

**K200154 B&J DVT Compression Devices MHH800/MHH800SQ**Jul 8, 2020  
168 days to decisionK200154 · Product code: **JOW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k200154/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Jan 22, 2020
Decision date	Jul 8, 2020
Days to decision	168 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>B&amp;J Manufacturing , Ltd.</b>
Location	Shenzhen, CN
Contact	Billy Yu
510(k) history	4 submissions · 4 cleared · 2020-2020

**REGULATORY CONSULTANT**

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Consulting firm	<b>Shenzhen Joyantech Consulting Co., Ltd.</b>
Contact	Fu Ailing

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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