FDA 510(k) Premarket Notification

K821096 A-LINER

CLEARED

May 14, 1982
24 days to decision

K821096 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k821096/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Apr 20, 1982
Decision date	May 14, 1982
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Martell Medical Products, Inc.**
Location	Walker, MI, US
510(k) history	7 submissions · 7 cleared · 1981-1995