FDA 510(k) Premarket Notification

K896892 KAWASUMI Y-TYPE BLOOD ADMINISTRATION SET

CLEARED

Sep 12, 1990
278 days to decision

K896892 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k896892/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 8, 1989
Decision date	Sep 12, 1990
Days to decision	278 days
Third-party review	No

APPLICANT

Company	**Kawasumi Laboratories Co., Ltd.**
Location	Canoga Park, CA, US
Contact	KENJIRO TANI
510(k) history	18 submissions · 18 cleared · 1987-2000