FDA 510(k) Premarket Notification

K834353 OK FLASH STERILIZATION INDICATOR

CLEARED

Mar 19, 1984
97 days to decision

K834353 · Product code: **JOJ** · General Hospital
Source: 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	**Propper Mfg. Co., Inc.**
Location	Mchenry, IL, US
510(k) history	25 submissions · 25 cleared · 1977-2012