FDA 510(k) Premarket Notification

K897120 EQUIP COVERS, STERILE

CLEARED

Jan 4, 1990
9 days to decision

K897120 · Product code: **KKX** · General Hospital
Source: https://www.510kdatabase.net/k897120/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Dec 26, 1989
Decision date	Jan 4, 1990
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Amedic USA**
Location	Phoenix, AZ, US
Contact	THEO RUSSO-LARSSON
510(k) history	10 submissions · 10 cleared · 1987-1990