

**K842357 PM 2000 DENTAL UNIT**Sep 12, 1984  
82 days to decisionK842357 · Product code: **EIA** · DentalSource: <https://www.510kdatabase.net/k842357/>**SUBMISSION DETAILS**

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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**

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Company	<b>Planmeca USA, Inc.</b>
Location	Mchenry, IL, US
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/k842357/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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