days to decision

K011698 · Product code: **JEY** · Dental
Source: https://www.510kdatabase.net/k011698/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Jun 1, 2001
Decision date	Aug 23, 2001
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Institut Straumann AG**
Location	Basel, CH
Contact	LINDA JALBERT
Website	https://www.straumann.com
510(k) history	90 submissions · 90 cleared · 1990-2026

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k011698/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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