

## K262047 Grappler Suture Anchor System

Aug 3, 2026  
46 days to decision

K262047 · Product code: **MBI** · Orthopedic  
Source: <https://www.510kdatabase.net/k262047/>

### SUBMISSION DETAILS

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|                       |                                                      |
|-----------------------|------------------------------------------------------|
| Decision              | Substantially Equivalent (Cleared)                   |
| Submission type       | Special                                              |
| Device classification | Fastener, Fixation, Nondegradable, Soft Tissue (MBI) |
| Date received         | Jun 18, 2026                                         |
| Decision date         | Aug 3, 2026                                          |
| Days to decision      | 46 days                                              |
| Third-party review    | No                                                   |
| Combination product   | No                                                   |
| PCCP authorized       | No                                                   |
| Summary / Statement   | Summary                                              |

### APPLICANT

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|                |                                                           |
|----------------|-----------------------------------------------------------|
| Company        | <b>Paragon 28, Inc.</b>                                   |
| Location       | Englewood, CO, US                                         |
| Contact        | Brittanie Dogue                                           |
| Website        | <a href="https://paragon28.com">https://paragon28.com</a> |
| 510(k) history | 53 submissions · 53 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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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k262047/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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