Cataract surgery in eyes with an idiopathic macular epiretinal membrane: the effect on the epiretinal membrane and the surgical outcome

Published: 25-04-2019 Last updated: 30-01-2025

To quantify progression of macular ERM, and the degree of correlation with phaco-energy, after phacoemulsification.

Ethical review Approved WMO **Status** Recruiting

Health condition type Eye disorders NEC

Study type Observational non invasive

Summary

ID

NL-OMON48514

Source

ToetsingOnline

Brief title

Cataract surgery & ERM

Condition

Eye disorders NEC

Synonym

macular epiretinal membrane, macular pucker

Research involving

Human

Sponsors and support

Primary sponsor: Oogziekenhuis Rotterdam

Source(s) of monetary or material Support: Stichting wetenschappelijk onderzoek

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Oogziekenhuis

Intervention

Keyword: Cataract, Epiretinal membrane, Pars plana vitrectomy

Outcome measures

Primary outcome

Progression of ERM and phaco-energy.

Secondary outcome

Stage of ERM (OCT).

Central foveal thickness (CFT; OCT).

Progression of ERM after phacoemulsification.

Presence of CME (OCT).

Visual acuity (Snellen).

Metamorphopsia (M-charts, see Matsumoto et al. 2003).

Indication for vitrectomy and ERM peeling.

Complications.

Study description

Background summary

There is, as yet, no consensus among cataract surgeons and vitreoretinal surgeons with respect to the optimal treatment strategy for patients with cataract and an epiretinal membrane (ERM). For ERM-related symptoms, pars plana vitrectomy (PPV) is usually combined with cataract extraction but when cataract surgery is indicated for these patients, the option of phacovitrectomy, although perhaps equally beneficial for some of them, is less often taken into consideration.

Study objective

To quantify progression of macular ERM, and the degree of correlation with phaco-energy, after phacoemulsification.

Study design

Prospective observational.

Study burden and risks

Participants do not benefit, risks and inconvenience are negligible.

Contacts

Public

Oogziekenhuis Rotterdam

Schiedamse Vest 180 Rotterdam 3011 BH NL

Scientific

Oogziekenhuis Rotterdam

Schiedamse Vest 180 Rotterdam 3011 BH NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years)

Inclusion criteria

Age >= 50 years.

Informed consent.

Senile cataract and idiopathic macular epiretinal membrane, stage 1, 2 or 3, on OCT.

Exclusion criteria

Secondary macular ERM.

Diabetic retinopathy or other vascular retinopathy.

Pseudoexfoliation syndrome.

Advanced glaucoma (with visual field defects).

Any ocular opacity that may prevent reliable OCT scans.

Any kind of maculopathy such as pucker and severe (subjective) metamorphopsia.

Study design

Design

Study type: Observational non invasive

Masking: Open (masking not used)

Control: Uncontrolled

Primary purpose: Diagnostic

Recruitment

NL

Recruitment status: Recruiting

Start date (anticipated): 16-08-2019

Enrollment: 120

Type: Actual

Ethics review

Approved WMO

Date: 25-04-2019

Application type: First submission

Review commission: METC Erasmus MC, Universitair Medisch Centrum Rotterdam

(Rotterdam)

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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 ID

CCMO NL69328.078.19