

**K170275 Tri Auto ZX2**Sep 11, 2017  
224 days to decisionK170275 · Product code: **EKX** · Dental  
Source: <https://www.510kdatabase.net/k170275/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Direct Drive, Ac-powered (EKX)
Date received	Jan 30, 2017
Decision date	Sep 11, 2017
Days to decision	224 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>J. Morita USA, Inc.</b>
Location	Irvine, CA, US
Contact	KEISUKE MORI
510(k) history	52 submissions · 52 cleared · 1988-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k170275/> 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 14, 2026