FDA 510(k) Premarket Notification

K260301 Vial Adapter

CLEARED

Jun 5, 2026
126 days to decision

K260301 · Product code: **LHI** · General Hospital
Source: https://www.510kdatabase.net/k260301/

SUBMISSION DETAILS

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

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