FDA 510(k) Premarket Notification

K842357 PM 2000 DENTAL UNIT

CLEARED

Sep 12, 1984
82 days to decision

K842357 · Product code: **EIA** · Dental
Source: https://www.510kdatabase.net/k842357/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Operative Dental (EIA)
Date received	Jun 22, 1984
Decision date	Sep 12, 1984
Days to decision	82 days
Third-party review	No

APPLICANT

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