

**K834353 OK FLASH STERILIZATION INDICATOR**Mar 19, 1984  
97 days to decisionK834353 · Product code: **JOJ** · General HospitalSource: <https://www.510kdatabase.net/k834353/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Physical/chemical Sterilization Process (JOJ)
Date received	Dec 13, 1983
Decision date	Mar 19, 1984
Days to decision	97 days
Third-party review	No

**APPLICANT**

---

Company	<b>Propper Mfg. Co., Inc.</b>
Location	Mchenry, IL, US
510(k) history	25 submissions · 25 cleared · 1977-2012

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k834353/> 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 26, 2026