

**K190220 Vista FS, Vista FS Liquid**Jun 10, 2019  
125 days to decisionK190220 · Product code: **MVL** · Dental  
Source: <https://www.510kdatabase.net/k190220/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Cord, Retraction (MVL)             |
| Date received         | Feb 5, 2019                        |
| Decision date         | Jun 10, 2019                       |
| Days to decision      | 125 days                           |
| Third-party review    | No                                 |
| Combination product   | No                                 |
| PCCP authorized       | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|--|
| Company        | <b>Inter-Med/Vista Dental Products</b> |
| Location       | Racine, WI, US                         |
| Contact        | Alex Johnson                           |
| 510(k) history | 6 submissions · 6 cleared · 2017-2020  |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k190220/> 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 June 15, 2026