FDA 510(k) Premarket Notification

K792096 2 FR KELMAN CHAMBER MAINTAINER

CLEARED

Nov 16, 1979
29 days to decision

K792096 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k792096/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Oct 18, 1979
Decision date	Nov 16, 1979
Days to decision	29 days
Third-party review	No

APPLICANT

Company	**Heyer Schulte Corp.**
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1976-1979