

K830739 PRO-GUARDMar 29, 1983
21 days to decisionK830739 · 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

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Device record: <https://www.510kdatabase.net/k830739/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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