FDA 510(k) Premarket Notification

K902482 VITALMIX (SET & BAG)

CLEARED

Jul 2, 1990
28 days to decision

K902482 · Product code: **LHI** · General Hospital
Source: https://www.510kdatabase.net/k902482/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Jun 4, 1990
Decision date	Jul 2, 1990
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Pacific Device, Inc.**
Location	San Diego, CA, US
Contact	MIKE SHANNON
510(k) history	5 submissions · 5 cleared · 1990-1997