

**K210885 Dayspring**Apr 23, 2021  
29 days to decisionK210885 · Product code: **JOW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k210885/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Mar 25, 2021
Decision date	Apr 23, 2021
Days to decision	29 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Koya Medical, Inc.</b>
Location	Oakland, CA, US
Contact	Andy Doraiswamy
510(k) history	3 submissions · 3 cleared · 2021-2024

**REGULATORY CONSULTANT**

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Consulting firm	<b>Biodesign Regulatory Services, LLC</b>
Contact	Alex Chang

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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