

K840342 PDI PREPMASTER WET PACK PATIENT PREP KITMar 27, 1984
81 days to decisionK840342 · Product code: **FRG** · General Hospital
Source: <https://www.510kdatabase.net/k840342/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Jan 6, 1984
Decision date	Mar 27, 1984
Days to decision	81 days
Third-party review	No

APPLICANT

Company	Nice-Pak Products, Inc.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1978-1998

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Device record: <https://www.510kdatabase.net/k840342/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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