FDA 510(k) Premarket Notification

K905436 NOBELPHARMA DRILLING EQUIPMENT DEC 500

CLEARED

Mar 4, 1991
90 days to decision

K905436 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k905436/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 4, 1990
Decision date	Mar 4, 1991
Days to decision	90 days
Third-party review	No

APPLICANT

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