

**K003512 LASER PACK I, LASER PACK II, PHACO PACK I,
PHACO PACK II**Feb 8, 2001
86 days to decisionK003512 · Product code: **HQC** · Ophthalmic
Source: <https://www.510kdatabase.net/k003512/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Nov 14, 2000
Decision date	Feb 8, 2001
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	A.R.C. Laser AG
Location	Salt Lake City, UT, US
Contact	DANIEL HOEFER
510(k) history	3 submissions · 3 cleared · 2000-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003512/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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