

**K072213 PERIMETRICS PERIOMETER**Aug 29, 2008  
386 days to decisionK072213 · Product code: **EKX** · Dental  
Source: <https://www.510kdatabase.net/k072213/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Direct Drive, Ac-powered (EKX)
Date received	Aug 9, 2007
Decision date	Aug 29, 2008
Days to decision	386 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Perimetrics, Inc.</b>
Location	Mission Viejo, CA, US
Contact	ALBERT REGO
510(k) history	3 submissions · 3 cleared · 2008-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072213/> 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 26, 2026