FDA 510(k) Premarket Notification

K020062 XERAFIT

CLEARED

Feb 14, 2002
37 days to decision

K020062 · Product code: **EJH** · Dental
Source: https://www.510kdatabase.net/k020062/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alloy, Metal, Base (EJH)
Date received	Jan 8, 2002
Decision date	Feb 14, 2002
Days to decision	37 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Metalor Technologies USA**
Location	North Attleborough, MA, US
Contact	BRUCE A BARTON
510(k) history	4 submissions · 4 cleared · 2001-2002