

**K222688 The Trust**Aug 9, 2024  
703 days to decisionK222688 · Product code: **EKX** · Dental  
Source: <https://www.510kdatabase.net/k222688/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Direct Drive, Ac-powered (EKX)
Date received	Sep 6, 2022
Decision date	Aug 9, 2024
Days to decision	703 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Genoss Co., Ltd.</b>
Location	Laguna Niguel, CA, US
Contact	Myeong Yujin
510(k) history	18 submissions · 18 cleared · 2007-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222688/> 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