

**K990220 MODIFICATION OF GIP/MEDI-GLOBE ULTRASOUND NEEDLE SYSTEM**

Mar 31, 1999  
68 days to decision

K990220 · Product code: **FCG** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k990220/>

**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Biopsy Needle (FCG)                |
| Date received         | Jan 22, 1999                       |
| Decision date         | Mar 31, 1999                       |
| Days to decision      | 68 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|                |   |
|----------------|---|
| Company        | <b>Medi-Globe Corp.</b>                 |
| Location       | Irvine, CA, US                          |
| Contact        | MICHELLE L FIELDS                       |
| 510(k) history | 18 submissions · 17 cleared · 1990-2006 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k990220/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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