FDA 510(k) Premarket Notification

K854877 ACU-SYRINGE W/WO NEEDLES

CLEARED

Feb 19, 1986
75 days to decision

K854877 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k854877/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Dec 6, 1985
Decision date	Feb 19, 1986
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**Acuderm, Inc.**
Location	Mchenry, IL, US
Contact	CHARLES YEH
510(k) history	13 submissions · 13 cleared · 1983-1994