FDA 510(k) Premarket Notification

K980512 LORENZ RECONSTRUCTION SYSTEM WITH MODULAR SCREW

CLEARED

May 11, 1998
90 days to decision

K980512 · Product code: **JEY** · Dental
Source: https://www.510kdatabase.net/k980512/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Feb 10, 1998
Decision date	May 11, 1998
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Walter Lorenz Surgical, Inc.**
Location	Rockville, MD, US
Contact	DIANA PRESTON
510(k) history	56 submissions · 56 cleared · 1986-2007