

**K213881 Celerity Incubator**Jul 21, 2022  
220 days to decisionK213881 · Product code: **FRC** · General Hospital  
Source: <https://www.510kdatabase.net/k213881/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Biological Sterilization Process (FRC)
Date received	Dec 13, 2021
Decision date	Jul 21, 2022
Days to decision	220 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>STERIS Corporation</b>
Location	Mentor, OH, US
Contact	Gregory Land
510(k) history	204 submissions · 202 cleared · 1997-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k213881/> 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 13, 2026