

**K013802 FRC POSTEC**May 1, 2002  
167 days to decisionK013802 · Product code: **ELR** · DentalSource: <https://www.510kdatabase.net/k013802/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Post, Root Canal (ELR)
Date received	Nov 15, 2001
Decision date	May 1, 2002
Days to decision	167 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ivoclar Vivadent, Inc.</b>
Location	Amherst, NY, US
Contact	DONNA M HARTNETT
Website	<a href="https://www.ivoclar.com">https://www.ivoclar.com</a>
510(k) history	65 submissions · 65 cleared · 2001-2026

Ivoclar Vivadent, Inc. is a dental solutions provider based in Amherst, US. The company develops modern products for dental practitioners and laboratory technicians. The company has received FDA 510(k) clearances from total submissions since 2001. Dental devices represent 97% of its regulatory portfolio. The latest clearance was issued in 2026, confirming active market engagement. Recent cleared devices include restorative materials, adhesives, CAD-on restorations, bulk-fill composites, and curing lights. The product portfolio spans chairside and laboratory workflows for ...

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