FDA 510(k) Premarket Notification

K880963 HAC AUTOLOGY CENTERS DISPENSING KIT FOR FIB.CON.

CLEARED

Jun 24, 1988
108 days to decision

K880963 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k880963/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Mar 8, 1988
Decision date	Jun 24, 1988
Days to decision	108 days
Third-party review	No

APPLICANT

Company	**Buckman Co., Inc.**
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998