

**K222199 Collagen P.I.N. (Percutaneous Induction Needling)**Oct 21, 2022  
91 days to decisionK222199 · Product code: **QAI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k222199/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Microneedle Device (QAI)
Date received	Jul 22, 2022
Decision date	Oct 21, 2022
Days to decision	91 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Induction Therapies, LLC</b>
Location	Louisville, KY, US
Contact	Amelia Aslam
510(k) history	1 submissions · 1 cleared · 2022-2022

**REGULATORY CONSULTANT**

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Consulting firm	<b>Technology Sciences Group, Inc.</b>
Contact	Laurie A. Clarke

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222199/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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