FDA 510(k) Premarket Notification

K820788 CLIRANS PRIMING SET

CLEARED

Apr 6, 1982
14 days to decision

K820788 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k820788/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 23, 1982
Decision date	Apr 6, 1982
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Terumo Medical Corp.**
Location	Elkton, MD, US
510(k) history	143 submissions · 143 cleared · 1980-2011