

K220390 CreoDent Solidex Customized Abutment and ScrewDec 2, 2022
294 days to decisionK220390 · Product code: **NHA** · Dental
Source: <https://www.510kdatabase.net/k220390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Feb 11, 2022
Decision date	Dec 2, 2022
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Creodent Prosthetics, Ltd.
Location	New York, NY, US
Contact	Calvin Shim
510(k) history	9 submissions · 9 cleared · 2013-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k220390/> 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 May 26, 2026