

K213313 Veinoflow SCD (Model: LBTK-M-I 5006)Feb 9, 2022
128 days to decisionK213313 · Product code: **JOW** · Cardiovascular
Source: <https://www.510kdatabase.net/k213313/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Oct 4, 2021
Decision date	Feb 9, 2022
Days to decision	128 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Dalian Labtek Science & Development Co., Ltd.
Location	Shanghai, CN
Contact	Yao Cui
510(k) history	3 submissions · 3 cleared · 2012-2022

REGULATORY CONSULTANT

Consulting firm	Shanghai CV Technology Co., Ltd.
Contact	Doris Dong

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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