

K842405 DR. GOULD HAND PACK SUSTOM STERILAug 15, 1984
57 days to decisionK842405 · Product code: **FRG** · General Hospital
Source: <https://www.510kdatabase.net/k842405/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Jun 19, 1984
Decision date	Aug 15, 1984
Days to decision	57 days
Third-party review	No

APPLICANT

Company	Pro-Lab, Inc.
Location	Mchenry, IL, US
510(k) history	27 submissions · 26 cleared · 1984-1995

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Device record: <https://www.510kdatabase.net/k842405/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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