

K251862 External Fixation Bone DistractorAug 11, 2025
55 days to decisionK251862 · Product code: **KTT** · Orthopedic
Source: <https://www.510kdatabase.net/k251862/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Jun 17, 2025
Decision date	Aug 11, 2025
Days to decision	55 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Paragon 28, Inc.
Location	Englewood, CO, US
Contact	Jacqueline Sloan
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...

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Device record: <https://www.510kdatabase.net/k251862/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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