FDA 510(k) Premarket Notification

K222137 OK Plus Indicator

CLEARED

Dec 5, 2022
139 days to decision

K222137 · Product code: **JOJ** · General Hospital
Source: https://www.510kdatabase.net/k222137/

SUBMISSION DETAILS

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

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