

**K871672 STERILIZATION INDICATOR FOR FLASH  
STERILIZATION**Jun 1, 1987  
34 days to decisionK871672 · Product code: LRT · General Hospital  
Source: <https://www.510kdatabase.net/k871672/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Sterilization (LRT)
Date received	Apr 28, 1987
Decision date	Jun 1, 1987
Days to decision	34 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871672/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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