

**K221070 DP4 Microneedling device**Dec 20, 2022  
252 days to decisionK221070 · Product code: **QAI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k221070/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Microneedle Device (QAI)
Date received	Apr 12, 2022
Decision date	Dec 20, 2022
Days to decision	252 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Equipped USA, LLC</b>
Location	Newport Beach, CA, US
Contact	Shaun Kerrigan
510(k) history	1 submissions · 1 cleared · 2022-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k221070/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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