

K992589 RESOLUTION 5X (SPHERICAL) AND ULTRA GEL 5X (TORIC)(HIOXIFILCON A) SOFT DAILY WEAR CONTACT LENS, (CLEAR & BLUE VISIBILITY)

Aug 26, 1999
24 days to decision

K992589 · Product code: LPL · Ophthalmic
Source: <https://www.510kdatabase.net/k992589/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	Aug 2, 1999
Decision date	Aug 26, 1999
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Opti-Centre Laboratories
Location	Grand Junction, CO, US
Contact	MARTIN DALRING
510(k) history	6 submissions · 6 cleared · 1998-2005

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Device record: <https://www.510kdatabase.net/k992589/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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