

**K222137 OK Plus Indicator**Dec 5, 2022  
139 days to decisionK222137 · Product code: **JOJ** · General Hospital  
Source: <https://www.510kdatabase.net/k222137/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Indicator, Physical/chemical Sterilization Process (JOJ)
Date received	Jul 19, 2022
Decision date	Dec 5, 2022
Days to decision	139 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Propper Manufacturing Co., Inc.</b>
Location	Long Island, NY, US
Contact	Andrew Sharavara
510(k) history	7 submissions · 7 cleared · 2014-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222137/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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