

**K210903 IS Multi Unit Abutment System**Jul 23, 2021  
119 days to decisionK210903 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k210903/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Mar 26, 2021
Decision date	Jul 23, 2021
Days to decision	119 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neobiotech Co., Ltd.</b>
Location	Santa Fe Springs, CA, US
Contact	Young-Ku Heo
510(k) history	17 submissions · 17 cleared · 2004-2024

**REGULATORY CONSULTANT**

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Consulting firm	<b>Withus Group, Inc.</b>
Contact	April Lee

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA [accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k210903/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 25, 2026