

**K062482 VENTED VIAL ADAPTER TRANSFER DEVICE**

Nov 3, 2006  
71 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k062482/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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