FDA 510(k) Premarket Notification

K904948 PM2002 CC AND PM2002 EC

CLEARED

Jan 30, 1991
90 days to decision

K904948 · Product code: **EIA** · Dental
Source: https://www.510kdatabase.net/k904948/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Operative Dental (EIA)
Date received	Nov 1, 1990
Decision date	Jan 30, 1991
Days to decision	90 days
Third-party review	No

APPLICANT

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