# Straylight as an additional indicator in surgical refractive procedures.

Published: 09-06-2011 Last updated: 29-04-2024

To evaluate whether straylight assessments [log(s)] can contribute to develop/refine the decision tree for the indications AC-PIOL, RLE or CE.

| Ethical review        | Approved WMO               |
|-----------------------|----------------------------|
| Status                | Pending                    |
| Health condition type | Vision disorders           |
| Study type            | Observational non invasive |

## Summary

#### ID

NL-OMON36178

**Source** ToetsingOnline

**Brief title** Straylight and surgical refractive procedures.

## Condition

• Vision disorders

Synonym refractive error

**Research involving** Human

## **Sponsors and support**

Primary sponsor: Oogziekenhuis Rotterdam Source(s) of monetary or material Support: SWOO

#### Intervention

Keyword: Anterior Chamber Phakic Intra-Ocular Lens, Refractive Lens Exchange, Straylight

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#### **Outcome measures**

#### **Primary outcome**

Straylight parameter s (logs[s])

Corrected distance visual acuity (CDVA), monocular and binocular

Contrast sensitivity

#### Secondary outcome

Optical characteristics of the eye.

Outcome of questionnaires.

# **Study description**

#### **Background summary**

Various means for the correction of refractive errors, including surgical procedures, are available nowadays. In the absence of early cataract, phakic intraocular lens (PIOL) implant is one of the suited procedures, whereas in the presence of early cataract, refractive lens exchange (RLE) is suited. The absence or presence of early cataract is assessed by ophthalmologists using slitlamp examination and classical parameters of visual function (visual acuity [VA] and contrast sensitivity [CS]). However, subclinical lens opacities may not be identified by slitlamp examination. Although these subclinical opacities may already cause a functional effect, it may not yet affect VA and CS tests. However, it may already affect the straylight parameter. In the presence of early cataract there may be a discrepancy between ophthalmologist's assessment of the degree of cataract and the patient\*s subjective complaints (e.g. functionally insignificant versus significant cataract). The main cause for such a discrepancy is glare caused by forward light scatter (i.e. straylight). Improvement of visual function after refractive correction may, in part, be due to a reduction of straylight after replacement of the lens, together with its opacities. It is expected that straylight measurement can be used to distinguish the relative contribution of refractive error and glare to visual function impairment.

#### **Study objective**

To evaluate whether straylight assessments [log(s)] can contribute to

develop/refine the decision tree for the indications AC-PIOL, RLE or CE.

#### Study design

Prospective observational case series (pre-post design)

#### Study burden and risks

Participants do not benefit. Risks are negligible. Patients will be subjected to study-related measurements twice (i.e. pre- and postoperatively). Measurements will be performed at the time of regular visits to Focus Clinic/The Rotterdam Eye Hospital. For the straylight measurements pupils will be dilated by eyedrops. On average, the extra time required will be 1 hour every visit.

# Contacts

#### Public

Oogziekenhuis Rotterdam

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# **Trial sites**

## Listed location countries

Netherlands

# **Eligibility criteria**

Age Adults (18-64 years)

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Elderly (65 years and older)

#### **Inclusion criteria**

Age > 18 years Informed consent Bilateral refractive treatment

#### **Exclusion criteria**

Ocular pathology other than lens opacities (e.g. cornea opacities,) Per- or postoperative complications Posterior capsule opacification Linguistic barrier

## Study design

## Design

| Study type:         | Observational non invasive      |
|---------------------|---------------------------------|
| Intervention model: | Other                           |
| Allocation:         | Non-randomized controlled trial |
| Masking:            | Open (masking not used)         |
| Control:            | Active                          |
| Primary purpose:    | Diagnostic                      |

#### Recruitment

| NL                        |             |
|---------------------------|-------------|
| Recruitment status:       | Pending     |
| Start date (anticipated): | 01-07-2011  |
| Enrollment:               | 60          |
| Туре:                     | Anticipated |

## **Ethics review**

Approved WMO

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| Date:              | 09-06-2011   |
|--------------------|--|
| Application type:  | First submission   |
| Review commission: | METC Erasmus MC, Universitair Medisch Centrum Rotterdam<br>(Rotterdam) |

# **Study registrations**

## Followed up by the following (possibly more current) registration

No registrations found.

## Other (possibly less up-to-date) registrations in this register

No registrations found.

## In other registers

**Register** CCMO **ID** NL36666.078.11