FDA 510(k) Premarket Notification

K800213 PHARMACY ADDITIVE SYRINGE SET

CLEARED

Feb 26, 1980
25 days to decision

K800213 · Product code: **KYW** · General Hospital
Source: https://www.510kdatabase.net/k800213/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Liquid Medication, Graduated (KYW)
Date received	Feb 1, 1980
Decision date	Feb 26, 1980
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Cutter Laboratories, Inc.**
Location	Mchenry, IL, US
Website	https://www.bayer.com
510(k) history	39 submissions · 39 cleared · 1976-1986