

**K923462 DELIGHT**Dec 21, 1992  
160 days to decisionK923462 · Product code: **EAZ** · DentalSource: <https://www.510kdatabase.net/k923462/>**SUBMISSION DETAILS**

---

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

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923462/> 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