FDA 510(k) Premarket Notification

K923462 DELIGHT

CLEARED

Dec 21, 1992
160 days to decision

K923462 · Product code: **EAZ** · Dental
Source: https://www.510kdatabase.net/k923462/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Operating, Dental (EAZ)
Date received	Jul 14, 1992
Decision date	Dec 21, 1992
Days to decision	160 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Planmeca USA, Inc.**
Location	Mchenry, IL, US
Contact	ISMO SEPPA
510(k) history	13 submissions · 13 cleared · 1984-2011