

**K220251 Neodent Implant System - GM Narrow Implant System**Sep 6, 2022  
218 days to decisionK220251 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k220251/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 31, 2022
Decision date	Sep 6, 2022
Days to decision	218 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Jjgc Industria E Comercio DE Materiais Dentarios S.A.</b>
Location	Curitiba, BR
Contact	Mariana Soares Hartmann
510(k) history	28 submissions · 28 cleared · 2016-2023

**REGULATORY CONSULTANT**

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Consulting firm	<b>Straumann USA, LLC</b>
Contact	Jennifer M. Jackson

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/k220251/> 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 13, 2026