

Topical antihistamines and mast cell stabilisers for treating seasonal and perennial allergic conjunctivitis (Review)

Castillo M, Scott NW, Mustafa MZ, Mustafa MS, Azuara-Blanco A



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TABLE OF CONTENTS

HEADER	1
ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
BACKGROUND	3
OBJECTIVES	4
METHODS	4
RESULTS	6
Figure 1.	8
Figure 2.	9
Figure 3.	10
Figure 4.	11
Figure 5.	20
Figure 6.	20
DISCUSSION	29
AUTHORS' CONCLUSIONS	29
ACKNOWLEDGEMENTS	29
REFERENCES	30
CHARACTERISTICS OF STUDIES	33
DATA AND ANALYSES	69
Analysis 1.1. Comparison 1 Olopatadine versus ketotifen, Outcome 1 Itching at 14 days.	69
Analysis 1.2. Comparison 1 Olopatadine versus ketotifen, Outcome 2 Tearing at 14 days.	70
ADDITIONAL TABLES	70
APPENDICES	72
CONTRIBUTIONS OF AUTHORS	76
DECLARATIONS OF INTEREST	76
SOURCES OF SUPPORT	76
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	77
NOTES	77

[Intervention Review]

Topical antihistamines and mast cell stabilisers for treating seasonal and perennial allergic conjunctivitis

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ABSTRACT

Background

Seasonal/perennial allergic conjunctivitis is the most common allergic conjunctivitis, usually with acute manifestations when a person is exposed to allergens and with typical signs and symptoms including itching, redness, and tearing. The clinical signs and symptoms of allergic conjunctivitis are mediated by the release of histamine by mast cells. Histamine antagonists (also called antihistamines) inhibit the action of histamine by blocking histamine H1 receptors, antagonising the vasoconstrictor, and to a lesser extent, the vasodilator effects of histamine. Mast cell stabilisers inhibit degranulation and consequently the release of histamine by interrupting the normal chain of intracellular signals.

Topical treatments include eye drops with antihistamines, mast cell stabilisers, non-steroidal anti-inflammatory drugs, combinations of the previous treatments, and corticosteroids. Standard treatment is based on topical antihistamines alone or topical mast cell stabilisers alone or a combination of treatments. There is clinical uncertainty about the relative efficacy and safety of topical treatment.

Objectives

The objective of this review was to assess the effects of topical antihistamines and mast cell stabilisers, alone or in combination, for use in treating seasonal and perennial allergic conjunctivitis.

Search methods

We searched CENTRAL (which contains the Cochrane Eyes and Vision Trials Register) (2014, Issue 7), Ovid MEDLINE, Ovid MEDLINE In-Process and Other Non-Indexed Citations, Ovid MEDLINE Daily, Ovid OLDMEDLINE (January 1946 to July 2014), EMBASE (January 1980 to July 2014), the *meta*Register of Controlled Trials (*mRCT*) (www.controlled-trials.com), ClinicalTrials.gov (www.clinicaltrials.gov) and the World Health Organization (WHO) International Clinical Trials Registry Platform (ICTRP) (www.who.int/ictrp/search/en). We did not use any date or language restrictions in the electronic searches for trials. We last searched the electronic databases on 17 July 2014. We also searched the reference lists of review articles and relevant trial reports for details of further relevant publications.

Topical antihistamines and mast cell stabilisers for treating seasonal and perennial allergic conjunctivitis (Review)

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1

Selection criteria

We included randomised controlled trials (RCTs) comparing topical antihistamine and mast cell stabilisers, alone or in combination, with placebo, no treatment or to any other antihistamine or mast cell stabiliser, or both, that examined people with seasonal or perennial allergic conjunctivitis, or both. The primary outcome was any participant-reported evaluation (by questionnaire) of severity of four main ocular symptoms: itching, irritation, watering eye (tearing), and photophobia (dislike of light), both separately and, if possible, by an overall symptom score. We considered any follow-up time between one week and one year.

Data collection and analysis

Two review authors independently extracted data and assessed risk of bias. Disagreements were resolved by discussion among review authors and the involvement of a third review author. We followed standard methodological approaches used by Cochrane.

Main results

We identified 30 trials with a total of 4344 participants randomised, with 17 different drugs or treatment comparisons. The following antihistamines and mast cell stabilisers were evaluated in at least one RCT: nedocromil sodium or sodium cromoglycate, olopatadine, ketotifen, azelastine, emedastine, levocabastine (or levocabastine), mequitazine, bepotastine besilate, combination of antazoline and tetryzoline, combination of levocabastine and pemirolast potassium. The most common comparison was azelastine versus placebo (nine studies).

We observed a large variability in reporting outcomes. The quality of the studies and reporting was variable, but overall the risk of bias was low. Trials evaluated only short-term effects, with a range of treatment of one to eight weeks. Meta-analysis was only possible in one comparison (olopatadine versus ketotifen). There was some evidence to support that topical antihistamines and mast cell stabilisers reduce symptoms and signs of seasonal allergic conjunctivitis when compared with placebo. There were no reported serious adverse events related to the use of topical antihistamine and mast cell stabilisers treatment.

Authors' conclusions

It seems that all reported topical antihistamines and mast cell stabilisers reduce symptoms and signs of seasonal allergic conjunctivitis when compared with placebo in the short term. However, there is no long-term data on their efficacy. Direct comparisons of different antihistamines and mast cell stabilisers need to be interpreted with caution. Overall, topical antihistamines and mast cell stabilisers appear to be safe and well tolerated. We observed a large variability in outcomes reported. Poor quality of reporting challenged the synthesis of evidence.

PLAIN LANGUAGE SUMMARY

Topical antihistamines and mast cell stabilisers for treating seasonal and perennial allergic conjunctivitis

Review question

Are treatments with eye drops of antihistamines and mast cell stabilisers, alone or in combination, effective and safe in people with seasonal and allergic conjunctivitis? The main outcome measure was eye symptoms reported by participants, including eye itching, irritation (burning sensation), watering eyes (tearing), and photophobia (dislike of light). We found 30 trials.

Background

Conjunctivitis refers to inflammation of the conjunctiva, which is the thin tissue that covers the sclera (white part of the eye). Seasonal and perennial allergic conjunctivitis is the most common type of allergic conjunctivitis. Although this condition does not cause sight loss, it can cause intense itching and eye watering. Eye drops with antihistamines or mast cell stabilisers, or both are commonly used.

Study characteristics

The evidence is current to July 2014. Among the 30 studies reviewed there were 17 different comparisons, including 4344 participants ranging in age between 4 and 85 years. The duration of treatment ranged from one to eight weeks. Ten out of 30 studies were funded by the drug manufacturer (8 totally and 2 partially funded); 20 studies did not report any source of funding. There was inconsistency in the way the effect of treatment was measured and reported. Overall risk of bias was low.

Key results

Topical antihistamines and mast cell stabilisers, alone or in combination, are safe and effective for reducing symptoms of seasonal and perennial allergic conjunctivitis. We found insufficient evidence to discern which topical antihistamines and mast cell stabilisers are the most effective.

BACKGROUND

Description of the condition

The conjunctiva forms the mucous membrane component of the ocular surface and is continually exposed to a wide variety of allergens. In sensitised individuals, pollen, plant, house dust mite or animal dander allergens dissolve in the tear film, cross the conjunctival epithelium, and activate mast cells in the substantia propria leading to the prevalent and unpleasant conditions known as seasonal allergic conjunctivitis (SAC) and perennial allergic conjunctivitis (PAC) (Anderson 2001).

SAC is the most common allergic conjunctivitis, usually with an acute or subacute manifestation characterised by peaks of self-limiting signs and symptoms that become persistent in repeated allergen stimulations during the pollen season. The hallmark signs and symptoms are itching, redness, and lid swelling. Patients may also complain of epiphora (watering eye), mucous (translucent) discharge, rhinitis (runny or stuffy nose), and eye-burning sensation (Leonardi 2008).

In PAC, the non-specific signs and symptoms of redness, burning, and chemosis (swelling of the conjunctiva) may persist with varying severity for months, accompanied by a low level of itching (Leonardi 2008).

Seasonal and perennial allergic conjunctivitis rarely cause visual disability.

Epidemiology

SAC is the most common allergic eye disorder, affecting up to 20% of the population in the United Kingdom at some point in their lives (Buckley 1998; Dart 1986). It has a varying prevalence in countries throughout the world (Beasley 1998). PAC is much rarer and typically results in milder symptoms than those seen in the seasonal condition. People with allergic conjunctivitis often have a personal history, family history, or both of atopic disorders. A recent epidemiological survey noted that up to 40% of 13 and 14-year-olds worldwide reported the symptoms of allergic conjunctivitis (see 'Clinical presentation' below) (Strachan 1997). It has been estimated that ocular symptoms are present in 40% to 60% of the allergic population (people with atopy, that is asthma

and eczema) (Ono 2005). Visual loss and severe complications are rare.

Biological mechanism

Allergic conjunctivitis results from a Type 1 (immediate) immunoglobulin E (IgE)-mediated hypersensitivity reaction. The allergen particles bind to and cross-link IgE molecules that are attached to tissue mast cells, resulting in mast cell degranulation and release of histamines and other inflammatory mediators into the tissue. Histamine is the primary contributor to the development of early-phase signs and symptoms of SAC (Abelson 1979). A comprehensive summary of these is provided in Hingorani 1997.

Clinical presentation

Allergic conjunctivitis is usually bilateral. The main defining feature of this type of conjunctivitis is intense itching. More than 75% of patients report this symptom when seeking treatment (Whitcup 2006). Other signs and symptoms include epiphora, hyperaemia (engorgement of blood vessels of the conjunctiva), micropapillae (little bumps on the inner surface of the eyelid), conjunctival chemosis (swelling of the conjunctiva), and thin mucous discharge.

If accompanied by rhinitis (sneezing, rhinorrhoea (runny nose), nasal itch or blockage), the symptom complex is sometimes referred to as rhinoconjunctivitis. Coughing and wheezing due to asthma may also co-exist. Diagnosis is usually on the basis of a careful allergy history and examination of the conjunctiva. Skin-prick tests, serum-specific IgE, and conjunctival allergen challenge may be used to confirm the allergic trigger.

Description of the intervention

A variety of drugs have been used to treat allergic conjunctivitis, which can be administered orally or applied directly to the eye. Systemic treatment with oral antihistamines is usually indicated when other symptoms are being treated along with the conjunctivitis, such as rhinitis. Topical treatments include antihistamines, mast cell stabilisers, non-steroidal anti-inflammatory drugs, and corticosteroids, the last being reserved for severe symptoms that

are unresponsive to other treatments and given for a short period of time.

How the intervention might work

The pharmacological effects of antihistamines are based on their abilities to block histamine H1 receptors. H1 receptors are activated by histamine, which has many actions. Histamine mediates the tissue response to injury (for example mechanical, thermal, infections, etc.). It is also a mediator of gastric acid secretion and may serve as a neurotransmitter. With respect to conjunctivitis, the action of antihistamines is to antagonise the vasoconstrictor, and, to a lesser extent, the vasodilator effects of histamine. Some antihistamines have central effects like sedation, which can influence their use when administered systemically.

Mast cell stabilisers inhibit degranulation by interrupting the normal chain of intracellular signals resulting from the cross-linking and activation of FcεRI by allergen (Cook 2002). They inhibit mast cell degranulation, the release of histamine, and the other preformed mediators and the arachidonic acid cascade (Leonardi 2005).

Why it is important to do this review

As allergic conjunctivitis is considered to be a leading cause of acute red eye presentation to hospital, optometrists, and general practitioners, it has a significant impact on health service provision. The spectrum of disease varies from mild to severe and can therefore interfere with quality of life. It also has the potential to compromise visual function, although this is rarely seen.

Standard treatment is topical antihistamines alone or in combination with topical mast cell stabilisers. There is clinical uncertainty about the relative efficacy of this treatment.

OBJECTIVES

The objective of this review was to assess the effects of topical antihistamines and mast cell stabilisers, alone or in combination, for use in treating seasonal and perennial allergic conjunctivitis.

METHODS

Criteria for considering studies for this review

Types of studies

We included randomised controlled trials (RCTs) only.

We excluded within-person trials (in which eyes were randomly allocated to different treatments).

Types of participants

We included trials that recruited children or adults with seasonal allergic conjunctivitis (SAC) or perennial allergic conjunctivitis (PAC). This review did not cover other allergic conjunctivitis entities such as vernal keratoconjunctivitis, atopic keratoconjunctivitis, and giant papillary conjunctivitis. We attempted to obtain separate data for participants with SAC or PAC where trials also included participants with other types of conjunctivitis.

The diagnosis was done clinically based on the presence of typical symptoms and signs.

We excluded trials conducted with the use of conjunctival allergen challenge or conjunctival provocation testing, as it would be difficult to compare the data with trials that have participants who were recruited with active symptomatic disease without provocation tests. In addition, information from trials using provocation or challenge might not be representative and generalisable of the condition observed in routine practice.

We excluded people with allergic eye disease associated with atopy.

Types of interventions

This review included trials comparing topical antihistamines and mast cell stabilisers, alone or in combination, with placebo, no treatment, or any other antihistamine and/or mast cell stabiliser. These included the medications listed in Appendix 1 and any other pure antihistamine alone or in combination with a mast cell stabiliser not originally listed.

We placed no restriction on the duration of treatment in the trials. We did not include any trials that used herbal, oral, nasal antihistamines and any other medication.

Types of outcome measures

Primary outcomes

The primary outcome was any participant-reported evaluation (by questionnaire) of severity of four main ocular symptoms: itching, irritation, watering eye, and photophobia (dislike of light), both separately and, if possible, by an overall symptom score. We considered any follow-up time between one week and one year.

As there are a variety of different questionnaires evaluating ocular symptoms, we tried to transform data to common measurements where possible to facilitate meta-analysis using the standardised mean difference. We used this only if studies utilised different participant-reported outcomes measuring a comparable range of symptoms.

Secondary outcomes

Secondary outcomes were measured at any reported follow-up time and included:

- Adverse events.
- Signs of hyperaemia and redness, chemosis, tarsal papillae (little bumps on inner surface of eyelid). Signs were clinician/investigator assessed and analysed using categorisation where appropriate.
- Duration of symptoms (days) of acute episodes.
- Incidence of acute episodes (per year).

A variety of numeric rating scales were used to assess participant- and clinician-reported symptoms in the included studies. Unless stated otherwise, higher scores represent higher levels of symptoms than lower scores.

Search methods for identification of studies

Electronic searches

We searched CENTRAL (which contains the Cochrane Eyes and Vision Trials Register) (2014, Issue 7), Ovid MEDLINE, Ovid MEDLINE In-Process and Other Non-Indexed Citations, Ovid MEDLINE Daily, Ovid OLDMEDLINE (January 1946 to July 2014), EMBASE (January 1980 to July 2014), the *metaRegister* of Controlled Trials (*mRCT*) (www.controlled-trials.com), ClinicalTrials.gov (www.clinicaltrials.gov) and the World Health Organization (WHO) International Clinical Trials Registry Platform (ICTRP) (www.who.int/ictcp/search/en). We did not use any date or language restrictions in the electronic searches for trials. We last searched the electronic databases on 17 July 2014.

See: Appendices for details of search strategies for CENTRAL (Appendix 2), MEDLINE (Appendix 3), EMBASE (Appendix 4), *mRCT* (Appendix 5), ClinicalTrials.gov (Appendix 6), and the ICTRP (Appendix 7).

Searching other resources

We searched the reference lists of review articles, book chapters, and relevant trial reports for details of further relevant publications. We planned to contact authors for data queries. We used the Science Citation Index to search for references that cite the studies that were included in the review.

Data collection and analysis

Selection of studies

Two review authors (MSM, MC/MZM) independently assessed the titles and abstracts resulting from the manual and electronic

searches for selection of studies for inclusion. We obtained full copies of all relevant or potentially relevant trials and assessed these according to the 'Criteria for considering studies for this review' section. The investigators were not masked to the names of authors, institutions, journal of publication, or results when they made their assessments. We resolved disagreements about whether a trial should be included by discussion with a third review author (AAB) and consensus. In cases where we needed additional information before we could make a decision about whether or not to include a trial, we planned to obtain this information by contacting the trial authors when author contact details were available. We attempted to obtain further information about any trial that had been published only as an abstract by contacting authors if contact details were available. If a full report was not available, we planned to assess and include data from abstracts for the review and meta-analysis. If there was not sufficient abstract data for the meta-analysis, we still planned to present the abstract data in the review to ensure no significant information was lost.

Using a form developed to document the process, we divided the full copies into two groups: 1) definitely include and 2) definitely exclude.

Agreement between the two review authors was recorded. At this stage, we excluded only those papers that both review authors had assigned to the 'definitely exclude' group. We documented and reported these exclusions in the review. We assessed all other papers for methodological quality.

Data extraction and management

As well as recording information about the methods used in the trial, we extracted the following information from the trial reports using a standardised form:

- details of participants (age, gender, setting, number in each group, comparability at baseline);
- details of interventions (dosage, schedule, compliance, comparison group, timing);
- outcomes (primary and secondary outcomes, adverse effects);
- other information (source of funding, declaration of interest).

Two review authors (MSM, MC/MZM) independently extracted the data for the primary and secondary outcomes on to a standardised form. One review author (MC) entered data into RevMan (RevMan 2014), and a second review author (AAB) checked the data entered into RevMan to ensure that no mistakes had been made. The review authors resolved any differences by discussion. If there was any doubt about the data of a trial, the review authors contacted the authors of the trial. Where studies were reported in more than one publication, we extracted data from each report separately. We then collated the information from the multiple data collection forms.

Assessment of risk of bias in included studies

Two review authors (MC, AAB) evaluated the risk of bias independently. They met after completion to identify disagreements. Review authors assessed trial quality according to the methods set out in Chapter 8 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011a). We used Cochrane's 'Risk of bias tool' and considered six domains: sequence generation (randomisation), allocation concealment, blinding (masking), incomplete outcome data (completeness of follow-up), selective outcome reporting, and free from other bias.

Two review authors (MC, AAB) assessed the risk of bias for each parameter and judged each parameter as low risk of bias, high risk of bias, or unclear (uncertain risk of bias where there was insufficient information to be able to judge).

We planned to perform a sensitivity analysis, if possible, excluding studies with high risk of bias to determine the effect of this decision on the results (Higgins 2011b).

Measures of treatment effect

Dichotomous data

For dichotomous data, we calculated odds ratios and their 95% confidence intervals.

Continuous data

For continuous data presented using a common scale, we calculated the mean difference and its 95% confidence interval. For overall symptom scores, we used the standardised mean difference was used.

Unit of analysis issues

The unit of randomisation was the individual participant (not eye) due to potential systemic absorption and effect in the fellow eye; we therefore only included studies that used the participant as the unit of randomisation. We considered cross-over trials if there was an adequate washout (48 hours) between the two treatment periods and if the treatment was compared with a control placebo. H1 antihistamine terminal elimination half-life values range from 2 hours for acrivastine to 27 hours for desloratadine (Simons 2002).

Dealing with missing data

If data were not obtainable we considered the potential impact of the missing data on the results and highlighted the potential impact in the 'Discussion' section of the review.

Assessment of heterogeneity

Before combining studies we planned to assess clinical heterogeneity by examination of the study details and statistical heterogeneity using the Chi^2 test and the I^2 statistic. Where we were unable to pool results we planned to provide a descriptive summary.

Assessment of reporting biases

We minimised reporting bias by maximising our search strategy to include research that had been published in different languages as well as unpublished research.

If appropriate, we planned to investigate publication bias by looking at a funnel plot of the data.

Data synthesis

When there were sufficient trials available without substantial heterogeneity, we combined studies in a meta-analysis using a random-effects model. Where this was not possible, we performed a narrative synthesis. As we evaluated a network of treatments, we also planned a network meta-analysis, provided suitable data were available.

Subgroup analysis and investigation of heterogeneity

Not applicable for this review.

Sensitivity analysis

We planned to perform sensitivity analyses, if possible, to assess how robust the results were to changes in methods such as:

1. excluding studies of lower methodological quality;
2. excluding unpublished studies;
3. excluding studies that assumed that eyes within a participant are independent.

RESULTS

Description of studies

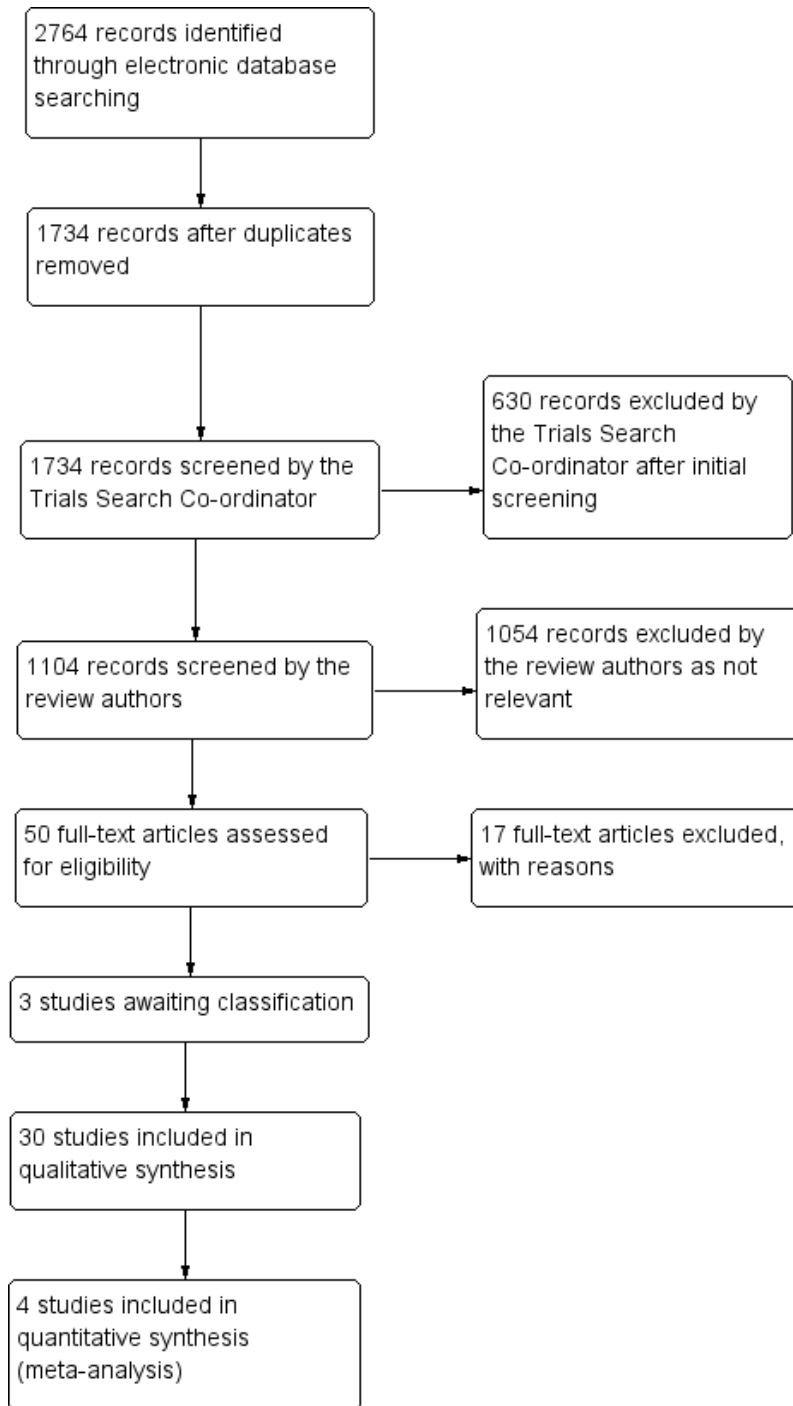
We have described the studies within the context of each comparison (see 'Effects of interventions' section).

Results of the search

The electronic searches yielded a total of 2764 references (Figure 1). The Trials Search Co-ordinator removed 1030 duplicate records, screened 1734 records, and removed 630 references that

were not relevant to the scope of this review. We screened the remaining 1104 references and discarded 1054 reports as not relevant. We obtained 50 full-text reports for potential inclusion in the review and included 30 studies (see 'Characteristics of included studies' section) and excluded 17 studies (see 'Characteristics of excluded studies' section). Currently three studies are still awaiting classification: [Scandashree 2013](#) has insufficient details on outcome measures, and [Jia 2012](#) and [Dharmistha 2013](#) have insufficient information on methods of allocation. If we are able to obtain further information on these studies, we will assess them in future updates of this review.

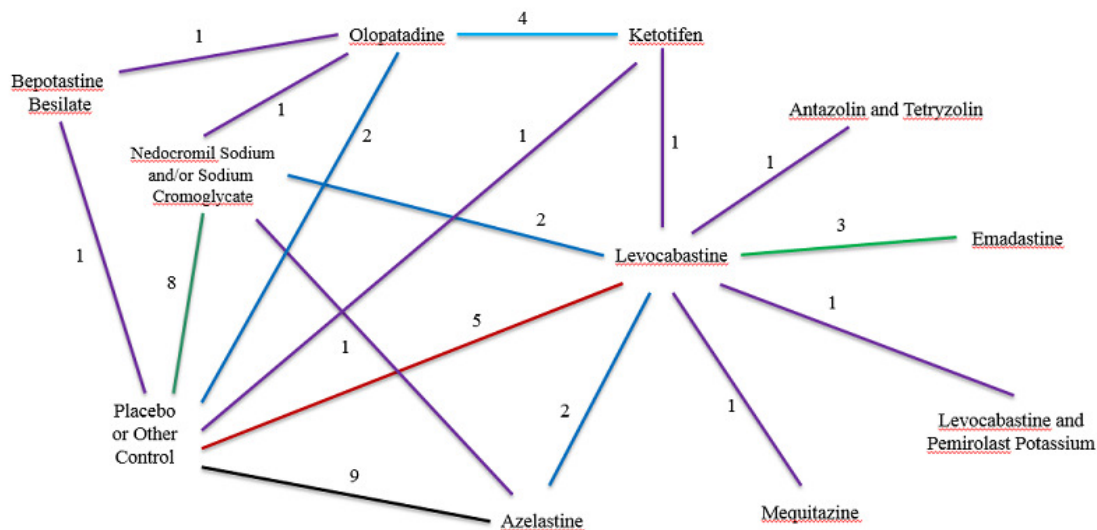
Figure 1. Results of searching for studies for inclusion in the review



Included studies

We included 30 trials with 17 different comparisons (see [Figure 2](#), a network diagram showing the number of studies contributing to each of the 17 comparisons).

Figure 2. Network diagram: Number of studies by treatment comparison



Excluded studies

We excluded 17 studies that did not meet the inclusion criteria (see 'Characteristics of excluded studies' section).

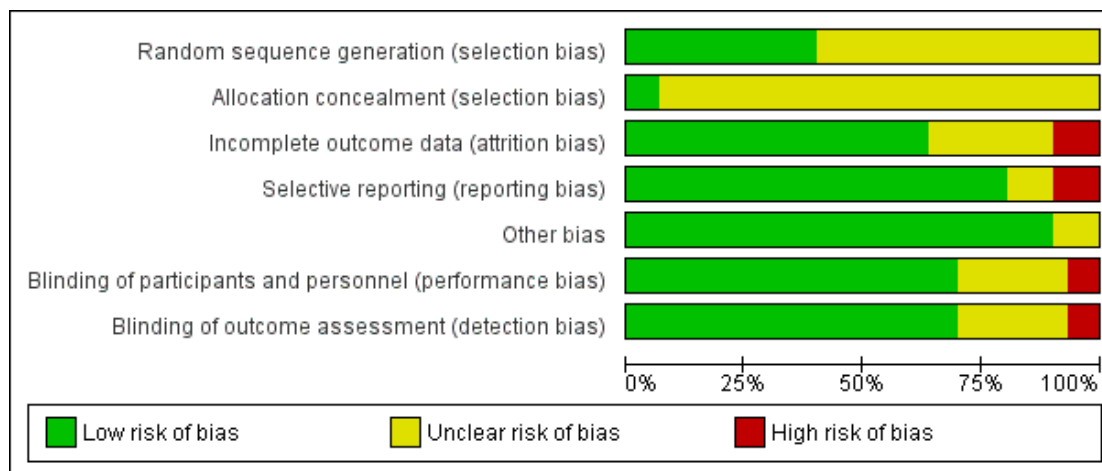
Risk of bias in included studies

We summarised risk of bias in [Figure 3](#) and [Figure 4](#). Most studies had low risk of bias regarding the masking of participants and investigators and selective reporting. Information on sequence generation and allocation concealment was frequently missing.

Figure 3. 'Risk of bias' summary: review authors' judgements about each 'Risk of bias' item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)
Avunduk 2005	+	?	+	+	+	+	+
Azevedo 1991	+	?	+	+	+	+	+
Canonica 2003	+	?	+	+	+	+	+
Carr 2013	+	?	+	+	+	+	+
Davies 1993	?	?	-	-	?	+	+
Fujishima 2008	+	?	+	+	+	?	?
Giede-Tuch 1998	?	?	+	+	+	+	+
Graue 1994	?	?	+	+	+	+	+
Hechanova 1984	?	?	-	-	+	+	+
Höfling-Lima 2001	+	?	+	+	+	+	+
James 2003	?	?	+	+	+	?	?
Katellaris 2002	+	?	?	+	+	+	+
Kidd 2003	+	?	?	+	+	+	+
Lanier 2001	?	?	?	+	+	-	-
Leino 1992	?	+	-	+	+	+	+
Lenhard 1997	?	?	?	+	+	?	?
McCabe 2012	+	?	+	+	+	-	-
Melamed 1994	?	?	+	+	+	+	+
Melamed 2000	?	?	+	+	+	+	+
Möller 1994	+	+	+	+	+	+	+
Nazarov 2003	?	?	+	+	+	+	+
Petzold 2002	?	?	?	?	?	?	?
Sabbah 1998	?	?	+	+	+	?	?
Sarker 2011	+	?	+	+	+	+	+
Secchi 2000a	?	?	+	+	+	+	+
Secchi 2000b	?	?	+	?	+	+	+
Trinquand 1999	?	?	?	?	?	?	?
Varguez-Rodriguez 2009	?	?	+	+	+	+	+
Verin 2001	?	?	?	+	+	+	+
Wertheimer 1997	+	?	?	-	+	?	?

Figure 4. 'Risk of bias' graph: review authors' judgements about each 'Risk of bias' item presented as percentages across all included studies.



Allocation

We judged the method of randomisation to be at low risk of selection bias in approximately 37% of the studies (Figure 3).

Allocation concealment was the item with the lowest proportion (less than 10%) of studies judged to be at low risk of bias. We judged more than 90% of studies as at unclear risk of selection bias on the basis of inadequate or no information provided on the method of allocation concealment.

Blinding

We judged the majority (approximately 62%) of studies as at low risk of detection bias (masking of outcome assessment) and considered a similar proportion as at low risk of performance bias (masking of participants and investigators) (Figure 3).

Incomplete outcome data

Over 50% of the studies have low risk of bias for this domain. We judged a lower proportion of studies (approximately 36%) with unclear risk of bias, as some of the trials did not conduct an intention-to-treat analysis, or no reasons for dropouts were described (Figure 3).

Selective reporting

We judged a highest proportion of studies (75%) with low risk of bias for this domain.

Other potential sources of bias

Not applicable for this review.

Effects of interventions

Drug comparisons

We have used the following order to describe the comparisons: placebo-controlled studies with mast cell stabilisers (comparison 1); placebo-controlled studies with antihistamines (comparisons 2 and 3, in order of number of studies available); and comparative studies of antihistamines and/or mast cell stabilisers (comparisons 4 to 16, in order of number of studies available). We described primary and safety outcomes for each comparison. Secondary outcomes were either not reported (duration of symptoms of acute episodes and incidence of acute episodes) or uncommonly reported and using different measures and scales (signs evaluated by an investigator) and thus were not analysed.

I. Nedocromil sodium/sodium cromoglycate versus placebo

The search identified eight eligible studies comparing the mast cell stabilisers nedocromil sodium or sodium cromoglycate against placebo (Azevedo 1991; Davies 1993; Hechanova 1984; James 2003; Leino 1992; Melamed 1994; Melamed 2000; Möller 1994). One study was a three-arm trial comparing both nedocromil sodium and sodium cromoglycate with placebo (Leino 1992). Of the remaining seven studies, four examined sodium cromoglycate (Azevedo 1991; Davies 1993; Hechanova 1984; James 2003), and two examined nedocromil sodium (Melamed 2000; Möller 1994). Three studies also included a third treatment arm: either levocabastine, in Azevedo 1991 and Davies 1993, or azelastine (James 2003). One study was reported in two publications (Melamed 1994; Melamed 2000); Melamed 2000 reported combined results from two separate studies, one of which had been previously reported in Melamed 1994.

The number of participants randomised (or analysed if numbers randomised not known) to nedocromil sodium/sodium cromoglycate and placebo were 21 versus 21 (Azevedo 1991), 32 versus 32 (Davies 1993), 20 versus 20 (Hechanova 1984), 50 versus 49 (James 2003), 61 versus 64 (Leino 1992), 43 versus 43 (Melamed 1994), 94 versus 95 (Melamed 2000), and 77 versus 72 (Möller 1994), respectively.

Duration of treatment was two weeks in one study (James 2003), four weeks in five studies, and eight weeks in one study (Melamed 1994; Melamed 2000).

Primary outcomes

Although all eight studies reported at least one of the four symptoms prespecified as primary outcomes (itching, irritation, watering eyes, or photophobia), there was considerable variation in how these outcomes were reported (Table 1).

Six studies collected data on itching. Three of these studies collected participant-reported itching scores using a 0-4 scale, however as no studies also reported a standard deviation, we considered it not possible to perform formal meta-analysis. All studies did report less itching for the active treatment group compared with placebo. Hechanova 1984 presented mean participant-reported scores of 0.96 (sodium cromoglycate) versus 2.10 (placebo) at 14 days, with even larger differences in favour of the active treatment at 21 and 28 days. Melamed 1994 reported scores (it was unclear whether this was the mean or median) of 1.14 (nedocromil sodium) versus 1.48 (placebo) at 14 days. Melamed 2000 reported only change scores from baseline to the peak pollen period and found a statistically significant difference in favour of nedocromil sodium. Möller 1994 reported mean scores of 0.9 (nedocromil sodium) versus 1.4 (placebo) during peak pollen challenge in a graph. Leino 1992 reported that both the sodium cromoglycate and nedocromil sodium groups had statistically significantly less itching than the placebo group, but presented no data. James 2003 only reported itching as part of an overall symptom score plus

the proportion of participants with improvement in itching in each group between days 0 and 3. Although Davies 1993 collected daily data on itch from participant diaries, this publication did not specifically report any data.

Three studies presented data for participant-reported ocular irritation (including grittiness and foreign body sensation), but overall the evidence for the effectiveness of the active treatment was limited. At 14 days, Hechanova 1984 reported mean grittiness scores of 0.58 (sodium cromoglycate) versus 1.31 (placebo) using a 0-4 scale, a difference that was considered statistically significant. Azevedo 1991 presented daily mean irritation using a 0-4 scale in a graph: at 14 days, mean irritation was around 1 for both groups. Between-group differences for the median area under the curve were also similar when the entire 28-day follow-up period was considered. In the study by Möller 1994, mean grittiness (estimated from a graph) during peak pollen challenge was 0.2 (nedocromil sodium) versus 0.5 (placebo). Leino 1992 reported statistically significant differences in favour of both treatment groups versus placebo when considering the change from baseline to four weeks, but not at one week; once again, no actual data were presented. James 2003 only reported foreign body sensation in terms of improvement from day 0 to day 3. Davies 1993 also collected information on ocular irritation, but again did not report this information directly.

Data on watering eyes was not commonly reported. Using a 0-4 scale, Melamed 1994 reported scores of 0.71 (nedocromil sodium) versus 1.08 (placebo) at 14 days, but it was unclear whether these were mean values. In the later publication by this group (Melamed 2000), both treatment groups in fact experienced an increase in tearing during the peak pollen period. Using a 0-4 scale, Möller 1994 reported mean watering scores of 0.4 (nedocromil sodium) versus 0.7 (placebo) during peak pollen challenge. Azevedo 1991 only reported changes in tearing over 28 days, and again James 2003 reported improvement in tearing from baseline to day 3. Davies 1993 and Leino 1992 did not report any information relating to this outcome, even though this had been collected using participant diaries.

Six studies were known to have collected data on photophobia, but there was no clear evidence available concerning this outcome. Two studies did not report any information on this outcome (Davies 1993; Leino 1992), and two studies reported change scores (Azevedo 1991; James 2003). Hechanova 1984 reported a statistically significant difference in favour of sodium cromoglycate, but only at the 14-day time point. The remaining study reported mean scores of around 0.3 on a 0-4 scale for both the nedocromil sodium and placebo groups (Möller 1994).

Overall symptom score

Five articles representing four studies reported some kind of global evaluation or overall symptom score (Table 1). Melamed 2000 reported a participant-reported composite symptom score during

the peak pollen period, and similar separate results for one of the two constituent substudies were also reported (Melamed 1994). James 2003 reported both investigator- and participant-reported composite symptom scores up to 14 days. Leino 1992 collected similar composite scores from investigators and participants but only reported that there were no statistically significant differences. Davies 1993 reported the participants' global evaluation of treatment efficacy using a 4-point scale. Although all available results from these studies tended to favour the active treatment, no standard deviations were reported and no formal meta-analysis was conducted.

No serious adverse events were observed. Overall, placebo, nedocromil sodium, and sodium cromoglycate were well tolerated.

Summary

There was some evidence from individual trials that nedocromil sodium or sodium cromoglycate is more effective than placebo in improving ocular symptoms. However, it was not possible to perform formal meta-analyses for this comparison due to variation in how outcomes were reported and the lack of suitable data (especially standard deviations (SDs)).

Safety

Table 1. Nedocromil sodium or sodium cromoglycate versus placebo. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Nedocromil sodium or sodium cromoglycate	Placebo	Number of participants randomised (n) and comments
Davies 1993	Participants' global evaluation of treatment efficacy (4-point scale)	28 days	Cromoglycate: Excellent/good: 68% Moderate/poor: 32%	Excellent/good: 63% Moderate/poor: 37%	n = 95 Individual categories may be estimated from graph, and mean score could be calculated if considered appropriate
James 2003	Investigator's composite symptom score (itching, tearing, conjunctival redness) (range 0-9), participant's composite symptom score (itching, redness, tearing) (range 0-9)	14 days	Cromoglycate: Investigator: Mean 2.2 Participant: Mean 1.8	Investigator: Mean 2.9 Participant: Mean 2.8	n = 144 Estimated from graph (no SD). Days 3 and 7 also available (investigator-reported); days 1-14 available (participant-reported)
Leino 1992	Overall assessment by participants and investigators (4-point scale)	28 days	Cromoglycate and nedocromil: No data presented	No data presented	n = 195 No significant differences between the 3 groups (except for subgroup analysis by centre)
Melamed 1994	Participant-reported composite symptom score: itchy	14 days	Nedocromil: Mean 3.8	Mean 5.1	n = 86 No SD. Estimated from graph in Fig-

(Continued)

	eyes, burning eyes, tearing eyes, overall eye condition (range 0-16)				ure 2 of the original report. Slightly unclear which symptoms contributed to this score
Melamed 2000	Participant-reported composite symptom score: itchy eyes, burning eyes, tearing eyes, overall eye condition (range 0-16)	“peak pollen period”	Nedocromil: Mean 3.95	Mean 4.92	n = 189 No SD. The time point varied by included study. One of the two studies is Melamed 1994, above

Footnotes

SD: standard deviation

2. Azelastine versus placebo

The search strategy identified nine eligible studies comparing the antihistamine azelastine against placebo (Canonica 2003; Giede-Tuch 1998; James 2003; Lenhard 1997; Nazarov 2003; Petzold 2002; Sabbah 1998). Petzold 2002 summarised data from three different RCTs, which we analysed as individual studies. Three studies reported a third treatment arm, comparing azelastine against levocabastine, in Canonica 2003 and Sabbah 1998, and azelastine against sodium cromoglycate (James 2003). All studies were available as full-text papers, except for Petzold 2002, which was available only as an abstract. Drug concentration was the same in all studies (azelastine 0.05%), however two studies compared two different concentrations (azelastine 0.05% and 0.025%) (Giede-Tuch 1998; Lenhard 1997). Azelastine was administered twice a day.

The number of participants randomised (or analysed if numbers randomised not known) to azelastine and placebo were 57 versus 56 (Canonica 2003), 99 versus 52 (Giede-Tuch 1998), 45 versus 49 (James 2003), 92 versus 94 (Lenhard 1997), 58 versus 58 (Nazarov 2003), 160 versus 80 (Petzold 2002), 99 versus 46 (Petzold 2002), 49 versus 29 (Petzold 2002), and 51 versus 30 (Sabbah 1998), respectively.

Duration of treatment was six weeks in three studies (Canonica 2003; Nazarov 2003; Sabbah 1998), two or four weeks in three trials reported by Petzold 2002, and two weeks in three other studies (Giede-Tuch 1998; James 2003; Lenhard 1997).

Primary outcomes

Although all nine studies reported at least one of the four pre-specified primary outcomes (itching, irritation, watering eyes, or

photophobia), there were some variations in how these outcomes were reported across the included studies.

Data on itching were collected as mean scores in five studies using a 0-3 scale, where higher scores represented worse itching, but as SD was not reported, meta-analysis was not feasible (Giede-Tuch 1998; James 2003; Lenhard 1997; Nazarov 2003; Sabbah 1998). All studies reported less itching with azelastine compared to placebo, but in some studies it was not possible to confirm whether the results were statistically significant (Lenhard 1997). Giede-Tuch 1998 presented participant-reported mean scores of 0.75 (azelastine 0.05%) and 0.90 (azelastine 0.025%) versus 1.15 (placebo) at 14 days. SD was not available. Petzold 2002 summarised data from three RCTs assessing investigators' score on severity of itching; one of these three trials (study ID 3021), with mean scores of 0.53 (azelastine) versus 1.39 (placebo), had statistically significant results in favour of the active treatment. The differences reported in the other two RCTs were not statistically significant, that is study ID 3062 reported mean scores 0.73 (azelastine) and 0.76 (placebo), and study ID 3034 reported mean scores 0.76 (azelastine) and 1.10 (placebo).

Four studies presented data for watering eyes or tearing using a 0-3 scale or sum scores of symptoms (Giede-Tuch 1998; James 2003; Lenhard 1997; Sabbah 1998). Giede-Tuch 1998 presented participant-reported tearing as mean scores (data estimated from graph) of 0.45 and 0.35 (azelastine 0.05% and 0.025%, respectively) versus 0.55 (placebo) at 14 days. No SD was reported. Sabbah 1998 used response rates (calculated from participants' diaries), and Lenhard 1997 reported participants' tearing scores using the same 0-3 scale as for itching.

Data on ocular irritation (described as foreign body sensation) and photophobia were not frequently reported. In James 2003, there was improvement in both study and control groups, although the results were not statistically significant. Some studies reported ocu-

lar irritation and photophobia as part of a composite investigator-reported sum score, for example Sabbah 1998, in which foreign body sensation was part of an 8-symptom sum score at days 3 and 14.

Overall symptom scores

One study, Sabbah 1998, reported itching as composite sum score and response rates on three eye symptoms (itching, tearing, and conjunctival redness), calculated from participants' diaries. Two studies reported participant composite sum score based on two symptoms (itching and redness) using sum mean score (0-6 scale) (Canonica 2003; Nazarov 2003), however neither of these studies reported SD. Nazarov 2003 reported sum score of 1.9 (azelastine) versus 3.0 (placebo), estimated from a published graph at day 14 and the differences consistently increased in favour of the active treatment with longer follow-up, until day 42. These results were consistent with investigators' assessment reported as statistically significant. Canonica 2003 reported investigators' sum scores on itching and redness with mean scores and SD of 1.8 (1.4) (azelas-

tine) versus 3.1 (1.5) (placebo), with change at day 7 statistically significant.

Safety outcomes

No serious adverse events or changes in vital signs were reported during the treatment. Overall, azelastine and placebo were well tolerated by participants across all the studies; few cases withdrew from azelastine or placebo groups due to insufficient tolerability (burning sensation or bitter taste, or both).

Summary

There was some evidence from individual studies that azelastine improved some symptoms more than placebo. However, it was not possible to perform formal meta-analyses for this treatment comparison due to variations of outcomes reported and the lack of suitable data.

Table 2. Azelastine versus placebo. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Azelastine	Placebo	Number of participants randomised (n) and comment
Canonica 2003	Investigator's assessment of change in clinical sum score (itching, redness) (range 0-6)	7 days	Investigator: day 7 Mean sum scores (SD) 1.8 (1.4) Participants: Mean score (itching and conjunctiva redness) 1.65 at day 14	Investigator: day 7 Mean sum scores (SD) 3.1 (1.5) Participants: Mean score (itching and c. redness) 3.85 at day 14	n = 139 Main variable: Time course of sum score of main eye symptoms at day 7 Secondary outcomes: symptoms day 21, 42
Giede-Tuch 1998	Participants' (responders) rate sum score (itching, lacrimation, redness) decreased by at least 3 score points between day 0-3; Investigators' composite symptom mean score (itching, lacrimation, redness)	14 days	Participants' response rate at day 3 (0.025% and 0.05%): 73% and 82%, respectively Investigator: Mean score of three symptoms (0.05% dose) 7 days 2.4	Participants' response rate: 56% Investigator: Mean score of 3 symptoms, 7 days 3.5	n = 151 Investigators' mean estimated from graph (no SD)

(Continued)

James 2003	Investigator's composite sum symptom score (itching, tearing, conjunctival redness) (range 0-9), participant's composite symptom score (itching, redness, tearing) (range 0-9)	14 days	Investigator: Mean score 2.2 Participant: Mean score 1.9	Investigator: Mean score 2.9 Participant: Mean score 2.8	n = 144 Mean scores estimated from the graphs (no SD). Days 3 and 7 also available (investigator-reported); days 1-14 available (participant-reported)
Lenhard 1997	Investigator's composite sum symptom score (itching, tearing, conjunctival redness) (range 0-9) (Secondary analysis)	7 and 14 days	Investigator: Mean score of 3 symptoms (itching, lacrimation, and c. redness) 2.0 at day 14	Investigator: Mean score of 3 symptoms (itching, lacrimation, and c. redness) 2.1 at day 14	n = 278 Mean scores (no SD) due to high variability for itching and marked placebo response, a secondary, more objective analysis was performed Participant-reported itching, c. redness, lacrimation, and swollen eyelids on a 0-3 scale
Nazarov 2003	Investigator's composite sum symptom score (itching, conjunctival redness) on day 7	7 days	Investigator: sum score 1.9 day 14 (estimated from graph)	Investigator: sum score 3.0 day 14 (estimated from graph)	n = 116 Secondary variables: outcomes measured days 21, 42 Participant-reported symptoms: used to corroborate clinical assessments
Petzold 2002 (ID 3021)	Investigator's assessment of severity score for both (itching, redness) and separated for each symptom (itching, redness) (range 0-3)	3 and 14 days	Investigator: Mean score day 14 0.53	Investigator: Mean score day 14 1.39	ID 3021: n = 78 ID 3062: n = 145 ID 3034: n = 240 Treatment duration of 14 days (2 studies) or 28 days (1 study)
Petzold 2002 (ID 3062)			Investigator: Mean score day 14 0.73	Investigator: Mean score day 14 0.76	
Petzold 2002 (ID 3034)			Investigator: Mean score day 14 0.76	Investigator: Mean score day 14 1.10	
Sabbah 1998	Investigator's responders rate based on decrease of at	3 and 14 days	Investigator: Mean sum scores 1.85 (no SD) at day 14	Investigator: Mean sum scores 2.45 (no SD) at day 14	n = 113 Investigators' mean estimated from graph

(Continued)

	<p>least 3 points in the sum symptom score (itching, conjunctival redness, lacrimation) (range 0-9) between day 0 to 3; participant's composite sum symptom score (itching, conjunctival redness, lacrimation) (range 0-9)</p>		<p>Investigator: Difference in the mean sum score (itching, conjunctival redness, and lacrimation) -5.4 (2.3) at day 14</p>	<p>Investigator: Difference in the mean sum score (itching, conjunctival redness, and lacrimation) -3.4 (3.0) at day 14</p>	<p>(no SD) Participant-reported data as response rate (n, %) for composite sum symptom score (itching, conjunctival redness, and lacrimation) at days 3 and 14</p>
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Footnotes

SD: standard deviation

3. Levocabastine versus placebo

The search strategy identified five eligible studies comparing the antihistamine levocabastine versus placebo (Azevedo 1991; Canonica 2003; Davies 1993; Graue 1994; Sabbah 1998). Four of these studies included a third arm comparing levocabastine against azelastine, in Canonica 2003 and Sabbah 1998, and levocabastine against nedocromil sodium or sodium cromoglycate (Azevedo 1991; Davies 1993). Drug concentration was the same in all the studies, 0.05% or 0.5 mg/ml instilled two to four times a day, but Canonica 2003 did not specify dose or concentrations.

The number of participants randomised (or analysed if numbers randomised not known) to levocabastine and placebo were 18 versus 21 (Azevedo 1991), 26 versus 52 (Cannonica 2003), 31 versus 32 (Davies 1993), 20 versus 20 (Graue 1994), and 32 versus 30 (Sabbah 1998), respectively.

Duration of treatment was variable: six weeks in two studies (Cannonica 2003; Sabbah 1998), four weeks in two studies (Azevedo 1991; Davies 1993), and one week in one study (Graue 1994).

Primary outcomes

Data on itching were reported in four studies with some variations in how this symptom was reported. One study did not report itching as a main eye symptom but reported the other three primary outcomes (Azevedo 1991). Grass pollen counts were counted daily during the treatment period of some studies (Azevedo 1991; Davies 1993). Graue 1994 reported only the percentages in various severity categories using graphs. There were no statistically significant differences between the levocabastine and placebo groups (P = 0.45) for improvement of itching at 7 days.

Azevedo 1991 collected data on participant-reported ocular irrita-

tion as percentage of days with absence of ocular irritation. Sixty-three percent had symptom-free days in the levocabastine group versus 44% after placebo (P < 0.06). Itching was also reported in a graph using median area under the curve: 14% (levocabastine) and 29% (placebo) (data estimated from graph). Graue 1994 reported ocular irritation (foreign body sensation), but there was no statistically significant difference between groups (P = 0.178).

Azevedo 1991 reported tearing as the percentage of days free of tearing, reporting larger differences during peak pollen days, with tearing absent in 88% (levocabastine) of the days under treatment versus 58% (placebo) (P = 0.01). Graue 1994 also reported statistically significant results with improvement of tearing scores (P = 0.006).

Data on photophobia were collected and reported as individual symptom or as composite sum score. Graue 1994 reported statistically significant results (P = 0.06) in favour of levocabastine compared with placebo.

Overall symptom scores

Some studies reported sum scores including itching (Cannonica 2003; Sabbah 1998). Cannonica 2003 used the mean score for itching and conjunctival redness (0-6 scale) with mean scores 1.5 (levocabastine) versus 3.85 (placebo) (estimated from a graph), but SD was not given. Davies 1993 reported percentages of global efficacy of treatment with significant participant-reported grading in which participants considered treatment as excellent or good in 87% (levocabastine) against 63% (placebo) (P = 0.05); similar results were observed from investigators' assessed overall score of efficacy of treatment (91% levocabastine versus 68% placebo).

Safety outcomes

Overall, levocabastine and placebo were well tolerated by participants across the studies with no reports of serious adverse events. No difference was reported in the incidence of adverse events between the active and placebo groups, however one study reported higher incidence of (mild) adverse events in the placebo group compared to the levocabastine group (Graue 1994).

There was some evidence from individual studies that levocabastine-treated participants had better outcomes than those who received placebo. However, it was not possible to perform formal meta-analyses for this comparison due to variations of outcomes reported and lack of suitable data.

Summary

Table 3. Levocabastine versus placebo. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Levocabastine	Placebo	Number of participants randomised (n) and comment
Azevedo 1991	Investigator's assessment sum score (ocular irritation, c. redness, photophobia, tearing, swollen eyelids, and conjunctival oedema) (range 0-3); participant-reported same symptoms using VAS scale	14 and 28 days	Investigators: Mean severity scores for each symptom (irritation, tearing, and photophobia) (data not reported, only change score from baseline) Participants: median area under the curve (%) 14%	Investigators: Mean severity scores for each symptom (irritation, tearing, and photophobia) (data not reported, only change score from baseline) Participants: median area under the curve (%) 29%	n = 63 Investigators' and participants' global assessment of efficacy at the end of treatment
Canonica 2003	Investigator's assessment of change in clinical sum score (itching, redness) (range 0-6)	7 days	Investigator: day 7 Mean sum scores (SD) 2.2 (1.2) Participants: Mean score 1.5 at day 14 (itching and conjunctival redness)	Investigator: day 7 Mean sum scores (SD) 3.1 (1.5) Participants: Mean score 3.85 at day 14 (itching and conjunctival redness)	n = 139 Main variable: Time course of sum score of main eye symptoms at day 7 Participants' mean score (no SD) estimated from graphs Secondary outcomes: symptoms day 21, 42
Davies 1993	Area under the curve for participant-reported symptoms (VAS score 0-100, 0 = no symptoms, 100 = extremely severe symptoms); Investigator's symptoms score (ocular irritation, c. redness, itching, photophobia, tear-	14 days	Participants: VAS ≤ 10 symptom-free 37% (P < 0.01); VAS ≤ 50 symptom-free 40% (P < 0.1)	Participants: VAS ≤ 10 symptom-free 4% (P < 0.01); VAS ≤ 50 symptom-free 13%	n = 95 VAS percentages (estimated from graph) Investigator's symptoms severity 0-3 scale (0 = absent, 3 = severe)

(Continued)

	ing, lid and conjunctival oedema) (range 0-3)				
Graue 1994	Investigator-reported symptoms (absent, mild, moderate, or severe scale) Participant-reported symptoms (VAS scale, 0 = no symptoms and 10 = worst possible symptoms)	7 days	Symptoms improvement: 58% first 2 hours 88% day 1 Percentage at day 7 to be estimated from graphs	Symptoms improvement: 33.5% first 2 hours 61% day 1 Percentage at day 7 to be estimated from graphs	n = 40 Unclear whether graph is participant or investigator assessment Percentages of improved symptoms reported (no means and SD)
Sabbah 1998	Investigator's responders rate based on decrease of at least 3 points in the sum symptom score (itching, conjunctival redness, lacrimation) (range 0-9) between day 0 to 3; participant's composite sum symptom score (itching, conjunctival redness, lacrimation) (range 0-9)	3 and 14 days	Investigator: Mean sum scores 1.25 (no SD) at day 14 Investigator: Difference in the mean sum score (itching, conjunctival redness, and lacrimation) -5.4 (2.3) at day 14	Investigator: Mean sum scores 2.45 (no SD) at day 14 Investigator: Difference in the mean sum score (itching, conjunctival redness, and tearing) -3.4 (3.0) at day 14	n = 113 Investigators' mean estimated from graph (no SD) Participant-reported data as response rate (n, %) for composite sum symptom score (itching, conjunctival redness, and tearing) at days 3 and 14

Footnotes

SD: standard deviation
VAS: visual analogue scale

versus 16 (Avunduk 2005), 20 versus 20 (Höffling-Lima 2001), 46 versus 46 (Sarker 2011), and 20 versus 20 (Varguez-Rodriguez 2009), respectively.

4. Olopatadine versus ketotifen

The search identified four eligible studies comparing the antihistamines olopatadine and ketotifen (Avunduk 2005; Höffling-Lima 2001; Sarker 2011; Varguez-Rodriguez 2009). One study was a three-arm trial comparing both olopatadine and ketotifen with placebo (Avunduk 2005). Drug concentration was the same in all studies (olopatadine 0.1% and ketotifen 0.025%), except Höffling-Lima 2001, which used ketotifen 0.05%. Duration of treatment was four weeks in three studies, Avunduk 2005, Höffling-Lima 2001, and Varguez-Rodriguez 2009, and two weeks in one study (Sarker 2011).

In all studies the sample size was relatively small. The number of participants randomised to olopatadine and ketotifen were 16

Primary outcomes

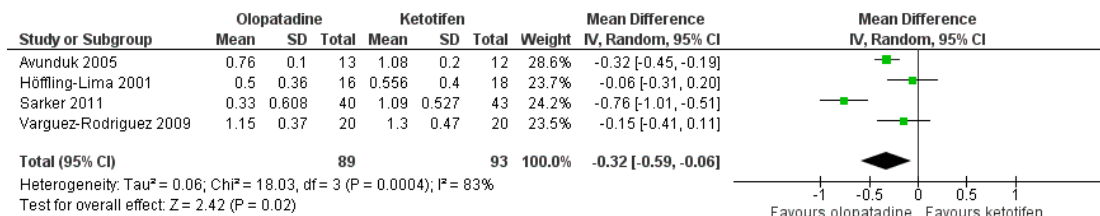
Although four studies reported at least two of the four symptoms prespecified as primary outcomes (itching and tearing), there was some variation in how these outcomes were reported.

All four studies collected data on participant-reported itching using a 0-3 scale. Two studies reported mean and SD values (Avunduk 2005; Sarker 2011).

Two studies did not find any differences between olopatadine and ketotifen in itching reporting (Avunduk 2005; Höffling-Lima 2001), while two studies found a greater reduction in itching with olopatadine than with ketotifen after two weeks of treatment (Sarker 2011; Varguez-Rodriguez 2009). Sarker 2011 reported

two-week mean scores of 1.09 (SD 0.53) with ketotifen and 0.33 (SD 0.60) with olopatadine. A random-effects meta-analysis of these four studies showed evidence of a statistically significant difference in favour of olopatadine in the reduction of itching at 14 days (mean difference (MD) -0.32, 95% confidence interval (CI) -0.59 to -0.06) (Figure 5). However, there was high statistical heterogeneity ($I^2 = 83\%$).

Figure 5. Forest plot of comparison: I Olopatadine versus ketotifen, outcome: I.1 Itching at 14 days (0-3 scale)

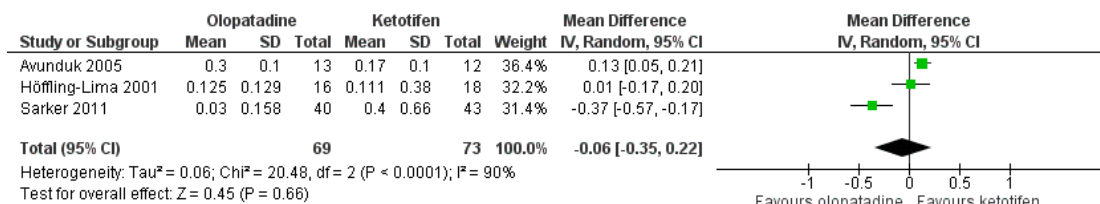


Two studies presented data for participant-reported ocular irritation (described in both studies as burning) (Höfling-Lima 2001; Varguez-Rodriguez 2009). Höfling-Lima 2001 did not find any differences between groups, while Varguez-Rodriguez 2009 reported a statistically significant greater reduction in burning in the olopatadine group ($P < 0.05$).

All four studies reported data on tearing. Only one study reported differences between groups after two weeks (Sarker 2011), with the group treated with olopatadine having less tearing (mean 0.03, SD

0.16) than the one treated with ketotifen (mean 0.40, SD 0.66). Three studies assessed tearing at 14 days (Avunduk 2005; Höfling-Lima 2001; Sarker 2011). A random-effects meta-analysis of these studies found no evidence of a difference between olopatadine and ketotifen (MD -0.06, 95% CI -0.35 to 0.22). There was no evidence of a difference in the reduction of tearing scores at 14 days between the 2 groups (Figure 6). Once again, there was high statistical heterogeneity between the studies ($I^2 = 90\%$).

Figure 6. Forest plot of comparison: I Olopatadine versus ketotifen, outcome: I.2 Tearing at 14 days (0-3 scale)



We knew of two studies that collected data on photophobia (Höfling-Lima 2001; Sarker 2011). There were no differences between groups.

Safety outcomes

No serious adverse events were reported in the four papers. Three

studies did not report any side effects (Avunduk 2005; Höfling-Lima 2001; Varguez-Rodriguez 2009). In one study, ketotifen was associated with a mild stinging sensation of short duration (less than 30 minutes) in 13 out of 43 participants; no participants

treated with olopatadine reported such discomfort (Sarker 2011).

Overall summary

There was some evidence from individual trials that olopatadine may be more effective than ketotifen in improving some ocular symptoms such as itching.

Both drugs are safe.

Table 4. Olopatadine versus ketotifen. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Olopatadine	Ketotifen	Number of participants randomised (n) and comment
Avunduk 2005	Participant-reported (itching and tearing; range 0-3) and ocular signs (redness, eyelid swelling, chemosis; range 0-3)	15 days	Itching: Mean (SD) 0.76 (0.1) Tearing: Mean (SD) 0.30 (0.1)	Itching: Mean (SD) 1.08 (0.2) Tearing: Mean (SD) 0.17 (0.1)	n = 39 SD estimated from graph
Höffling-Lima 2001	Participant-reported (itching, burning, watering, discharge, photophobia; range 0-3) Signs, investigator examination (range 0-3)	14 days	Itching: Mean 0.50 Watering: Mean 0.15 Burning: Mean 0.18 Photophobia: Mean 0.00	Itching: Mean 0.55 Watering: Mean 0.11 Burning: Mean 0.11 Photophobia: Mean 0.05	n = 40
Sarker 2011	Participant-reported (itching, watering, photophobia; range 0-3) (redness; range 0-3)	14 days	Itching: Mean (SD): 0.33 (0.60) Watering: Mean (SD): 0.03 (0.15) Photophobia: Mean (SD): 0.05 (0.22)	Itching: Mean (SD): 1.09 (0.52) Watering: Mean (SD): 0.40 (0.66) Photophobia: Mean (SD): 0.26 (0.44)	n = 92
Varguez-Rodriguez 2009	Participant-reported (itching, watering, burning; range 0-3) Signs, investigator examination (redness, chemosis; range: present or ab-	14 days	Data not extractable. Only P value available.	Data not extractable. Only P value available.	n = 40 Olopatadine induced greater reduction of burning than ketotifen at 2 weeks. There was no difference in itching and tearing at 2 weeks

(Continued)

sent)

Footnotes

SD: standard deviation

5. Emedastine versus levocabastine

We identified three randomised studies that compared the antihistamines emedastine and levocabastine (or levocabastine) (Secchi 2000a; Secchi 2000b; Verin 2001). There was substantial overlap in the authorship of these three studies, and they had some similarities in study design, outcome measures, and the time points for clinic visits (days 3, 7, 14, 30, and 42). One study, Secchi 2000a, was done entirely in a paediatric population (range 4 to 16 years); the other two studies, although including mainly adults, recruited participants from a wide range of age groups (range 4 to 76 years). The same treatment dose (0.05% twice a day) was used in each study, and treatment lasted for six weeks.

The number of participants randomised to emedastine and levocabastine was 20 versus 22 (Secchi 2000a), 97 versus 105 (Secchi 2000b), and 97 versus 105 (Verin 2001), respectively.

Primary outcomes

All three studies evaluated itching. Secchi 2000a evaluated itching up to 42 days using a 0-9 scale. At 14 days, mean scores of 0.7 (emedastine) and 2.5 (levocabastine) were obtained (no SD, estimated from graph). Maximum itching scores from participant diaries were also reported as always lower for emedastine for this study, but no actual data were reported. Secchi 2000b also collected itching data at visits and using participant diaries but did not report any specific data in this article. The authors stated that emedastine was statistically significantly better than levocabastine at 4 of the 5 time points evaluated. Using graphs, Verin 2001 reported mean itching and maximum itching (participant diaries)

up to 42 days using a 0-4 scale. At 14 days, mean scores were 1.5 (emedastine) versus 2.3 (levocabastine). For this study, participant diary scores for maximum itching were 2.0 (emedastine) versus 2.7 (levocabastine).

None of the studies directly evaluated the other primary outcomes of this review (irritation, watering eyes, and photophobia).

Overall symptom score

Secchi 2000a measured the physician's overall impression score at each visit. At 14 days this was 1.1 (emedastine) versus 1.9 (levocabastine) (no SD, estimated from graph). Secchi 2000b measured the physician's overall assessment at the same time points, stating that emedastine was statistically significantly better than levocabastine at all 5 time points, but presented no data.

Conclusion

There was only limited information on our primary outcomes, as these three papers focused on chemosis and eyelid swelling. The results suggested a benefit in favour of emedastine compared with levocabastine for itching and overall symptom relief, but despite some similarities in study design between the three papers, it was not possible to conduct formal meta-analysis, either because the studies presented no data or because they provided no SDs. It was not possible to investigate irritation, watering eyes, or photophobia as no data were available.

Table 5. Emedastine versus levocabastine. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Emedastine	levocabastine	Number of participants randomised (n) and comment
Secchi 2000a	Mean physician's impression score (0-4 scale)	14 days	1.1	1.9	n = 42 Estimated from graph, no standard deviation
Secchi 2000b	Mean physician's impression score (0-4 scale)	14 days	Not reported	Not reported	n = 202 "statistically significant" differences

6. Nedocromil sodium/sodium cromoglycate versus levocabastine

We identified two randomised studies comparing the mast cell stabiliser sodium cromoglycate with the antihistamine levocabastine (levocabastine) (Azevedo 1991; Davies 1993). Duration of treatment was four weeks in both studies, and both studies also included a placebo arm. We identified no studies comparing nedocromil sodium versus levocabastine.

The number of participants randomised (or analysed if numbers randomised not known) to sodium cromoglycate and levocabastine were 21 versus 18 in Azevedo 1991 and 32 versus 31 in Davies 1993, respectively.

Primary outcomes

Neither study presented data on itching. Davies 1993 collected daily data on itching from participant diaries, but specifically reported no data.

Azevedo 1991 presented a graph showing daily mean irritation using a 0-4 scale: at 14 days mean itching was around 1.0 for sodium cromoglycate and 0.8 for levocabastine, and there were statistically significant between-group differences in the median area under the curve for the entire 28-day follow-up period in favour of levocabastine. Davies 1993 collected information on ocular irritation but again did not report this information directly. Azevedo 1991 reported that the median area under the curve for tearing was similar in each group. Although the study provided no data, Davies 1993 reported that tearing was statistically significantly lower for levocabastine.

For photophobia, Azevedo 1991 reported similar change scores

from baseline for the two groups. Davies 1993 did not report any information on this outcome.

Overall symptom score

Davies 1993 collected the participant and investigator global evaluation of treatment efficacy using a 4-point scale (Table 6). Azevedo 1991 reported the investigator assessment using a similar scale. In each study there was a similar statistically significant difference in the proportion rated as excellent or good by the investigator in favour of levocabastine: Azevedo 1991 (nedocromil sodium: 67%, levocabastine: 89%); Davies 1993 (nedocromil sodium: 68%, levocabastine: 91%) (Table 6).

Davies 1993 also reported that participants treated with levocabastine were statistically significantly more likely to be “virtually symptom-free”.

Overall summary

We identified only two randomised studies for this comparison. Although these studies collected data for most of the primary outcomes of this review, the data were either not reported or not given in a format suitable for meta-analysis. Both studies reported a difference of over 20% in favour of levocabastine in those rating the overall treatment efficacy as good or excellent. Although some statistically significant differences were reported suggesting increased efficacy of levocabastine compared with nedocromil sodium, this finding needs to be treated with caution.

Table 6. Nedocromil sodium/sodium cromoglycate versus levocabastine. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Nedocromil sodium / sodium cromoglycate	Levocabastine	Number of participants randomised (n) and comment
Azevedo 1991	Investigator's global assessment	28 days	Excellent/good response: 67% (14/21)	Excellent/good response: 89% (16/18)	n = 63 Response at last available assessment. Percentage of symptom-free days also reported
Davies 1993	Participants' and investigator's global evaluation of treatment efficacy (4-point scale)	28 days	Participant evaluation: Excellent/good: 68%; Investigator evaluation: Excellent/good: 68%	Participant evaluation: Excellent/good: 87%; Investigator evaluation: Excellent/good: 91%	n = 95 Individual categories (participant evaluation) can be estimated from graph, and mean score could be calculated if considered appropriate

7. Azelastine versus levocabastine

We identified two randomised studies comparing the antihistamines azelastine and levocabastine (Canonica 2003; Sabbah 1998). Both studies also included a placebo group. The duration of treatment was six weeks in Canonica 2003 and two weeks in Sabbah 1998.

The number of participants randomised (or analysed if numbers randomised not known) to azelastine and levocabastine were 57 versus 26 in Canonica 2003 and 51 versus 32 in Sabbah 1998, respectively.

Primary outcomes

In both studies itching was assessed by both investigators and participants (using diaries), but no results were directly reported. Canonica 2003 reported only a composite score of itching and redness. Sabbah 1998 only reported itching as part of two composite scores comprising three and eight eye symptoms. Sabbah 1998 only reported the other primary outcomes as part of composite symptom scores. Tearing (tearing) was reported in both composite scores. Foreign body sensation and photophobia were included as part of the eight-item score.

Overall symptom score

The composite symptom scores reported by the two studies were different. Canonica 2003 used the sum of itching and redness (range 0-6). At 14 days the mean scores from the participant diaries (estimated from a graph) were around 1.6 (azelastine) versus 1.4 (levocabastine). The investigator-reported composite scores at 7 and 21 days were also similar in each group. The participant-reported scores were only reported in terms of the proportion of participants improving.

Sabbah 1998 used two composite scores: the first comprised three symptoms (itching, conjunctival redness, and tearing; range 0-9); the second added five additional symptoms (swollen eyelids, foreign body sensation, photophobia, soreness, and discharge/eyelids sticking together). At day 13 scores for the 3-item score (estimated from a graph) were around 1.8 (azelastine) versus 1.5 (levocabastine). The study presented additional results for the numbers responding by day three of treatment.

Overall summary

Overall, there was no clear evidence of differences between azelastine and levocabastine in either study (Table 7).

Table 7. Azelastine versus levocabastine. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Azelastine	Levocabastine	Number of participants randomised (n) and comment
Canonica 2003	Itching and redness (range 0-6)	14 days	1.6	1.4	n = 139 Estimated from a graph (no SD)
Sabbah 1998	Itching, conjunctival redness, and tearing (range 0-9)	13 days	1.8	1.5	n = 113 Estimated from a graph (no SD). A further composite score comprising eight symptoms was also presented

Footnotes

SD: standard deviation

8. Olopatadine versus placebo or other control

The search identified two eligible studies comparing the antihistamine olopatadine versus placebo, in Avunduk 2005, or other control, in Lanier 2001.

The Avunduk 2005 study was a three-arm trial comparing both olopatadine and ketotifen with placebo (Avunduk 2005). Duration of treatment was four weeks, but two-week data were available. The sample size was relatively small, with 16 participants randomised to olopatadine and 17 participants randomised to ar-

tificial tears (placebo). Mean and SD data were available.

Lanier 2001 compared the effect of topical olopatadine in people undergoing treatment with an oral antihistamine (loratadine). Duration of treatment was one week. A total of 94 participants (49 versus 45) were randomised but not masked. No SD data were available.

Primary outcomes

Avunduk 2005 reported two of the four symptoms prespecified as primary outcomes (participant-reported itching and watering eyes). Data on mean and SD were available. Olopatadine had

statistically significantly less severe itching (mean 0.76, SD 0.1) and tearing (mean 0.30, SD 0.1) than the placebo group (mean itching 1.85, SD 0.3; mean tearing 1.07, SD 0.2) after two weeks of treatment.

Lanier 2001 included participant-reported itching on a 4-point scale (from 1 to 4). After one week, reported itching was less in the group treated with olopatadine (mean 2.21 versus 2.74, $P = 0.044$).

Safety outcomes

There were no adverse events or side effects associated with olopatadine in either study.

Overall summary

There was evidence from two small trials that olopatadine may be effective in improving some ocular symptoms.

Table 8. Olopatadine versus control. Outcome definition, time points, and summary of results

Article	Outcome definition	Time point	Olopatadine drops	Placebo	Number of participants randomised (n) and comment
Avunduk 2005	Participant-reported (itching and tearing; range 0-3) and ocular signs (redness, eyelid swelling, chemosis; range 0-3)	30 days	Itching: Mean (SD) 0.76 (0.1) Tearing: Mean (SD) 0.30 (0.1)	Itching: Mean (SD) 1.85 (0.3) Tearing: Mean (SD) 1.07 (0.2)	n = 39 SD estimated from graph
Lanier 2001	Participant-reported itching (range 1-4) and clinician exam of redness	7 days	Itching: Mean 2.21 Redness: Mean 1.36	Itching: Mean 2.74 Redness: Mean 1.55	n = 94 No SD available

Footnotes

SD: standard deviation

9. Nedocromil sodium or sodium cromoglycate versus azelastine

We identified a single randomised study comparing the mast cell stabiliser sodium cromoglycate with the antihistamine azelastine (James 2003). This study also included a placebo arm. The original numbers randomised was not clear, but the analysis included 50 (sodium cromoglycate) versus 45 (azelastine) participants. Duration of treatment was two weeks.

Primary outcomes

Unfortunately this study did not directly report any of our pre-specified outcomes at follow-up, although it did report results for itching, foreign body sensation, and photophobia in terms of the improvement rates from baseline to day three. Improvement rates were higher for azelastine for all eight symptoms examined.

Overall symptom score

James 2003 also reported 2 composite symptom scores comprising 3 main symptoms (itching, redness, and tearing): an investigator-reported score and composite symptom scores from participant diaries up to 14 days. Scores were similar in each group. The study reported response rates to treatment by day 3 (a decrease of at least 3 points in the composite symptom score) to be 39 out of 47 (83%) for sodium cromoglycate and 35 out of 41 (85%) for azelastine.

Overall summary

The evidence for this comparison came from a single study, and although the study reported results for ocular symptoms, the results were not in the format prespecified for this review. Overall, there was no clear evidence of a difference between the two treatments.

10. Levocabastine versus antazoline plus tetryzoline

We identified a single randomised study comparing the antihistamine levocabastine (n = 35) versus the antihistamine antazoline plus tetryzoline (a derivative of imidazoline) (n = 34) (Wertheimer 1997).

Primary outcomes

The study reported no data on the four primary outcomes of this review (itching, irritation, watering eyes, and photophobia). However, all four symptoms were included as part of an overall symptom score.

Overall symptom score

The study presented a total symptom score (range 0-24) comprising four subjective (itching, foreign body feeling, tearing, and photophobia) and four objective (hyperaemia, follicles, chemosis, and swollen eyelid) symptoms. At day 15, mean scores were 2.0 for levocabastine and 3.0 for antazoline plus tetryzoline. These results were estimated from a graph, and no SDs were presented. There were no significant differences between the groups at days 4 and 15, although the antazoline/tetryzoline group was favoured when assessed 30 minutes after application.

Overall summary

Evidence for this comparison comes from one relatively small randomised study. There were no clear differences between the groups.

11. Ketotifen versus placebo

The search identified one eligible study comparing the antihistamine ketotifen versus placebo (Avunduk 2005). This study was a three-arm trial comparing both olopatadine and ketotifen with placebo.

Duration of treatment was four weeks.

The sample size was relatively small, with 16 participants randomised to ketotifen and 17 participants randomised to artificial tears (placebo).

Primary outcomes

The study reported two of the four symptoms prespecified as primary outcomes (itching and watering eyes). Data on mean and SD were available.

Ketotifen had statistically significantly less severe itching (mean 1.08, SD 0.2) and tearing (mean 0.17, SD 0.1) than the placebo group (mean itching 1.85, SD 0.3; mean tearing 1.07, SD 0.2) after two weeks of treatment.

Safety outcomes

No adverse events or side effects were reported.

Overall summary

There is evidence from a small individual trial that ketotifen may be more effective than placebo in improving some ocular symptoms.

12. Olopatadine versus nedocromil sodium

The search identified one eligible study comparing the antihistamine olopatadine with the mast cell stabiliser nedocromil sodium in a parallel-group trial where one group used olopatadine and placebo and the other nedocromil sodium (Katelaris 2002). Duration of treatment was six weeks. The sample size was relatively large, with 91 participants randomised to the olopatadine group and 94 participants randomised to the nedocromil sodium group; analysis was only performed on 82 and 87 participants, respectively. In the olopatadine group, participants were treated with olopatadine 0.1% (in the morning and evening) and placebo (at noon and afternoon); in the nedocromil sodium group, participants were treated with nedocromil sodium 2%. The study provided no information on how many participants were randomised within each treatment group.

Primary outcomes

The study reported one of the four symptoms prespecified as primary outcomes (itching). Participants recorded itching and redness on a scale of 0 to 9 (9 being more severe). After two weeks, mean itching score of participants taking olopatadine was 1.9, compared to 2.5 in those taking nedocromil sodium ($P < 0.05$). However, there was no significant statistical difference in redness scores at 14 days.

The study reported that the likelihood of a day without eye redness and itching was 1.6 times greater on most days in participants treated with olopatadine compared to those treated with nedocromil sodium. This difference was recorded as constant over time ($P < 0.001$).

Safety outcomes

In the olopatadine group, four cases of treatment-related adverse events were recorded (ocular discharge, stinging, and blurred vision). Two participants experienced dry nose and taste perversion. In the nedocromil sodium group, five participants experienced treatment-related ocular adverse events (dry eye, stinging, itching, and tearing).

Overall summary

This study suggested there is some evidence supporting olopatadine as a more effective agent than nedocromil sodium in improving some ocular symptoms of allergic conjunctivitis.

13. Ketotifen versus levocabastine

The search identified one eligible study comparing the antihistamine ketotifen (0.025%) with the antihistamine levocabastine (0.05%) (Kidd 2003). The study was a three-arm trial comparing both ketotifen and levocabastine to placebo in a large, multicentre trial.

Duration of treatment was four weeks.

The sample size was large, with a total of 519 participants being randomised to 1 of 3 groups: 172 participants were randomised to ketotifen and 174 participants were randomised to levocabastine.

Primary outcome

This study reported only two of the prespecified primary outcome measures (participant-reported itching and watering). Participant-recorded data was only available for days one to four. Participants scored itching on a 5-point scale (4 being most severe) and watering on a 4-point scale (3 being severe).

At day 4, mean score of itching for ketotifen was the lowest (1.4), compared with 1.7 for both levocabastine and placebo ($P < 0.05$). The mean score of watery eyes for participants treated with ketotifen was the lowest (0.75) after 4 days compared with 1.2 for participants treated with both placebo and levocabastine ($P < 0.05$).

Safety outcomes

The study recorded four serious adverse events. In the placebo group, two participants experienced persistent photophobia and conjunctivitis with corneal ulcer. In the ketotifen group, two participants experienced spontaneous pneumothorax and abdominal pain, which was probably not due to the drug.

Overall summary

This study showed some evidence that ketotifen may be superior in alleviating some ocular symptoms of allergic conjunctivitis.

14. Combined levocabastine hydrochloride and pemirolast potassium versus levocabastine hydrochloride alone

The search identified one eligible study comparing a combination of the antihistamine levocabastine hydrochloride (0.025%) ophthalmic suspension and pemirolast potassium solution with levocabastine (0.025%) alone (Fujishima 2008).

Duration of treatment was one week.

The sample size was small, with 15 participants randomised to the combined treatment group and 17 participants randomised to single-agent treatment.

Primary outcomes

The study reported two of the four symptoms prespecified as primary outcomes (itching, tearing). The study collected data by participant diary and visual analogue scale quantified by the participant. Data on mean and SD was available.

After one week of treatment there was a reduction in mean (SD) itching in both combined-treatment (-4.6 (2.3)) and single-treatment (-2.8 (2.8)) groups. There was no significant difference in alleviation of symptoms between the groups ($P = 0.079$).

Only 11 participants in the single-treatment group and 8 participants in the combined-treatment group had symptoms of tearing. The degree of reduction in tearing (mean (SD)) was more statistically significantly in the combined-treatment group compared with the single-treatment group (-4.0 (2.6) vs. -1.5 (0.9), $P = 0.008$).

Safety outcomes

No adverse reactions were observed in either group.

Overall summary

There was some evidence from a single study with a small sample size that levocabastine in combination with pemirolast potassium may be more effective in eliminating some symptoms when compared with levocabastine alone.

15. Levocabastine versus mequitazine

The search identified one randomised study as an abstract (full text was not available) comparing the antihistamine levocabastine (0.05%) versus the antihistamine mequitazine (0.05%) eyedrops in a parallel-group trial (Trinquand 1999).

The study did not report the numbers randomised to each group, but reported that a total of 357 participants were randomised.

Duration of treatment was four weeks (doses were twice a day for the first week and two or three times daily for three weeks).

Primary outcomes

Ocular symptoms were assessed by the investigator and participants (diary cards). Investigator assessment was good or excellent in 72% of mequitazine-treated participants and 70% of levocabastine-treated participants.

Safety outcomes

The study reported fewer adverse events in the mequitazine group ($P < 0.05$).

The overall safety of mequitazine (0.05%) was significantly better than levocabastine.

Overall summary

There was some evidence from a single study with a large sample size ($n = 357$) that mequitazine is as effective as levocabastine in teenagers and adult patients.

16. Bepotastine besilate versus olopatadine

The search identified one eligible study comparing the antihistamine bepotastine besilate ophthalmic solution (BBOS) against the antihistamine olopatadine, an investigator-masked, single-centre, cross-over study that randomised 30 participants (McCabe 2012). Participants were not masked.

The duration of treatment was two weeks, followed by a seven-day washout period. After the washout period, participants were crossed-over to the alternative treatment for two additional weeks. Although a paired t-test was used to analyse this study, no paired data were presented. Some results were obtainable from graphs, but these were assumed to represent mean (SD) values after pooling data from both the first and second periods of the study. This meant that unit of analysis issues would have been a potential concern when combining with participant-randomised studies. However, no meta-analyses were possible for this comparison.

Primary outcomes

Participants assessed ocular itching during the three clinic visits using a 5-point Likert scale (1 = lowest relief, 5 = highest relief) and used a daily home diary to assess ocular itch relief twice a day (morning and evening) over two weeks of treatment.

Rather different mean (SD) results were obtained for the morning and evening (morning: BBOS 4.04 (0.12) versus olopatadine 4.10 (0.15); evening: BBOS 4.04 (0.12) versus olopatadine 3.90 (0.12)). These results appeared to be for both periods of the cross-over study combined.

Safety outcomes

The study reported no serious adverse events. About 10% of the participants treated with BBOS 1.5% reported a mild, temporary adverse taste after instillation.

Overall symptom score

The study reported the mean (SD) rating of each treatment's ability to relieve all ocular-related allergy symptoms (1 = lowest relief,

3 = highest relief) for morning and evening separately (morning: BBOS 2.30 (0.1) versus olopatadine 2.25 (0.15); evening: BBOS 2.30 (0.1) versus olopatadine 2.15 (0.15)). From the participants' diary responses, BBOS 1.5% was significantly more effective at relieving morning and evening ocular allergy symptoms ($P = 0.032$ and $P < 0.0001$, respectively) compared to olopatadine hydrochloride 0.2%.

Overall summary

There was insufficient evidence to compare the efficacy of BBOS against olopatadine, as we found only one study with a small sample size that was judged to be at high risk of bias, as participants were not masked to treatment allocation.

17. BBOS versus placebo

The search identified a single multicentre, randomised trial comparing the antihistamine BBOS against placebo (Carr 2013). The study randomised participants to receive either BBOS ($n = 123$) or placebo ($n = 122$) twice a day for a period of two weeks.

Primary outcomes

Participants assessed ocular itching twice a day using a 4-point scale (0 = absent, 3 = severe) that was part of the participant outcomes assessed from a Rhinconjunctivitis Quality of Life Questionnaire. The study reported results as percentages of improvement from baseline (instantaneous -- approximately 15 minutes before scoring, and reflective -- last dosing throughout the 2-week treatment period) of ocular itching scores over the treatment period. The mean change from baseline involved taking the daily average of a two-week period and comparing with the mean scores from a three-day baseline period. For reflective itching, mean improvements were 28.0% for BBOS and 21.1% for placebo. For instantaneous itching, mean improvements were 28.3% for BBOS and 20.3% for placebo.

Safety outcomes

The study reported no serious adverse events. More mild adverse events were reported in the BBOS group (29 participants) than in the placebo group (11 participants). The most frequently reported adverse events in the BBOS group were bitter taste (14 participants in the BBOS group, 1 participant in the placebo group) and instillation site pain.

Overall summary

There was some evidence from a single trial with a large sample size ($n = 245$) that BBOS is more effective at improving ocular itching than placebo. However, full results other than percentage change in itching scores were not available.

DISCUSSION

Summary of main results

We identified 30 trials with 17 different treatment comparisons evaluating the efficacy and safety of topical antihistamines and mast cell stabilisers, either alone or in combination. The following antihistamines and mast cell stabilisers were evaluated in at least one randomised controlled trial: nedocromil sodium or sodium cromoglycate, olopatadine, ketotifen, azelastine, emedastine, levocabastine or levocabastine, combination of antazoline and tetryzoline, combination of levocabastine and pemirolast potassium, and bepotastine besilate. The most common comparison was azelastine versus placebo (nine studies). Unfortunately, formal meta-analysis was only possible for two outcomes in one comparison (olopatadine versus ketotifen, itching and tearing at 14 days), and the results should be interpreted with caution due to the high statistical heterogeneity, both for itching ($I^2 = 83\%$) and tearing ($I^2 = 90\%$). For this comparison, one study differed from the other studies as it favoured olopatadine (Sarker 2011).

The inability to meta-analyse other results was mainly due to the variety of outcome measures reported and the fact that standard deviations were often not provided. Due to the presence of a network of different treatments in this review, this should have been an ideal situation to conduct a network meta-analysis of overall symptom scores, but the heterogeneity in outcome definitions and time points and the lack of standard deviations meant that sadly, this was not feasible. We did not do sensitivity analysis because of the small number of studies involved in the meta-analysis.

There was some evidence to support the ability of topical antihistamines to reduce symptoms and signs of seasonal allergic conjunctivitis when compared with placebo. There were no serious adverse events related to the use of topical antihistamine treatment.

When comparing different types of antihistamines and mast cell stabilisers, there were limited data to inform if some treatments are more effective than others. Results from a meta-analysis suggested that olopatadine may be more effective than ketotifen in relieving itching, although there was high statistical heterogeneity between the two studies. Results from individual studies suggested better outcomes with emedastine, sodium cromoglycate, and ketotifen when compared with levocabastine. A single study suggested a benefit of olopatadine over sodium cromoglycate.

Overall completeness and applicability of evidence

There are no long-term studies on the efficacy of topical antihistamines for the treatment of seasonal allergic conjunctivitis, and

little evidence to compare antihistamines.

Quality of the evidence

Poor quality of reporting challenged the synthesis of evidence. We observed a large variability in reporting outcomes. The overall quality of the studies and reporting was poor, and most studies had small sample sizes. Trials only evaluated short-term effects, with a range of treatment of one to eight weeks.

Potential biases in the review process

None. This review was performed to Cochrane standards.

Agreements and disagreements with other studies or reviews

We are not aware of any other similar reviews.

AUTHORS' CONCLUSIONS

Implications for practice

Topical antihistamines and mast cell stabilisers reduce symptoms and signs of seasonal allergic conjunctivitis when compared with placebo in the short term. Overall, topical antihistamines and mast cell stabilisers appear to be safe and well tolerated. There is poor evidence to compare efficacy among different antihistamines and mast cell stabilisers.

Implications for research

Methodological research to reach consensus on core outcome measures and how best to quantify them would facilitate research in this area. Large trials comparing the efficacy of different antihistamines and mast cell stabilisers would be required.

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* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Avunduk 2005

Methods	Parallel-group RCT
Participants	49 participants recruited with seasonal allergic conjunctivitis
Interventions	Three treatment arms: ketotifen 0.025% ophthalmic solution; olopatadine hydrochloride 0.1% ophthalmic solution; artificial tear substitute (preservative free). Duration of treatment 30 days, follow-up 30 days (2-month study)
Outcomes	Participant evaluation of ocular symptoms (itching, tearing) Investigator assessment of ocular signs (redness, eyelid swelling, chemosis) Mean scores (using a scale range 0-3; 0 = none, 3 = severe) Time points: at day 0, 15, and 30 of treatment
Country	Turkey
Number randomised, gender (male:female)	39 participants randomised. M:F 20:19
Age mean (SD), median, range	Overall range 18-61 years
Notes	Study conducted from April to May 2004. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated list (p.1394 - study medications): "Eligible patients were randomly assigned, in a 1:1 ratio using a computer-generated list of random numbers..."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported for each intervention group with 20% lost to follow-up (Figure 1 in the trial report), reasons were not given. A similar number of participants were lost to follow-up in each arm
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias

Avunduk 2005 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	Low risk	Masking of participants (p.1394 - study medications): "To maintain masking, all medications had identical packaging, color, consistency, pH, and texture"
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Investigators masked (p.1394 - efficacy assessments): "...ocular signs (redness, eyelid swelling, and chemosis) were graded, using slit-lamp examination and ordinal grading scales, by an investigator masked to treatment assignment."

Azevedo 1991

Methods	Parallel-group RCT
Participants	63 participants recruited with moderate or severe allergic conjunctivitis
Interventions	Three treatment arms: levocabastine 0.5 mg/ml eye drops; sodium cromoglycate 20 mg/ml eye drops; placebo eye drops. Duration of treatment 28 days
Outcomes	Participant assessment (ocular irritation, redness, photophobia, tearing, swollen eyelids, conjunctival oedema using a 4-point scale; 0 = none, 3 = severe) Investigator assessment (ocular irritation, redness, photophobia, tearing, swollen eyelids, conjunctival oedema, using a 4-point scale; 0 = none, 3 = severe) Time points: at baseline and day 14 and 28 of treatment
Country	Portugal
Number randomised, gender (male:female)	63 participants randomised, 60 participants analysed. M:F 24:36
Age mean (SD), median, range	Median (range): levocabastine 27 years (13-55); sodium cromoglycate 26 years (9-46); placebo 34 years (12-51)
Notes	Not reported when study was conducted. Source of funding not reported. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated list (p.691 - treatment and methods): "Patients were randomly allocated (on the basis of a computer-generated list) to three groups..."
Allocation concealment (selection bias)	Unclear risk	Method used to conceal the allocation sequence was not described

Azevedo 1991 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and the reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.691 - treatment and methods): "The trial featured a doubled-blind, parallel groups design"
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.691 -treatment and methods): "The trial featured a doubled-blind, parallel groups design"

Canonica 2003

Methods	Parallel-group RCT
Participants	139 participants recruited with moderate to severe perennial allergic conjunctivitis
Interventions	Three treatment arms: azelastine 0.05% eye drops; placebo eye drops; levocabastine eye drops. Duration of treatment 6 weeks
Outcomes	Investigator assessment (itching and redness). Change in clinical sum score (itching and redness using a scale range 0-6) Participant assessment of ocular symptoms (daily diaries using a 4-point scale; 0 = none, 3 = severe symptoms) Time points: at baseline and day 7, 21, and 42 of treatment
Country	France, Italy, Spain, Russia, and United Kingdom
Number randomised, gender (male:female)	139 participants randomised. M:F 65:74
Age mean (SD), median, range	Mean (SD): azelastine group 34 years (13.7); placebo group 36 years (13.0); levocabastine 34 years (13.2)
Notes	Study conducted from 2 December 1998 to 16 June 1999. Source of funding: University of Genoa, Italy and Clinical and Biometrical Development, VIATRIS GmbH & Co. KG. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
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Canonica 2003 (Continued)

Random sequence generation (selection bias)	Low risk	Randomisation code (p.323 - drugs): "A predefined randomisation code was used to assign azelastine, placebo or levocabastine treatment to qualified patients in an unbalanced fashion (ratio 2:2:1)."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and the reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.323 - drugs): "Both azelastine eye drops and the matching vehicle containing placebo were provided by VI-ATRIS GmbH in identical packaging."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.323 - drugs): "Both azelastine eye drops and the matching vehicle containing placebo were provided by VI-ATRIS GmbH in identical packaging."

Carr 2013

Methods	Parallel-group RCT
Participants	245 participants recruited with allergic conjunctivitis
Interventions	Two treatment arms: bepotastine besilate (BBOS) eye drops 1.5%; placebo eye drops. Duration of treatment 2 weeks
Outcomes	Participant assessment of ocular itching (instantaneous and reflective) mean change scores (using diaries and a 4-point scale; 0 = none, 3 = severe) Time points: at baseline, day 7 and 14 after treatment
Country	United States
Number randomised, gender (male:female)	245 participants randomised. M:F 92:153
Age mean (SD), median, range	Mean (SD) BBOS group 38.7 (13.80) range 12-85; placebo group 41.6 (15.03) range 12-72
Notes	Source of funding: ISTA Pharmacologicals. The main investigators declared they had no financial interests in the outcome of the clinical trial. Carr and Ratner also declared that they were speakers and consultants for ISTA and Meda Pharmaceuticals

<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated randomisation list (p.249 - clinical trial design): "...were assigned to receive BBOS 1.5% or placebo in a 1:1 ratio as determined by a computer-generated randomisation list."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10 % lost to follow-up, and the reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Masking of participants (p.249 - clinical trial design): "Both test agents were provided in identical packaging and manufactured by Bausch & Lomb (Tampa, FL)."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Study conducted from July to October 2010. Investigators masking (p.249 - clinical trial design): "Subjects were instructed to instill the double-masked test agent..."

Davies 1993

Methods	Parallel-group RCT
Participants	95 participants recruited with allergic conjunctivitis
Interventions	Three treatment arms: levocabastine 0.5 mg/ml; sodium cromoglycate 20 mg/ml; placebo eye drops. Duration of treatment 28 days
Outcomes	Investigators' and participants' assessment of ocular symptoms (4-point qualitative scale: excellent, good, moderate, poor) Time points: at baseline, day 14 and 28 after treatment
Country	United Kingdom
Number randomised, gender (male:female)	95 participants randomised. M:F 40:49
Age mean (SD), median, range	Mean (range) levocabastine 30 years (7-66); sodium cromoglycate 31 years (10-69); placebo 30 years (13-61)

Davies 1993 (Continued)

Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	High risk	Percentages given (numerators/denominators unclear)
Selective reporting (reporting bias)	High risk	P values given for selected outcomes
Other bias	Unclear risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.520 - methods): "...95 patients participated in this double-blind, parallel group trial."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.520 - methods): "...95 patients participated in this double-blind, parallel group trial."

Fujishima 2008

Methods	Parallel-group RCT
Participants	32 participants recruited with seasonal and perennial allergic conjunctivitis
Interventions	Two treatment arms: levocabastine; levocabastine and pemirolast potassium. Duration of treatment 1 week
Outcomes	Participants used a diary to quantify ocular symptoms (itching, foreign body sensation, tearing, eye discharge, and hyperaemia) using visual analogue scale Investigators used a 4-point scale (0 = none, 3 = severe) for quantifying ocular signs: conjunctival hyperaemia, conjunctival oedema, eyelid oedema, conjunctival papilla formation, and superficial punctate keratopathy Time points: at baseline and 1 week after treatment
Country	Japan

Fujishima 2008 (Continued)

Number randomised, gender (male:female)	32 participants randomised, 30 participants analysed. M:F 9:23
Age mean (SD), median, range	Mean (range): levocabastine 50.9 years (12-81); levocabastine and pemirolast potassium 40.5 years (7-73)
Notes	Not reported when study was conducted. Source of funding unclear. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Sealed-envelope technique (p.233 - test agents): "The patients were randomly allocated to one of two groups by a sealed-enveloped technique..."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data of all participants randomised was analysed and reported
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	The study did not describe details on masking of participants on the intervention allocations
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	The study did not describe details on masking of personnel/ investigators on the intervention allocations

Giede-Tuch 1998

Methods	Parallel-group RCT
Participants	151 participants with seasonal allergic conjunctivitis or rhinoconjunctivitis for at least 1 year
Interventions	Three treatment arms: azelastine 0.025%; azelastine 0.05%; placebo. Duration of treatment 14 days
Outcomes	Participant-assessed itching, tearing, and redness. Sum score (itching, tearing, redness) and rates of decreased scores by at least 3 score points between day 0-3 using a 4-point scale (0 = none, 3 = severe) were used

Giede-Tuch 1998 (Continued)

	Investigator-assessed itching, tearing, and redness. A composite sum symptom mean score (itching, tearing, redness) using a 4-point scale (0 = none, 3 = severe) was analysed Time points: at baseline, day 3, 7, and 14 after treatment
Country	Germany
Number randomised, gender (male:female)	151 participants randomised, 129 participants analysed. M:F 66:85
Age mean (SD), median, range	Mean (SD) azelastine 0.025% 35.4 years (11.4); azelastine 0.05% 35.2 years (10.7); placebo 35.9 years (11.5)
Notes	Study conducted from March to September 1994. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 15 % lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.858 - study design): "The investigation was performed as a double-blind, randomized, placebo-controlled study..."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.858 - study design): "The investigation was performed as a double-blind, randomized, placebo-controlled study..."

Graue 1994

Methods	Parallel-group RCT
Participants	Participants from 5 to 20 years old with seasonal conjunctivitis
Interventions	Two treatment arms: levocabastine 0.5 mg/ml; placebo. Duration of treatment 7 days

Graue 1994 (Continued)

Outcomes	Participants' assessment of ocular symptoms (visual analogue scale; 0 = none, 10 = worse possible) Investigator-evaluated symptoms (absent, mild, moderate, and severe) Time points: at baseline and day 7 after treatment
Country	Mexico
Number randomised, gender (male:female)	40 participants randomised. M:F 30:10
Age mean (SD), median, range	Age mean: levocabastine 9.1 years; placebo 10.1 years
Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described (p.37 - material and methods): "patients were divided randomly into two groups..."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.37 - material and methods): "The study design is double-blind, prospective and comparative..." "both drugs were provided by Janssen Pharmaceutical in similar plastic bottles of 4ml each, labeled as levocabastine (including the ones that contained placebo) and with an identification code number..."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.37 - material and methods): "The study design is double-blind, prospective and comparative..." "both drugs were provided by Janssen Pharmaceutical in similar plastic bottles of 4ml each, labeled as levocabastine (including the ones that contained placebo) and with an identification code number..."

Hechanova 1984

Methods	Parallel-group RCT
Participants	40 participants with allergic conjunctivitis
Interventions	Two treatment arms: sodium cromoglycate; placebo. Duration of treatment 4 weeks
Outcomes	Participants' and clinicians' assessment of ocular symptoms and opinion of efficacy of treatment: specifically they assessed itching, soreness, redness, grittiness, photophobia using a 5-point scale (0 = none, 5 = very severe) Time points: at 1, 2, and 4 weeks after treatment Results of photophobia were not reported
Country	Philippines
Number randomised, gender (male:female)	40 participants randomised. M:F 20:20
Age mean (SD), median, range	Mean (range) 35 years (5-71)
Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described (p.60 - material and methods): "...with patients randomly allocated to receive..."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	High risk	High rates of withdrawals
Selective reporting (reporting bias)	High risk	Only selected results were presented
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.60 - material and methods): "The trial was a double-blind group comparison..."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.60 -material and methods): "The trial was a double-blind group comparison..."

Höfling-Lima 2001

Methods	Parallel-group RCT
Participants	40 participants with allergic conjunctivitis
Interventions	Two treatment arms: ketotifen 0.05%; olopatadine 0.1%. Duration of treatment 30 days
Outcomes	Participants' assessment of severity of ocular symptoms: itching, burning, watering, discharge, photophobia (4-point scale; 0 = none, 3 = severe) Investigator assessment of ocular signs Time points: at days 1, 2, 7, 14, and 30 after treatment
Country	Brazil
Number randomised, gender (male:female)	40 participants randomised, 34 participants analysed. M:F 15:19
Age mean (SD), median, range	Mean (range) 27 years (5-73)
Notes	Study conducted from 1 February to 30 June 1999. Source of funding not stated. The authors declared no commercial interest in the drugs evaluated and did not receive any direct or indirect benefit or financial support for this study

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation (p.416 - methods): "A masked and randomised clinical study was conducted..." "the introduction of treatment was randomized beforehand..."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with 15% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Masking of participants (p.416 - methods): "A masked and randomised clinical study was conducted..." "... and bottles with topical medications were masked by ophthalmos laboratory."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Masking of investigator (p.417 - methods): "only the medical monitors delivered the drugs to patients..."

James 2003

Methods	Parallel-group RCT
Participants	144 participants with seasonal allergic conjunctivitis
Interventions	Three treatment arms: azelastine; sodium cromoglycate; placebo. Duration of treatment 14 days
Outcomes	Participants' assessment of symptoms (composite score): itching, redness, tearing, foreign body sensation, photophobia, soreness, discharge (range 0-9) using participant's diary during treatment (14 days) Investigator assessment of itching, tearing, conjunctival redness (range 0-9, composite sum score) Time points: at day 3, 7, and 14 after treatment
Country	Germany and United Kingdom
Number randomised, gender (male:female)	144 participants randomised, 136 participants analysed. M:F 51:93
Age mean (SD), median, range	Mean (range): azelastine 37.1 years (16-65); sodium cromoglycate 35.5 years (18-65); placebo 35.8 years (18-64)
Notes	Study conducted from April to September 1995. Source of funding: ASTA Medica AG (now VIATRIS GmbH & Co. KG). Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Partial double-masked (p.314 - study design): "partial double-blind, parallel-group... The appearance and application regimen of placebo eye drops was identical to azelastine and the study was double-blind in this respect."

James 2003 (Continued)

Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Partial double-masked (p.314 - study design): "However, the study was open in respect of sodium cromoglycate-treated patients where the bottle size and dosage regimen identified this treatment to the investigator."
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Katellaris 2002

Methods	Parallel-group RCT
Participants	188 participants with seasonal allergic conjunctivitis
Interventions	Two treatment arms: olopatadine hydrochloride 0.1%; cromolyn sodium 2% (nedocromil sodium). Duration of treatment 6 weeks
Outcomes	Investigator's assessment of ocular signs and symptoms: mean ocular symptom scores of itching and redness and physicians' impression scale scores, per protocol data (5-point scale; 0 = none, 4 = very frequent) Participants' assessment of itching and redness using diary (10-point scale; 0 = none, 9 = severe) Time points: at days 1 to 14, and 30 to 42 after treatment
Country	6 European countries and Australia
Number randomised, gender (male:female)	188 participants randomised, 185 participants analysed. M:F 103:82
Age mean (SD), median, range	Mean (SD): olopatadine 33.0 years (19.3); cromolyn sodium 36.8 years (20.9)
Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated (p.1563 - study procedures): "A computer-generated randomization schedule was used to assign sequential patients to the study treatments in an equal ratio."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Analysis per-protocol participants not intention-to-treat analysis. Less than 10% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of adequate outcome reporting

Katellaris 2002 (Continued)

Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masking (p.1563 - study procedures): "Double-masking was ensured through the use of identical opaque bottles and similar-appearing contents and labeling."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double masking (p.1563 - study procedures): "Double-masking was ensured through the use of identical opaque bottles and similar-appearing contents and labeling."

Kidd 2003

Methods	Parallel-group RCT
Participants	519 participants with seasonal allergic conjunctivitis
Interventions	Three treatment arms: ketotifen fumarate 0.025%; levocabastine 0.05%; placebo. Duration of treatment 4 weeks
Outcomes	Participant assessment (mean scores) of ocular itching (using a 5-point scale; 0 = none, 4 = severe) and watering (scale 0 = none, 3 = severe) within the first 4 days of treatment Investigator assessment of signs (redness, eyelid swelling, chemosis) after treatment (using a 5-point scale; 0 = none, 4 = severe)
Country	Australia
Number randomised, gender (male:female)	519 participants randomised, 348 participants analysed. M:F 269:254
Age mean (SD), median, range	Mean (SD): ketotifen 46.3 years (17.0); levocabastine 49.5 years (17.4); placebo 47.9 years (17.4)
Notes	Not reported when study was conducted. Source of funding Novartis. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	RANCODE software (p.1207 - study design): "Eligible subjects were randomised (RANCODE version 3.6) to one of the three treatment groups..."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described

Kidd 2003 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Not intention-to-treat analysis, no reasons reported per protocol deviations (large numbers). Number of participants randomised (519) does not match the sum of male and female (523)
Selective reporting (reporting bias)	Low risk	Evidence was available of adequate outcome reporting.
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Identical packing (p.1207 - study design): "The packaging of all trial medications was identical in appearance."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Identical packing (p.1207 - study design): "The packaging of all trial medications was identical in appearance."

Lanier 2001

Methods	Parallel-group RCT
Participants	94 participants with seasonal allergic conjunctivitis
Interventions	Two treatment arms: olopatadine and oral loratadine; oral loratadine. Duration of treatment 1 week
Outcomes	Participants' and physicians' assessment of ocular signs and symptoms Mean scores for ocular itching and redness using diaries and a 4-point scale Time points: at baseline, day 3 and 7 after treatment
Country	United States
Number randomised, gender (male:female)	94 participants randomised, 72 participants analysed. M:F 33:61
Age mean (SD), median, range	Mean (range): olopatadine and oral loratadine 39 years (7-74); oral loratadine 37 years (9-74)
Notes	Study conducted from May to November 1998. Source of funding Alcon Pharmaceutical. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described

Lanier 2001 (Continued)

Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Exclusion of 22 participants after randomisation (p.644 - results): "Ten patients were not evaluable for efficacy because they did not meet inclusion/exclusion criteria or had no follow up examination... Twelve patients exited the study prematurely..."
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	High risk	(p.643 - study design): "Because there were no control eye drops, patients were aware of their treatment groups, but clinicians performing the evaluations in the clinic were not."
Blinding of outcome assessment (detection bias) All outcomes	High risk	(p.643 - study design): "Because there were no control eye drops, patients were aware of their treatment groups, but clinicians performing the evaluations in the clinic were not."

Leino 1992

Methods	Parallel-group RCT
Participants	195 participants with seasonal allergic conjunctivitis
Interventions	Three treatment arms: nedocromil sodium 2%; sodium cromoglycate 2%; placebo. Duration of treatment 4 weeks
Outcomes	Overall assessment by participants and investigators using a 4-point scale
Country	Finland
Number randomised, gender (male:female)	195 participants randomised, 185 participants analysed. M:F not stated
Age mean (SD), median, range	Mean: nedocromil 20.8 years; sodium cromoglycate 19.3 years; placebo 19.7 years
Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described

Leino 1992 (Continued)

Allocation concealment (selection bias)	Low risk	(p.930 - material and methods): "After 1-2 weeks the patients were issued their appropriate eye drops by randomized code number..."
Incomplete outcome data (attrition bias) All outcomes	High risk	Withdrawal/exclusion of participants (p.931 - results): "Data were returned from 195 patients, 10 of whom withdrew without taking test treatment and were excluded." "A further 12 patients (5 nedocromil sodium, 3 sodium cromoglycate and 4 placebo) withdrew from the study without completing the 4-week treatment period."
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Identical drugs packaging (p.930 - medication): "All test drugs were supplied in identical 10 ml plastic bottles."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Identical drugs packaging (p.930 - medication): "All test drugs were supplied in identical 10 ml plastic bottles."

Lenhard 1997

Methods	Parallel-group RCT
Participants	278 participants with seasonal allergic conjunctivitis or rhinoconjunctivitis
Interventions	Three treatment arms: azelastine 0.025%; azelastine 0.05%; placebo. Duration of treatment 14 days
Outcomes	Investigator mean score of 3 symptoms (itching, tearing, and redness (secondary analysis)) using a 4-point scale Participants' assessment of symptoms (itching primary variable) using diaries and a 4-point scale Time points: at baseline, at treatment day 7 and 14
Country	France, Italy, Poland, and Slovenia
Number randomised, gender (male:female)	278 participants randomised, 226 participants analysed as per protocol. M:F 114:164
Age mean (SD), median, range	Mean (SD) azelastine 0.025% 31.6 years (10.6); azelastine 0.05% 31.7 years (11.7); placebo 33.9 years (11.9)

Lenhard 1997 (Continued)

Notes	Not reported when study was conducted. Source of funding ASTA Medica AG. Declaration of interest by the authors was not stated	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Of 278 recruited participants (p.24 - results): "226 patients completed the study as planned..." No reasons were given. (p.24 - results): "Twenty-four of the 278 recruited patients discontinued the study prematurely."
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	The study did not describe details on masking of participants on the intervention allocations
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	-The study did not describe details on masking of personnel/ investigators on the intervention allocations

McCabe 2012

Methods	Parallel-group crossed-over RCT
Participants	30 participants recruited with allergic conjunctivitis
Interventions	Two treatment arms: bepotastine besilate 1.5% ophthalmic solution; olopatadine hydrochloride 0.2%
Outcomes	Participants' assessment of ocular itching (using a 5-point Likert scale; 1 = lowest relief, 5 = highest relief) at baseline and 2 other clinic visits but also using daily diary to assess morning and evening ocular itch relief during 2 weeks of treatment
Country	United States
Number randomised, gender (male:female)	30 participants randomised, M:F 10:20

McCabe 2012 (Continued)

Age mean (SD), median, range	Mean (SD) 49.8 (2.76), range 23-75	
Notes	Study conducted in September 2011. Source of funding not stated. No authors have financial or proprietary interest in any material or method mentioned in this work	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomisation list (p.1733 - materials and methods): "The enrolled patients were assigned sequentially according to a computer-generated randomization list..."
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Complete outcome data reported, all the participants randomised were analysed
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	High risk	Participants were not masked to treatment allocation (p.1733 - material and methods): "Each treatment was provided in the packaging originally approved by the Food and Drug Administration..."
Blinding of outcome assessment (detection bias) All outcomes	High risk	Investigator was masked to treatment allocation (p. 1733 - material and methods): "...but the single investigator was masked as to which treatment the patient was currently using."

Melamed 1994

Methods	Parallel-group RCT
Participants	86 participants with seasonal allergic conjunctivitis
Interventions	Two treatment arms: nedocromil sodium; placebo. Duration of treatment 8 weeks
Outcomes	Participants' assessment of symptoms -- sum score itching, burning, tearing, overall eye condition Clinicians' assessment of ocular signs and symptoms (using a scale 0 = none, 4 = very severe) Time points: at baseline and peak pollen period during the 8 weeks of treatment

Melamed 1994 (Continued)

Country	United States
Number randomised, gender (male:female)	86 participants randomised, 85 participants analysed. M:F 49:36
Age mean (SD), median, range	Mean (range) nedocromil sodium 33.1 years (13-60); placebo 31.1 years (13-57)
Notes	Not reported when study was conducted. Source of funding: support in part by Fisons Pharmaceuticals. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked and identical package (p.58 - study design): "The study was designed as a multicenter, randomized, double-masked... Active and placebo drugs were packaged identically and labeled in opaque polyethylene bottles..."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked and identical package (p. 58 - study design): "The study was designed as a multicenter, randomized, double-masked... Active and placebo drugs were packaged identically and labeled in opaque polyethylene bottles..."

Melamed 2000

Methods	Parallel-group RCT
Participants	189 participants with seasonal allergic conjunctivitis
Interventions	Two treatment arms: nedocromil sodium 2%; vehicle. Duration of treatment 8 weeks

Melamed 2000 (Continued)

Outcomes	Participants' assessment of various symptoms (composite score itching, burning, tearing, overall eye condition) and diary cards (using a scale 0 = none, 4 = very severe) to assess individual symptoms at treatment Clinicians' assessment of ocular signs and symptoms, and clinician and participant overall opinions of treatment effectiveness at baseline and peak pollen season Time scale: during the 8 weeks of treatment	
Country	United States	
Number randomised, gender (male:female)	189 participants randomised. M:F 104:85	
Age mean (SD), median, range	Mean (range): nedocromil sodium 33.4 years (12-65); vehicle 31.0 years (13-67)	
Notes	Study conducted in August 1986. Source of funding: in part by Fisons Pharmaceuticals. The authors did not have a financial interest in the drugs discussed within this manuscript	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up with similar percentage of participants lost to follow-up in each arm, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.236 - study design): "The nine week studies used a double-blind, placebo-controlled..." Identical bottles (p.236 - study protocol): "At the end of the baseline week, patients were randomised to receive either one drop of nedocromil sodium 2% or vehicle solution b.i.d in each eye, delivered from identical opaque bottles."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.236 - study design): "The nine week studies used a double-blind, placebo-controlled..." Identical bottles (p.236 - study protocol): "At the end of the baseline week, patients were randomised to receive either one drop of nedocromil sodium 2% or vehicle solution b.i.d in each eye, delivered from identical opaque bottles."

Möller 1994

Methods	Parallel-group RCT
Participants	149 participants with seasonal allergic conjunctivitis
Interventions	Two treatment arms: nedocromil sodium; placebo. Duration of treatment 4 weeks
Outcomes	Investigator's and participants' (or their parents') assessment of severity of symptoms. Mean score of itching, redness, soreness, grittiness, photophobia, and general eye condition (using a 5-point scale; 0 = none, 4 = very severe) Time points: during peak pollen in the 4 weeks of treatment
Country	Sweden
Number randomised, gender (male:female)	149 participants randomised, 146 participants analysed. M:F 92:57
Age mean (SD), median, range	Mean (range): nedocromil sodium 12 years (6-16); placebo 13 years (6-16)
Notes	Not reported when study was conducted. Source of funding: Fisons Pharmaceuticals. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Use of predetermined randomised code (p.885 - patients and methods): "Patients were then allocated, by a predetermined randomised code... Trial supplies were coded at source, by computer-generated numbers randomised in a balanced blocks of four"
Allocation concealment (selection bias)	Low risk	Conceal allocation described (p.885 - patients and methods): "Patients entering the trial were assigned to treatment numbers in sequence. The codes were held in individual sealed envelopes by the investigator, to be returned unopened after the trial."
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.884 - summary): "This was a multicentre, double-blind..." (p.885 - patients and methods): "active or placebo eye drops (both contained... in addition, the placebo eye drops contained

Möller 1994 (Continued)

		0.0005% rivoflavin as a yellow colorant to match the 2% nedocromil sodium content of the active eye drops).”
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.884 - summary): “This was a multicentre, double-blind...” (p.885 - patients and methods): “active or placebo eye drops (both contained... in addition, the placebo eye drops contained 0.0005% rivoflavin as a yellow colorant to match the 2% nedocromil sodium content of the active eye drops).”

Nazarov 2003

Methods	Parallel-group RCT
Participants	116 participants with moderate to severe perennial allergic conjunctivitis
Interventions	Two treatment arms: azelastine 0.015 mg; placebo. Duration of treatment 6 weeks
Outcomes	Participants’ and clinicians’ assessment of ocular symptoms (sum score itching and redness, using a 4-point scale; 0 = none, 3 = severe) Time points: at baseline and day 7, 21, and 42 of treatment
Country	Germany
Number randomised, gender (male:female)	116 participants randomised and analysed. M:F 26:90
Age mean (SD), median, range	Mean (range): azelastine 35.7 years (17-63); placebo 31.7 years (17-59)
Notes	Study conducted from 19 December 1998 to 14 April 1999. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors’ judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting

Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.169 - patients, materials and methods): “The investigation was conducted as a double-blind, randomised...” Both drugs with identical packaging (p.169 - treatment): “Both azelastine eye drops and the matching vehicle containing placebo were provided by ASTA Medica AG in identical packaging.”
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.169 - patients, materials and methods): “The investigation was conducted as a double-blind, randomised...” Both drugs with identical packaging (p.169 - treatment): “Both azelastine eye drops and the matching vehicle containing placebo were provided by ASTA Medica AG in identical packaging.”

Petzold 2002

Methods	Parallel-group RCTs (3 RCTs reported together in abstract)
Participants	Study 3034: 78 participants; Study 3062: 145 participants; Study 3021: 240 participants with allergic conjunctivitis
Interventions	Two treatment arms: azelastine 0.05%; placebo. Duration of treatment 14 or 28 days
Outcomes	Investigators’ assessment of ocular symptoms (itching and redness) using a 4-point scale (0 = none, 3 = severe) Time scale: day 0, 3, and 14 of treatment
Country	Europe
Number randomised, gender (male:female)	Study 3034: 78 participants (number of participants in each arm uncertain) randomised Study 3062: 145 participants (number of participants in each arm uncertain) Study 3021: 240 participants (number of participants in each arm uncertain). Gender not stated (abstract)
Age mean (SD), median, range	Age range 4-12 years
Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors’ judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Only abstract available
Allocation concealment (selection bias)	Unclear risk	Only abstract available

Petzold 2002 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Only abstract available
Selective reporting (reporting bias)	Unclear risk	Only abstract available
Other bias	Unclear risk	Only abstract available
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Only abstract available
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Only abstract available

Sabbah 1998

Methods	Parallel-group RCT
Participants	113 participants with seasonal allergic conjunctivitis/rhinoconjunctivitis
Interventions	Three treatment arms: azelastine; levocabastine; placebo. Duration of treatment 14 days
Outcomes	Participants' (using diaries) and physicians' assessments (sum scores) based on a decrease of the average score > 3 units for three main eye symptoms (ocular itching, tearing, and redness) Time points: at baseline and day 3 and 14 of treatment
Country	France
Number randomised, gender (male:female)	113 participants randomised, 107 participants analysed. M:F 75:38
Age mean (SD), median, range	Mean (SD): azelastine 8.3 years (2.4); levocabastine 8.2 years (2.5); placebo 8.3 years (2.3)
Notes	Not reported when study was conducted. Source of funding Laboratoires ASTA Medica. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described

Sabbah 1998 (Continued)

Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	10% attrition, with similar numbers lost to follow-up between arms
Selective reporting (reporting bias)	Low risk	Primary outcome result was reported according to protocol
Other bias	Low risk	No evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	The study did not describe details on masking of participants on the intervention allocations. Masking was not done with levocabastine due to labelling, but double-masked with other interventions
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	The study did not describe details on masking of personnel/ investigators on the intervention allocations. Masking was not done with levocabastine due to labelling, but double-masked with other interventions

Sarker 2011

Methods	Parallel-group RCT	
Participants	92 participants with allergic conjunctivitis	
Interventions	Two treatment arms: ketotifen fumarate 0.025%; olopatadine hydrochloride 0.1%. Duration of treatment 2 weeks	
Outcomes	Assessment of ocular sign and symptoms (mean scores) for itching, tearing, hyperaemia, photophobia (using a 4-point scale; 0 = none, 3 = severe) Time points: at baseline and 2 weeks treatment period Unclear if outcomes were measured by participants or clinicians, or both	
Country	Bangladesh	
Number randomised, gender (male:female)	92 participants randomised, 83 participants analysed. M:F 36:47	
Age mean (SD), median, range	Mean (SD): ketotifen 28 years (12); olopatadine 28 years (11). Overall range 12-50 years	
Notes	Source of funding not stated. The authors have no relevant affiliations or financial involvement with any organization or entity with a financial interest in or financial conflict with the subject matter or materials discussed in the manuscript	
<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement

Random sequence generation (selection bias)	Low risk	Randomisation list (p.547 - treatment allocation and follow-up) : “Patients who were found to be eligible according to selection criteria were recruited in to one of the treatment groups according to a stratified randomisation list based on age and sex.”
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with less than 10% lost to follow-up, and reasons were given
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.547 - treatment allocation and follow-up): “Study medications were provided in identical containers so that both patients and investigators remained blinded.”
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Study conducted from 1 January to 31 December 2007. Double-masked (p.547 - treatment allocation and follow-up): “Study medications were provided in identical containers so that both patients and investigators remained blinded.”

Secchi 2000a

Methods	Parallel-group RCT
Participants	42 paediatric participants with allergic conjunctivitis
Interventions	Two treatment arms: emedastine 0.05%; levocabastine 0.05%. Duration of treatment 42 days
Outcomes	Investigator assessment of ocular signs (using a scale 0-4) at days 3, 7, 14, 30, 42, and overall progress Participant assessment of ocular symptoms itching and redness (using a visual analogue scale; 0 = none, 9 = severe) Time points: at days 7, 14, 30, 42 of treatment
Country	Italy
Number randomised, gender (male:female)	42 participants randomised, 38 participants analysed. Gender not stated
Age mean (SD), median, range	Age range 4-6 years: 11 participants Age range 7-16 years: 31 participants

Secchi 2000a (Continued)

Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported, with 10% lost to follow-up, reasons were given for one case only
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.42 - abstract, material and methods): "In a randomised, double-masked... Subjects who met all inclusion and exclusion criteria received masked study medication with instructions to instill drops..."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.42 - abstract, material and methods): "In a randomised, double-masked..."

Secchi 2000b

Methods	Parallel-group RCT
Participants	222 participants (including 42 paediatric) with seasonal allergic conjunctivitis
Interventions	Two treatment arms: emedastine 0.05%; levocabastine 0.05%. Duration of treatment 42 days
Outcomes	Investigator assessment of ocular signs (using a scale 0-4) at days 3, 7, 14, 30, 42 Participant assessment of ocular symptoms (using a visual analogue scale; 0 = none, 9 = severe) at days 7, 14, 30, 42
Country	Italy and United States
Number randomised, gender (male:female)	202 participants randomised. M:F 107:114

Secchi 2000b (Continued)

Age mean (SD), median, range	Average age (range): 30 years (4-76)
Notes	Not reported when study was conducted. Source of funding not stated. Declaration of interest by the authors was not stated

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data were nearly fully reported for each intervention group, with less than 20% lost to follow-up (Figure 1 of the trial report). A similar percentage of participants was lost to follow-up in each arm
Selective reporting (reporting bias)	Unclear risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Double-masked (p.49 - material and methods): "In a randomized, double-masked..."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Double-masked (p.49 - material and methods): "In a randomized, double-masked..."

Trinquand 1999

Methods	Parallel-group RCT
Participants	357 participants with perennial or seasonal allergic conjunctivitis
Interventions	Two treatment arms: mequitazine; levocabastine. Duration of treatment 4 weeks
Outcomes	Investigators' assessment of symptoms at day 7, 14, and 28 Participants' assessment of symptoms (diary cards)
Country	Not stated
Number randomised, gender (male:female)	357 participants randomised

Trinquand 1999 (Continued)

Age mean (SD), median, range	Not stated	
Notes	Not reported when study was conducted. Full-text paper not available, only abstract, therefore insufficient information to assess risk of bias. Source of funding and declaration of interest by the authors were not stated	
<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Only abstract available
Allocation concealment (selection bias)	Unclear risk	Only abstract available
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Only abstract available
Selective reporting (reporting bias)	Unclear risk	Only abstract available
Other bias	Unclear risk	Only abstract available
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Only abstract available
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Only abstract available

Varguez-Rodriguez 2009

Methods	Parallel-group RCT
Participants	40 participants (naive-treatment) with seasonal allergic conjunctivitis
Interventions	Two treatment arms: olopatadine 0.1%; ketotifen 0.025%. Duration of treatment 4 weeks
Outcomes	Assessment of ocular symptoms: itching, watering, burning (range 0-3; 0 = none, 3 = severe) and signs: redness, chemosis (absent, present) Unclear if outcomes were measured by participants or clinicians, or both
Country	Mexico
Number randomised, gender (male:female)	40 participants randomised. M:F 23:17

Varguez-Rodriguez 2009 (Continued)

Age mean (SD), median, range	Mean (SD): olopatadine 19.7 years (6.6); ketotifen 21.05 years (8.3)	
Notes	Source of funding not stated. Declaration of interest by the authors was not stated	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Low risk	Complete outcome data reported, all the participants randomised were analysed
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Masking of participants (p.401 - methods): "All the bottles had the same appearance and the patient did not know the treatment received."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Masking of investigators (p.401 - methods): "All the bottles had the same appearance and the patient did not know the treatment received."

Verin 2001

Methods	Parallel-group RCT
Participants	222 participants with seasonal allergic conjunctivitis
Interventions	Two treatment arms: levocabastine; emedastine. Duration of treatment 6 weeks
Outcomes	Participants' assessment of itching and redness using diaries and visual analogue scale (0 = none; 9 = severe) Physician assessment of itching, redness Time points: day 0-14 and day 30-42 of treatment
Country	Europe, South Africa, and Australia
Number randomised, gender (male:female)	222 participants randomised, 202 participants analysed. Gender not stated

Verin 2001 (Continued)

Age mean (SD), median, range	Mean (range) 30 years (4-76)	
Notes	Not reported when study was conducted. Source of funding Alcon Research Ltd. Declaration of interest by the authors was not stated	
<i>Risk of bias</i>		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	The method used to generate the allocation sequence was not described
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Not intention-to-treat analysis
Selective reporting (reporting bias)	Low risk	Evidence was available of proper outcome reporting
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Identical containers (p.692 - methods): "Study medications were provided in identical containers so that both patients and investigators remained masked."
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Identical containers (p.692 - methods): "Study medications were provided in identical containers so that both patients and investigators remained masked."

Wertheimer 1997

Methods	Parallel-group RCT
Participants	69 participants with acute allergic conjunctivitis
Interventions	Two treatment arms: antazoline 0.05% and tetrazoline 0.04%; levocabastine 0.05%. Duration of treatment 2 weeks
Outcomes	The total symptom score (Figure 1 of the trial report) of eight symptoms (each scored 0 = none; 3 = worst symptom). Four subjective and four objective symptoms; unclear which symptoms were assessed by participants or clinicians
Country	Germany
Number randomised, gender (male:female)	69 participants randomised. M:F 35:34

Wertheimer 1997 (Continued)

Age mean (SD), median, range	Mean (SD): antazoline and tetrazyoline 42.4 (15.4); levocabastine 43.1 (14.9)	
Notes	Study conducted from March to August 1995. Source of funding not stated. Declaration of interest by the authors was not stated	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Random sequence generation (p.94): "The randomisation schedule, created at random in blocks of four, assigned patients to the medications and application instructions listed in Table 2"
Allocation concealment (selection bias)	Unclear risk	The method used to conceal the allocation sequence was not described
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No dropouts were mentioned
Selective reporting (reporting bias)	High risk	Only P values that were statistically significant were reported (p. 95, paragraph 3). No results were reported for outcomes that were not statistically significant (p.94, paragraph 8). Numerical results were not reported in the text for any outcomes except for the numbers with blurred vision (p.95, paragraph 4)
Other bias	Low risk	No apparent evidence of other risk of bias
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	(p. 94, paragraph 3): "After 30 minutes the eyes were studied again, without the doctor knowing which drops were administered. The medication and the double-masked application instructions, which prevented early identification of the medication, were handed over to the patient at the end of the visit"
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Masked at 30-minute assessment, later assessments were not masked to participants, as treatments had different dose frequencies (2 per day versus 4 per day). Assessment masked to clinicians unclear

RCT: randomised controlled trial

SD: standard deviation

Characteristics of excluded studies *[ordered by study ID]*

Study	Reason for exclusion
Abelson 2003	Using conjunctival allergen challenge test
Abelson 2004	Using conjunctival allergen challenge test
Artal 2000	Unit of randomisation: eye (participants received different treatments on each eye)
Borazan 2009	Unit of randomisation: eye (participants received different treatments on each eye)
Garay 2001	Non-randomised controlled trial
Higuchi 1979	Non-randomised controlled trial
Kamis 2006	Unit of randomisation: eye (participants received different treatments on each eye)
Leino 1994	Comparing different concentrations and dosage of the same active drug
Leonardi 2004	Non-randomised controlled trial
Longo 1979	Non-randomised controlled trial
Merayo 2003	Non-randomised controlled trial
Möller 1990	Non-randomised controlled trial
Napoli 2005	Non-randomised controlled trial
Pinto 2001	Cost-effectiveness study (original study Verin 2001)
Scadding 1999	Non-randomised controlled trial
Scoper 2007	Non-randomised controlled trial
Torkildsen 2008	Unit of randomisation: eye (participants received different treatments in each eye)

Characteristics of studies awaiting assessment *[ordered by study ID]*

Dharmistha 2013

Methods	No details, unclear if it is a RCT
Participants	Allergic conjunctivitis
Interventions	Topical olopatadine hydrochloride 0.1% versus ketotifen fumarate 0.025%

Dharmistha 2013 (Continued)

Outcomes	Unclear
Notes	None

Jia 2012

Methods	No details, unclear if it is a RCT
Participants	Children with allergic conjunctivitis
Interventions	Azelastine
Outcomes	No details
Notes	In Chinese language

Scandashree 2013

Methods	RCT
Participants	Allergic conjunctivitis
Interventions	Olopatadine hydrochloride 0.2% ophthalmic solution once daily versus sodium cromoglycate 2% ophthalmic solution 4 times a day
Outcomes	Efficacy and tolerability. No further details
Notes	Reported at the 46th Annual Conference of the Indian Pharmacological Society, IPSCON Bangalore India, 2013

RCT: randomised controlled trial

DATA AND ANALYSES

Comparison 1. Olopatadine versus ketotifen

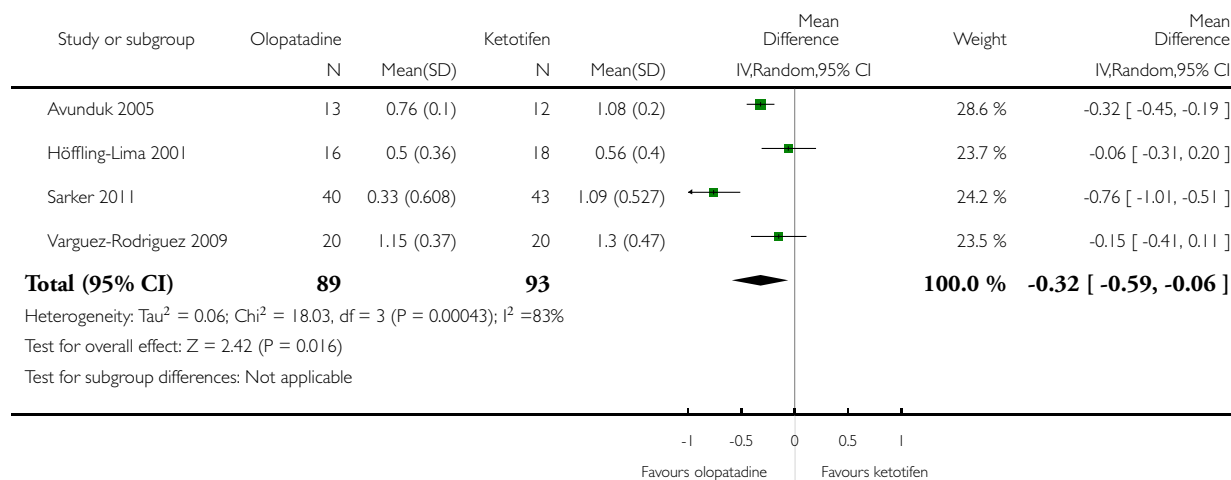
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Itching at 14 days	4	182	Mean Difference (IV, Random, 95% CI)	-0.32 [-0.59, -0.06]
2 Tearing at 14 days	3	142	Mean Difference (IV, Random, 95% CI)	-0.06 [-0.35, 0.22]

Analysis 1.1. Comparison 1 Olopatadine versus ketotifen, Outcome 1 Itching at 14 days.

Review: Topical antihistamines and mast cell stabilisers for treating seasonal and perennial allergic conjunctivitis

Comparison: 1 Olopatadine versus ketotifen

Outcome: 1 Itching at 14 days

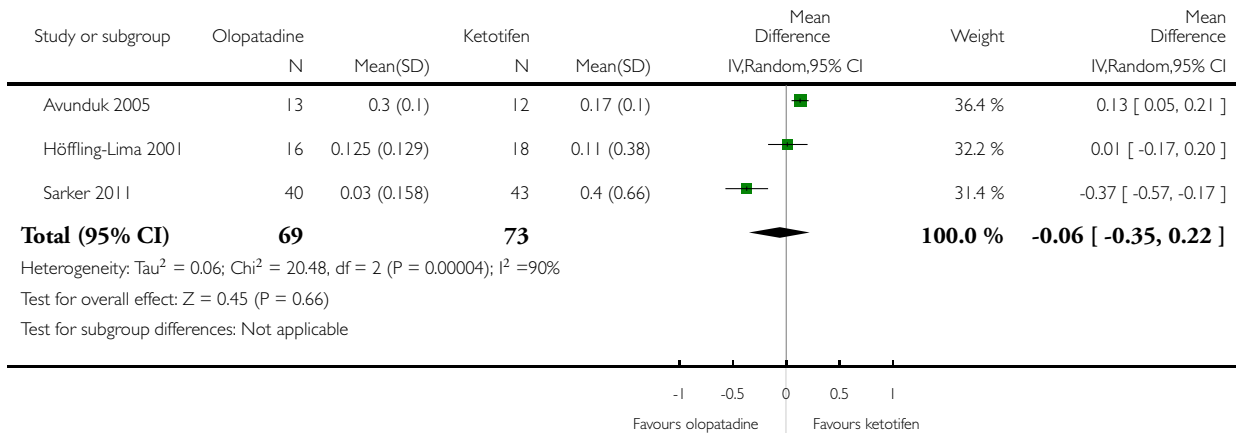


Analysis 1.2. Comparison 1 Olopatadine versus ketotifen, Outcome 2 Tearing at 14 days.

Review: Topical antihistamines and mast cell stabilisers for treating seasonal and perennial allergic conjunctivitis

Comparison: 1 Olopatadine versus ketotifen

Outcome: 2 Tearing at 14 days



ADDITIONAL TABLES

Table 1. Serious adverse events and withdrawals

Study	Serious adverse events	Number withdrawn due to side effects	Treatment arm from which withdrawn
Avunduk 2005	No serious adverse events reported	No participants recorded to have withdrawn	N/A
Azevedo 1991	No serious adverse events reported	1 participant withdrawn	1 from placebo group
Canonica 2003	No serious adverse events reported	2 participants withdrawn	1 from azelastine and 1 from placebo
Carr 2013	1 serious adverse event reported unrelated to the treatment	11 participants withdrawn	4 from bepotastine besilate and 7 from placebo
Davies 1993	No serious adverse events reported	5 participants withdrawn	2 from levocabastine group and 3 from placebo
Fujishima 2008	No serious adverse events reported	No participants recorded to have withdrawn	N/A

Table 1. Serious adverse events and withdrawals (Continued)

Giede-Tuch 1998	No serious adverse events reported	No participants withdrawn due to side effects	N/A
Graue 1994	No serious adverse events reported	2 participants withdrawn	1 from levocabastine and 1 from placebo
Hechanova 1984	No serious adverse events reported	12 participants withdrawn	2 from cromoglycate group and 10 from placebo group
Höffling-Lima 2001	No serious adverse events reported	No participants recorded to have withdrawn	N/A
James 2003	No serious adverse events reported	4 participants withdrawn due to side effects	1 from azelastine, 2 from cromoglycate, and 1 from placebo
Katellaris 2002	No serious adverse events reported	2 participants withdrawn	1 from olopatadine group, 1 from cromolyn group
Kidd 2003	4 serious adverse events: 1) persistent photophobia and 2) conjunctivitis with corneal ulcer	38 discontinued due to side effects	8 from ketotifen, 15 from levocabastine, 15 from placebo
Lanier 2001	No serious adverse events reported	No participants recorded to have withdrawn	N/A
Leino 1992	No serious adverse events reported	4 participants withdrawn	3 from nedocromil and 1 from cromoglycate
Lenhard 1997	No serious adverse events reported	4 participants withdrawn	2 from azelastine group and 2 from placebo group
McCabe 2012	No serious adverse events reported	No participants recorded to have withdrawn	N/A
Melamed 1994	No serious adverse events reported	No participants recorded to have withdrawn due to side effects	N/A
Melamed 2000	No serious adverse events reported	No participants recorded to have withdrawn due to side effects	N/A
Moller 1994	No serious adverse events reported	No participants recorded to have withdrawn due to side effects	N/A
Nazarov 2003	No serious adverse events reported	1 participant withdrawn due to side effects	1 participant from azelastine group

Table 1. Serious adverse events and withdrawals (Continued)

Petzold a,b,c 2002	No serious adverse events reported	No participants recorded to have withdrawn due to side effects	N/A
Sabbah 1998	No serious adverse events reported	1 participant withdrawn	1 participant from azelastine group
Sarker 2011	No serious adverse events reported	No mention of participants withdrawing due to side effects	N/A
Secchi a 2000	No serious adverse events reported	1 participant discontinued due to side effects	1 participant from emedastine
Secchi b 2000	No serious adverse events reported	No mention of participants withdrawing due to side effects	N/A
Trinquand 1999	No serious adverse events reported	No mention of participants withdrawing due to side effects	N/A
Varguez-Rodriguez 2009	No serious adverse events reported	No participants recorded to have withdrawn	N/A
Verin 2001	No serious adverse events reported	3 participants withdrawn	3 participants from emedastine group
Wertheimer 1997	No serious adverse events reported	No participants recorded to have withdrawn due to side effects	N/A

N/A: not applicable

APPENDICES

Appendix I. Listed medications and mechanism of action

Antazoline - pure antihistamine
 Azelastine - antihistamine + mast cell stabiliser
 Bepotastine besilate - antihistamine + mast cell stabiliser
 Carebastine - developing drug - pure antihistamine
 Levocabastine - pure antihistamine
 Emedastine - pure antihistamine
 Ketotifen - antihistamine + mast cell stabiliser
 Olopatadine - antihistamine + mast cell stabiliser
 Epinastine - antihistamine + mast cell stabiliser
 Sodium chromoglycate - antihistamine + mast cell stabiliser
 Nedocromil sodium - antihistamine + mast cell stabiliser

Lodoxamide - antihistamine + mast cell stabiliser

Tranilast - developing drug - antihistamine + mast cell stabiliser

Appendix 2. CENTRAL search strategy

#1 MeSH descriptor Conjunctivitis, Allergic

#2 conjunctivitis

#3 pollen near/3 allerg*

#4 hayfever

#5 hay near/2 fever

#6 (#1 OR #2 OR #3 OR #4 OR #5)

#7 MeSH descriptor Histamine H1 Antagonists

#8 antihistamin* or anti-histamin*

#9 antazolin*

#10 azelastin*

#11 bepotastin* or Senju

#12 carebastin*

#13 levocabastin*

#14 emedastin*

#15 MeSH descriptor Ketotifen

#16 ketotifen*

#17 olopatadin*

#18 epinastin*

#19 MeSH descriptor Cromolyn Sodium

#20 sodium chromoglycate

#21 MeSH descriptor Nedocromil

#22 nedocromil sodium

#23 lodoxamid*

#24 tranilast*

#25 (#7 OR #8 OR #9 OR #10 OR #11 OR #12 OR #13 OR #14 OR #15 OR #16 OR #17 OR #18 OR #19 OR #20 OR #21 OR #22 OR #23 OR #24)

#26 (#6 AND #25)

Appendix 3. MEDLINE (OvidSP) search strategy

1. randomized controlled trial.pt.

2. (randomized or randomised).ab,ti.

3. placebo.ab,ti.

4. dt.fs.

5. randomly.ab,ti.

6. trial.ab,ti.

7. groups.ab,ti.

8. or/1-7

9. exp animals/

10. exp humans/

11. 9 not (9 and 10)

12. 8 not 11

13. conjunctivitis, allergic/

14. conjunctivitis.tw.

15. (pollen adj3 allerg\$).tw.

16. hayfever.tw.

17. (hay adj2 fever).tw.

18. or/13-17
19. exp histamine h1 antagonists/
20. (antihistamin\$ or anti-histamin\$).tw.
21. antazolin\$.tw.
22. azelastin\$.tw.
23. (bepotastin\$ or Senju).tw.
24. carebastin\$.tw.
25. levocabastin\$.tw.
26. emedastin\$.tw.
27. Ketotifen/
28. ketotifen\$.tw.
29. olopatadin\$.tw.
30. epinastin\$.tw.
31. Cromolyn Sodium/
32. sodium chromoglycate.tw.
33. Nedocromil/
34. nedocromil sodium.tw.
35. lodoxamid\$.tw.
36. tranilast\$.tw.
37. or/19-36
38. 18 and 37
39. 12 and 38

The search filter for trials at the beginning of the MEDLINE strategy is from the published paper by Glanville ([Glanville 2006](#)).

Appendix 4. EMBASE (OvidSP) search strategy

1. exp randomized controlled trial/
2. exp randomization/
3. exp double blind procedure/
4. exp single blind procedure/
5. random\$.tw.
6. or/1-5
7. (animal or animal experiment).sh.
8. human.sh.
9. 7 and 8
10. 7 not 9
11. 6 not 10
12. exp clinical trial/
13. (clin\$ adj3 trial\$).tw.
14. ((singl\$ or doubl\$ or trebl\$ or tripl\$) adj3 (blind\$ or mask\$)).tw.
15. exp placebo/
16. placebo\$.tw.
17. random\$.tw.
18. exp experimental design/
19. exp crossover procedure/
20. exp control group/
21. exp latin square design/
22. or/12-21
23. 22 not 10
24. 23 not 11
25. exp comparative study/
26. exp evaluation/

27. exp prospective study/
28. (control\$ or prospectiv\$ or volunteer\$).tw.
29. or/25-28
30. 29 not 10
31. 30 not (11 or 23)
32. 11 or 24 or 31
33. exp allergic conjunctivitis/
34. conjunctivitis.tw.
35. (pollen adj3 allerg\$).tw.
36. hayfever.tw.
37. (hay adj2 fever).tw.
38. or/33-37
39. histamine H1 receptor antagonist/
40. (antihistamin\$ or anti-histamin\$).tw.
41. antazoline/
42. antazolin\$.tw.
43. azelastine/
44. azelastin\$.tw.
45. carebastine/
46. (bepotastin\$ or Senju).tw.
47. carebastin\$.tw.
48. levocabastine/
49. levocabastin\$.tw.
50. emedastine/
51. emedastin\$.tw.
52. ketotifen/
53. ketotifen\$.tw.
54. olopatadine/
55. olopatadin\$.tw.
56. epinastine/
57. epinastin\$.tw.
58. cromoglycate disodium/
59. sodium chromoglycate.tw.
60. nedocromil sodium/
61. nedocromil sodium.tw.
62. lodoxamide/
63. lodoxamid\$.tw.
64. tranilast/
65. tranilast\$.tw.
66. or/39-65
67. 38 and 66
68. 32 and 67

Appendix 5. *meta*Register of Controlled Trials search strategy

allergic conjunctivitis and antihistamine

Appendix 6. ClinicalTrials.gov search strategy

allergic conjunctivitis AND antihistamine

Appendix 7. ICTRP search strategy

allergic conjunctivitis AND antihistamine

CONTRIBUTIONS OF AUTHORS

Conceiving the review: MSM, MC, AAB

Designing the review: MSM, MC, AAB, NS

Drafting the protocol: MSM, MC, AAB

Coordinating the review: MC

Undertaking electronic and manual searches: MSM, MC, MZM

Screening search results: MSM, MC, MZM

Organising retrieved papers against inclusion criteria: MC

Appraising methodological quality of papers: MC, AAB, NS

Abstracting data from papers: MC, MSM, MZM, NS, AAB

Data management of the review: MC

Entering data into RevMan: MC, NS

Analysing and presenting results: MC, NS, AAB

Interpreting results: MC, NS, AAB

Writing the review: MC, NS, AAB

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We declare that there is no conflict of interest or financial interest in this systematic review by any of the review authors.

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DIFFERENCES BETWEEN PROTOCOL AND REVIEW

None.

NOTES

The current protocol is an update of a published protocol by Purohit G, Jayabalan P, Ratnasingham K, Patter A. Topical antihistamines for seasonal and perennial allergic conjunctivitis. Cochrane Database of Systematic Reviews 2006, Issue 1. The 'Background' and 'Objectives' sections of the protocol have been updated, and the 'Methods' section has been rewritten in line with updated Cochrane methodology. The protocol has also been peer reviewed again.