

K203117 TissueTak deviceJan 7, 2021
83 days to decisionK203117 · Product code: **GDW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k203117/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Oct 16, 2020
Decision date	Jan 7, 2021
Days to decision	83 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Via Surgical , Ltd.
Location	Providence, RI, US
Contact	Ofek Levin
510(k) history	4 submissions · 4 cleared · 2014-2021

REGULATORY CONSULTANT

Consulting firm	Orly Maor
Contact	Orly Maor

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k203117/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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