

**K020102 IQTEQ SPIROMETER**Mar 21, 2003  
434 days to decisionK020102 · Product code: **BZG** · Anesthesiology  
Source: <https://www.510kdatabase.net/k020102/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Spirometer, Diagnostic (BZG)       |
| Date received         | Jan 11, 2002                       |
| Decision date         | Mar 21, 2003                       |
| Days to decision      | 434 days                           |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Iqteq Development</b>              |
| Location       | Cape Town, Western Cape, ZA           |
| Contact        | RAY WRIGHT                            |
| 510(k) history | 1 submissions · 1 cleared · 2003-2003 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020102/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 25, 2026