FDA 510(k) Premarket Notification

K062482 VENTED VIAL ADAPTER TRANSFER DEVICE

CLEARED

Nov 3, 2006
71 days to decision

K062482 · Product code: **LHI** · General Hospital
Source: https://www.510kdatabase.net/k062482/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Aug 24, 2006
Decision date	Nov 3, 2006
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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