FDA 510(k) Premarket Notification

K771539 STEPPER

CLEARED

Oct 7, 1977
57 days to decision

K771539 · Product code: **KYX** · General Hospital
Source: https://www.510kdatabase.net/k771539/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dispenser, Liquid Medication (KYX)
Date received	Aug 11, 1977
Decision date	Oct 7, 1977
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Tridak**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977