

K063007 POWERTOME PERIOTOMEOct 17, 2006
15 days to decisionK063007 · Product code: **EKX** · Dental
Source: <https://www.510kdatabase.net/k063007/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Direct Drive, Ac-powered (EKX)
Date received	Oct 2, 2006
Decision date	Oct 17, 2006
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Westport Medical, Inc.
Location	Salem, OR, US
Contact	RALPH WELLS
510(k) history	1 submissions · 1 cleared · 2006-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k063007/> Data sourced from FDA 510(k) public records (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. - Generated June 29, 2026