

**K191024 xPORT Lens Fragmentation System**Aug 13, 2019  
117 days to decisionK191024 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k191024/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Apr 18, 2019
Decision date	Aug 13, 2019
Days to decision	117 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Carl Zeiss Meditec Cataract Technology, Inc.</b>
Location	Reno, NV, US
Contact	Gary Mocnik
510(k) history	7 submissions · 7 cleared · 2019-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k191024/> 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