FDA 510(k) Premarket Notification

# K252367 InbellaMulti System

**CLEARED**

Oct 15, 2025
77 days to decision

K252367 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k252367/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrosurgical, Cutting & Coagulation & Accessories (GEI) |
| Date received | Jul 30, 2025 |
| Decision date | Oct 15, 2025 |
| Days to decision | 77 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Inbella Medical, Inc.** |
| Location | Richmond Hill, CA |
| Contact | Michael Kreindel |
| 510(k) history | 4 submissions · 4 cleared · 2025-2026 |

## REGULATORY CONSULTANT

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|---|---|
| Consulting firm | **Hogan Lovells US LLP** |
| Contact | Janice Hogan |

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