FDA 510(k) Premarket Notification

K830739 PRO-GUARD

CLEARED

Mar 29, 1983
21 days to decision

K830739 · Product code: **FRG** · General Hospital
Source: https://www.510kdatabase.net/k830739/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Mar 8, 1983
Decision date	Mar 29, 1983
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Surgicot, Inc.**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1977-1997