

K202019 Maven Patient-Specific InstrumentationJan 14, 2021
177 days to decisionK202019 · Product code: **HSN** · Orthopedic
Source: <https://www.510kdatabase.net/k202019/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Ankle, Semi-constrained, Cemented, Metal/polymer (HSN)
Date received	Jul 21, 2020
Decision date	Jan 14, 2021
Days to decision	177 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Paragon 28, Inc.
Location	Englewood, CO, US
Contact	Haylie Hertz
Website	https://paragon28.com
510(k) history	50 submissions · 50 cleared · 2017-2026

Paragon 28, Inc. is a foot and ankle surgical device company based in Englewood, US. Established in 2010, the company specializes in innovative solutions for foot and ankle procedures. Paragon 28 has received FDA 510(k) clearances from total submissions since 2017. The company's portfolio is entirely focused on Orthopedic devices. Recent clearances include plating systems, nail systems, external fixation devices, and total ankle replacement systems. The latest FDA 510(k) clearance was in 2026, reflecting active ongoing regulatory engagement. The company's product range en...

REGULATORY CONSULTANT

Consulting firm	Musculoskeletal Clinical Regulatory Advisers, LLC
Contact	Haylie Hertz

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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Device record: <https://www.510kdatabase.net/k202019/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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