Ocular effect of TRPM8 agonist in patients with dry eye disease

Submission date	Recruitment status No longer recruiting	Prospectively registered		
17/03/2015		☐ Protocol		
Registration date 21/03/2015	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited 25/06/2020	Condition category Eye Diseases	Individual participant data		

Plain English summary of protocol

Background and study aims

Dry eye disease occurs when the eyes do not make enough tears or the tears evaporate too quickly, leading to the eyes drying out and becoming inflamed (red and swollen) and irritated. Our aim is to study the effect of topical administration of TRPM8 agonist in patients with mild to moderate dry eye disease.

Who can participate?

Patients with mild to moderate dry eye.

What does the study involve?

60 patients are randomly allocated to be treated with either TRPM8 agonist dissolved in distilled water, or distilled water only. Study medications will be topically applied twice on the upper eyelid. The severity of dry eye symptoms will be evaluated before and 1 hour after application.

What are the possible benefits and risks of participating? There are no benefits and risks involved in this study.

Where is the study run from?

Department of Ophthalmology, Chonnam National University Medical School and Hospital (South Korea).

When is the study starting and how long is it expected to run for? From January 2015 to March 2015.

Who is funding the study? Investigator initiated and funded (South Korea).

Who is the main contact? Pf. Kyung Chul Yoon kcyoon@jnu.ac.kr

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

Effect of topical administration of TRPM8 agonist in patients with dry eye disease: a single-center randomized double-masked vehicle-controlled study

Acronym

TRPM8 (transient receptor potential melastatin 8)

Study objectives

Topical administration of TRPM8 agonist may increase basal tear production in patients with mild to moderate dry eye disease. Also, it may provide short-term symptom relief of ocular dryness.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Institutional Review Board of Chonnam National University Hospital, 10/07/2014, IRB No. CNUH 2014-171

Study design

Single-center randomized double-masked vehicle-controlled study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Dry eye is a disorder of the tear film due to tear deficiency or excessive evaporation, which causes damage to the interpalpebral ocular surface and is associated with symptoms of ocular discomfort.

Interventions

TRPM8 agonist (1-(Diisopropyl-phosphinoyl)-nonane) dissolved in distilled water (2 mg/mL) or vehicle (distilled water) was topically delivered using the absorbent cotton gauze square (0.4 g rectangle (50 mm x 60 mm), CS-being, Daisan Cotton, Japan) and wiped twice across the closed eyelid. A loading volume of 0.5 mL of solution on cotton was used to wet the cotton.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

1-(Diisopropyl-phosphinoyl)-nonane

Primary outcome(s)

- 1. Basal tear secretion (baseline and every 20 minutes) assessed by Schirmer score
- 2. Dry eye symptom (baseline and after 60 minutes): using the questionnaire (0, no symptoms; 1, mild symptoms; 2, moderate symptoms; 3, severe symptoms; and 4, very severe symptoms)

Key secondary outcome(s))

- 1. Cooling sensation (baseline and every 5 minutes) assessed by visual analogue scale (VAS) (0 to 10)
- 2. Tear-film break up time (baseline and every 10 minutes) the time before the defect of fluorescein dye appeared in the stained tear film was measured and recorded (measured TBUT 3 times and averaged)
- 3. Corneal sensitivity (baseline and every 20 minutes) measured using the Cochet-Bonnet esthesiometer
- 4. Keratoepitheliopathy score (baseline and every 30 minutes) after staining the cornea with fluorescein dye, the score was obtained by multiplying the stained area (0-3) by stained density (0-3)

Area (0, no punctate staining; 1, area occupied less than 1/3 of the cornea; 2, area occupied 1/3 to 2/3 of the cornea; 3, area occupied greater than 2/3 of the cornea)

Density (0, no punctate staining; 1, sparse density; 2, moderate density; 3, high density and the overlapped lesions)

These outcomes were measured for 1 hour (60 minutes)

Completion date

01/03/2015

Eligibility

Key inclusion criteria

- 1. Dry eye symptoms for more than 3 months despite the use of artificial tears
- 2. Low tear film break-up time (TBUT) (≤ 7 seconds)
- 3. Low Schirmer score (≤ 10 mm/5 min)
- 4. Presence of corneal and conjunctival epithelial damage

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

- 1. History of any ocular disease other than DED
- 2. Meibomian gland dysfunction
- 3. Contact lens use
- 4. Ocular trauma or surgeries
- 5. Presence of an uncontrolled systemic disease that could affect ocular surface condition
- 6. Punctual plugs
- 7. Used any eye drops other than artificial tears
- 8. Used any systemic medication that can cause dry eye
- 9. Pregnant

Date of first enrolment

09/01/2015

Date of final enrolment

16/02/2015

Locations

Countries of recruitment

Korea, South

Study participating centre

Department of Ophthalmology, Chonnam National University Medical School and Hospital

42 Jebong-ro

Dong-gu

Gwangju

Korea, South

501-757

Sponsor information

Organisation

Chonnam National University Medical School and Hospital

ROR

https://ror.org/00f200z37

Funder(s)

Funder type

Other

Funder Name

Investigator initiated and funded (South Korea)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Other

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	26/06/2017	25/06/2020	Yes	No