

**K902482 VITALMIX (SET & BAG)**Jul 2, 1990  
28 days to decision

K902482 · Product code: LHI · General Hospital

Source: <https://www.510kdatabase.net/k902482/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902482/>; 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 May 26, 2026