

K903908 REPAK SURGICAL PACKSOct 26, 1990
64 days to decisionK903908 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k903908/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 7, 2026