

K081811 CARE3 PLUSJan 21, 2009
209 days to decisionK081811 · Product code: **EFB** · DentalSource: <https://www.510kdatabase.net/k081811/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Air-powered, Dental (EFB)
Date received	Jun 26, 2008
Decision date	Jan 21, 2009
Days to decision	209 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Nakanishi, Inc.
Location	Kanuma, JP
Contact	KEITH BARRITT
Website	https://www.nakanishi-inc.com
510(k) history	40 submissions · 40 cleared · 2001-2026

Nakanishi, Inc. is a precision medical device manufacturer based in Kanuma, Japan. The company specializes in motorized spindles and micro grinders for medical and industrial applications. Nakanishi has received FDA 510(k) clearances from total submissions since its first clearance in 2001. Dental devices represent the dominant category, accounting for 88% of the company's regulatory submissions. The company remains active, with its latest FDA 510(k) clearance in 2026. Recent cleared devices include motorized systems for dental surgery and scaling, air-powered polishing i...
