

**K031861 MIX2VIAL TRANSFER DEVICE**Jul 29, 2003  
43 days to decisionK031861 · Product code: LHI · General Hospital  
Source: <https://www.510kdatabase.net/k031861/>**SUBMISSION DETAILS**

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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**

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Company	<b>Medimop Medical Projects, Ltd.</b>
Location	Washington, DC, US
Contact	BENNY ARAZY
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/k031861/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 3, 2026