

**K112692 IDENTIFY**Aug 3, 2012  
323 days to decisionK112692 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k112692/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Accelerator, Linear, Medical (IYE) |
| Date received         | Sep 15, 2011                       |
| Decision date         | Aug 3, 2012                        |
| Days to decision      | 323 days                           |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Humediq GmbH</b>                   |
| Location       | Grafelfing, DE                        |
| Contact        | MORITZ HOYER                          |
| 510(k) history | 1 submissions · 1 cleared · 2012-2012 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k112692/> 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 17, 2026