

K923461 PM2002 PROLINE, PM2002 PROLINE ECDec 21, 1992
160 days to decisionK923461 · Product code: **KLC** · Dental
Source: <https://www.510kdatabase.net/k923461/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chair, Dental, With Operative Unit (KLC)
Date received	Jul 14, 1992
Decision date	Dec 21, 1992
Days to decision	160 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Planmeca USA, Inc.
Location	Mchenry, IL, US
Contact	ISMO SEPPA
510(k) history	13 submissions · 13 cleared · 1984-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923461/> Data sourced from FDA 510(k) public records (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. - Generated May 17, 2026