FDA 510(k) Premarket Notification

K903908 REPAK SURGICAL PACKS

CLEARED

Oct 26, 1990
64 days to decision

K903908 · Product code: **KKX** · General Hospital
Source: https://www.510kdatabase.net/k903908/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Aug 23, 1990
Decision date	Oct 26, 1990
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**Repack Surgical Enterprises, Inc.**
Location	Cincinnati, OH, US
Contact	BARBARA STANEWICK
510(k) history	2 submissions · 2 cleared · 1988-1990