

**K260301 Vial Adapter**Jun 5, 2026  
126 days to decision

K260301 · Product code: LHI · General Hospital

Source: <https://www.510kdatabase.net/k260301/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Jan 30, 2026
Decision date	Jun 5, 2026
Days to decision	126 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hangzhou Qiantang Longyue Biotechnology Co., Ltd.</b>
Location	Hangzhou, CN
Contact	Meijun Chen
510(k) history	5 submissions · 5 cleared · 2023-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k260301/> 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 9, 2026