FDA 510(k) Premarket Notification

K840342 PDI PREPMASTER WET PACK PATIENT PREP KIT

CLEARED

Mar 27, 1984
81 days to decision

K840342 · 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