FDA 510(k) Premarket Notification

K894433 BATES BLANKET

CLEARED

Aug 11, 1989
25 days to decision

K894433 · Product code: **KKX** · General Hospital
Source: https://www.510kdatabase.net/k894433/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Jul 17, 1989
Decision date	Aug 11, 1989
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Joe B. Bates, MD, FAAP**
Location	Tyler, TX, US
Contact	BATES, MD
510(k) history	1 submissions · 1 cleared · 1989-1989