

**K904948 PM2002 CC AND PM2002 EC**Jan 30, 1991  
90 days to decisionK904948 · Product code: **EIA** · DentalSource: <https://www.510kdatabase.net/k904948/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904948/>; 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 May 17, 2026