

K062930 LADARWAVE CUSTOMCORNEA WAVEFRONT SYSTEMNov 8, 2006
41 days to decisionK062930 · Product code: **NCF** · Ophthalmic
Source: <https://www.510kdatabase.net/k062930/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Abbreviated |
| Device classification | Aberrometer, Ophthalmic (NCF) |
| Date received | Sep 28, 2006 |
| Decision date | Nov 8, 2006 |
| Days to decision | 41 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Alcon Research, Ltd. |
| Location | Fort Worth, TX, US |
| Contact | JANET G JOHNSON |
| 510(k) history | 16 submissions · 16 cleared · 2000-2017 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062930/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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