

**K192540 Kitazato ET Catheters**May 1, 2020  
228 days to decisionK192540 · Product code: **MQF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k192540/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Assisted Reproduction (MQF)
Date received	Sep 16, 2019
Decision date	May 1, 2020
Days to decision	228 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kitazato Corporation</b>
Location	Tokyo, JP
Contact	Futoshi Inoue
510(k) history	13 submissions · 13 cleared · 2017-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k192540/> 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 25, 2026