FDA 510(k) Premarket Notification

K895890 AUTOPEN

CLEARED

Feb 16, 1990
134 days to decision

K895890 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k895890/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 5, 1989
Decision date	Feb 16, 1990
Days to decision	134 days
Third-party review	No

APPLICANT

Company	**Ulster Scientific, Inc.**
Location	Mchenry, IL, US
Contact	F CUNNINGHAM
510(k) history	20 submissions · 20 cleared · 1980-1994