FDA 510(k) Premarket Notification

K012308 KAVO QUATTROCARE

CLEARED

Jun 26, 2002
338 days to decision

K012308 · Product code: **EFB** · Dental
Source: https://www.510kdatabase.net/k012308/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Air-powered, Dental (EFB)
Date received	Jul 23, 2001
Decision date	Jun 26, 2002
Days to decision	338 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Kavo America**
Location	Washington, DC, US
Contact	JOHN WESTERMEIER
510(k) history	20 submissions · 19 cleared · 1991-2008