

K081382 DUCKWORTH & KENT, LTD., IMPLANTATION SYSTEM, MODEL DK7786Oct 23, 2008
160 days to decisionK081382 · Product code: **KYB** · Ophthalmic
Source: <https://www.510kdatabase.net/k081382/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	May 16, 2008
Decision date	Oct 23, 2008
Days to decision	160 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Duckworth & Kent, Ltd.
Location	Baldock, Hertfordshire, GB
Contact	MARTIN LOCK
510(k) history	6 submissions · 6 cleared · 2003-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081382/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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