

# A Study to Evaluate the Clinical Performance and Safety of UNICON Silicone Hydrogel Daily Disposable Soft Contact Lenses

NCT07322211

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Status	RECRUITING
Phase	Not Applicable
Sponsor	Unicon Optical Co. Ltd.
Enrollment	35 participants

## Key Eligibility Criteria

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### Inclusion (9)

- Adult with an age e 18;
- Be a currently adapted soft contact lens wearer who successful wear of soft contact lenses in both eyes for a minimum of 5 days per week and 6 hours per day within 3 months prior to study screening (by verbal confirmation);
- Participant must be stop wearing any contact lenses more than 7 days prior to screening;
- Able to wear contact lenses within a range of power from -2.00D to -6.00D (0.25 D steps) in both eyes;
- Astigmatism of 1.25D or less in both eyes;
- ... and 4 more (see full listing online)

### Exclusion (15)

- Women who are currently pregnant;
- Women who are lactating or planning a pregnancy at the time of enrollment (by verbal confirmation at the screening visit);
- Any use of systemic or ocular medications (e.g., non-steroidal anti-inflammatory eye drops, ophthalmic steroids) within 30 days prior to study enrollment for which contact lens wear could be contraindicated, as determined by the investigator;
- Any current or history ocular or systemic disease which may interfere with contact lens wear, as determined by the investigator;
- Any current or previous orthokeratology treatment within 90 days prior to study enrollment, or planned for orthokeratology treatment during the study;
- ... and 10 more (see full listing online)

## Locations (2 total)

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Kaohsiung Medical University Chung-Ho Memorial Hospital, Kaohsiung City, Taiwan  
Linkou Chang Gung Memorial Hospital, Taoyuan District, Taiwan

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<https://clinicaltrials.gov/study/NCT07322211>

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