FDA 510(k) Premarket Notification

K960639 ABUTMENT SELECTION KIT

CLEARED

May 14, 1996
90 days to decision

K960639 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k960639/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Feb 14, 1996
Decision date	May 14, 1996
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nobelpharma USA, Inc.**
Location	Chicago, IL, US
Contact	MARY EDWARDS
510(k) history	64 submissions · 64 cleared · 1990-1999