FDA 510(k) Premarket Notification

K052290 AQUASOFT ALL-DAY & ALL-DAY T (HIOXIFILCON A) DAILY WEAR CONTACT LENS

CLEARED

May 15, 2006
265 days to decision

K052290 · Product code: **LPL** · Ophthalmic
Source: https://www.510kdatabase.net/k052290/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	Aug 23, 2005
Decision date	May 15, 2006
Days to decision	265 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Clearlab Pte , Ltd.**
Location	Chicago, IL, US
Contact	MARTIN DALSING
510(k) history	4 submissions · 4 cleared · 2001-2006