

K181230 Largan 55 (methafilcon A) Daily Wear Soft (hydrophilic) Contact Lens, Largan 55 (methafilcon A) Daily Wear Soft (hydrophilic) Contact Lens for ASTIGMATISM, Largan 55 (methafilcon A) Daily Wear Soft (hydrophilic) Contact Lens for PRESBYOPIA

Jun 25, 2018
47 days to decision

K181230 · Product code: LPL · Ophthalmic
Source: <https://www.510kdatabase.net/k181230/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lenses, Soft Contact, Daily Wear (LPL)
Date received	May 9, 2018
Decision date	Jun 25, 2018
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Largan Medical Co., Ltd.
Location	Taichung, TW
Contact	Amy Tien
510(k) history	8 submissions · 8 cleared · 2017-2025

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k181230/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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