FDA 510(k) Premarket Notification

K770590 SURGI-KIT

CLEARED

Apr 14, 1977
17 days to decision

K770590 · Product code: **KCT** · General Hospital
Source: https://www.510kdatabase.net/k770590/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilization Wrap Containers, Trays, Cassettes & Other Accessories (KCT)
Date received	Mar 28, 1977
Decision date	Apr 14, 1977
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Ethox Corp.**
Location	Mchenry, IL, US
510(k) history	19 submissions · 18 cleared · 1977-2004

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k770590/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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