

**K081774 SOLX 790 TITANIUM SAPPHIRE LASER**Sep 12, 2008  
81 days to decisionK081774 · Product code: **HQF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k081774/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Laser, Ophthalmic (HQF)            |
| Date received         | Jun 23, 2008                       |
| Decision date         | Sep 12, 2008                       |
| Days to decision      | 81 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Solx, Inc.</b>                     |
| Location       | Waltham, MA, US                       |
| Contact        | DOUG ADAMS                            |
| 510(k) history | 1 submissions · 1 cleared · 2008-2008 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081774/> 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 24, 2026