

**K183229 Argen Clear Aligner**Jun 7, 2019  
199 days to decisionK183229 · Product code: **NXC** · Dental  
Source: <https://www.510kdatabase.net/k183229/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aligner, Sequential (NXC)
Date received	Nov 20, 2018
Decision date	Jun 7, 2019
Days to decision	199 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Argen Corporation</b>
Location	San Diego, CA, US
Contact	Paul Cascone
510(k) history	5 submissions · 5 cleared · 2016-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k183229/> 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 26, 2026