

# K173629 CELERITY 20 Steam Process Challenge Device for Gravity Cycles

Feb 21, 2018  
89 days to decision

K173629 · Product code: **FRC** · General Hospital  
Source: <https://www.510kdatabase.net/k173629/>

## SUBMISSION DETAILS

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Biological Sterilization Process (FRC)
Date received	Nov 24, 2017
Decision date	Feb 21, 2018
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

## APPLICANT

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Company	<b>STERIS Corporation</b>
Location	Mentor, OH, US
Contact	Anthony Piotrkowski
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/k173629/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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