

K232006 LEGACY® IPCAug 4, 2023
29 days to decisionK232006 · Product code: **JOW** · Cardiovascular
Source: <https://www.510kdatabase.net/k232006/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Jul 6, 2023
Decision date	Aug 4, 2023
Days to decision	29 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Ig Technology, Ltd.
Location	Sheffield, GB
Contact	Ivan Green
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	Third Party Review Group, LLC
Contact	Dave Yungvirt

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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