FDA 510(k) Premarket Notification

K072213 PERIMETRICS PERIOMETER

CLEARED

Aug 29, 2008
386 days to decision

K072213 · 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	**Perimetrics, Inc.**
Location	Mission Viejo, CA, US
Contact	ALBERT REGO
510(k) history	3 submissions · 3 cleared · 2008-2025