

**K020062 XERAFIT**Feb 14, 2002  
37 days to decisionK020062 · Product code: **EJH** · DentalSource: <https://www.510kdatabase.net/k020062/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020062/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 2, 2026