FDA 510(k) Premarket Notification

K031861 MIX2VIAL TRANSFER DEVICE

CLEARED

Jul 29, 2003
43 days to decision

K031861 · Product code: **LHI** · General Hospital
Source: https://www.510kdatabase.net/k031861/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Jun 16, 2003
Decision date	Jul 29, 2003
Days to decision	43 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medimop Medical Projects, Ltd.**
Location	Washington, DC, US
Contact	BENNY ARAZY
510(k) history	12 submissions · 12 cleared · 1996-2016