FDA 510(k) Premarket Notification

K222199 Collagen P.I.N. (Percutaneous Induction Needling)

CLEARED

Oct 21, 2022
91 days to decision

K222199 · Product code: **QAI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k222199/

SUBMISSION DETAILS

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

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

REGULATORY CONSULTANT

Consulting firm	**Technology Sciences Group, Inc.**
Contact	Laurie A. Clarke

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