Topical ophthalmic β blockers may cause release of histamine through cytotoxic effects on inflammatory cells

Luc M van Beek, Marcel Mulder, Nicolaas J van Haeringen, Aize Kijlstra

Abstract

Aim—To evaluate the effects of β blockers used in ophthalmology on the release of histamine from mixed cell preparations containing human leucocytes and basophils.

Methods—A mixed leucocyte and basophil preparation was obtained from venous blood of healthy non-atopic volunteers. Cell preparations were then incubated with betaxolol, metipranolol, timolol, or carteolol. After incubation for 1 hour the histamine content of the supernatant was analysed by automated fluorometric analysis. Cell viability was tested by measuring lactate dehydrogenase (LDH) concentrations.

Results—Betaxolol and metipranolol in concentrations between 10⁻² M and 10⁻⁷ M liberated histamine from human blood cells in a dose dependent manner. Carteolol and timolol had no effect on histamine at these concentrations. At the same concentrations LDH was also detected in the supernatants of cell suspensions incubated with metipranolol or betaxolol.

Conclusions—Betaxolol and metipranolol induce substantial histamine release from human leucocytes, probably as a result of their cytotoxic effect.

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Topical β blockers are widely used to lower intraocular pressure in glaucoma and ocular hypertension. Topical β blockers can cause serious systemic side effects;ocular side effects including stinging or burning upon instillation, foreign body sensation, dry eyes, periorbital dermatitis, conjunctivitis, keratitis, and uveitis have also been described.1 Mast cells, which are abundant in the conjunctiva, play a pivotal role in type I hypersensitivity. Mast cells release histamine and other preformed mediators upon stimulation. Histamine causes itching, hyperaemia, and increases vascular permeability. Earlier reports indicate that histamine is released from isolated rat mast cells after incubation with certain β blockers.2 This suggests that some β blockers may have pseudallergic properties which may account for some of the ocular side effects of these drugs.

Release of histamine from mast cells (degranulation) is the result of either IgE dependent or IgE independent mechanisms. IgE is bound to high affinity IgE receptors (FcεRI) on the membrane of mast cells and basophils. Cross linking of these IgE antibodies by a divalent antigen (allergen) results, via a molecular cascade, in an increase in the intracellular calcium concentration. This in turn leads to release of histamine and other preformed mediators from the granules of the mast cell. Cross linking of the IgE antibodies also activates the enzyme adenyl cyclase which promotes the synthesis of cyclic adenosine monophosphate (cAMP) from ATP. Cyclic AMP inhibits degranulation, thus forming a negative feedback loop.3

Mast cells and basophils express β receptors at their surface; these receptors are also coupled to adenylyl cyclase.7 Activation of β receptors increases the intracellular cAMP concentration which then acts as a second messenger. Blocking the β receptors may thus disturb the negative feedback loop, making the mast cells more susceptible to degranulation. There have been several reports of a worsening or recurrence of atopic or anaphylactic reactions after starting treatment with systemic β blockers.6

Betaxolol, carteolol, metipranolol, and timolol are β blocking agents currently used for the treatment of glaucoma. We investigated whether these compounds could induce histamine release. As a model we used human basophils which have similar histamine releasing properties to mast cells.7

Materials and methods

ISOLATION OF LEUCOCYTES

Blood samples from healthy volunteers were collected in citrate tubes and suspended in an ice cold hypotonic ammonium chloride buffer, causing lysis of erythrocytes. The leucocytes were then washed twice with ice cold HEPES buffer containing 132 mM NaCl, 6 mM KCl, 1 mM CaCl₂, 1 mM MgSO₄, 1.2 mM KH₂PO₄, 20 mM N-2-hydroxethylpiperazine-N-2-ethanesulphonic acid (Sigma Chemical Co, St Louis, MO, USA), 5.5 mM glucose, and 0.5% (w/v) human serum albumin, pH 7.4, osmolality 280–300 mOsm/l. After washing the leucocytes were suspended in HEPES buffer and incubated with various β blockers.

The volunteers (n = 14) were free of atopic disease and were not taking any medication. Several different concentrations (usually three) of β blockers were tested in duplicate on each blood sample. All volunteers gave their informed consent and the study conformed to the Universal Declaration of Human Rights and to the European Convention for the Protection of Human Rights and Fundamental Freedoms.

The number of leucocytes per aliquot ranged from 46.2 × 10⁶ to 73.5 × 10⁶. Approxi-
Release of histamine by topical ophthalmic β blockers

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To determine the release of histamine from isolated leucocytes from human donors, we incubated suspensions of human leucocytes with topical ophthalmic β blockers at the highest concentration (10–2 M). After 1 hour incubation with β blockers (0.01 M) used in treatment of glaucoma, we measured the LDH content of the supernatant.

### Results

Initially we tested four β blockers (carteolol, betaxolol, metipranolol, and timolol) for their ability to release histamine from isolated leucocytes. These β blockers are currently in use for the treatment of glaucoma. After incubation for 1 hour at a concentration of 0.01 M, betaxolol and metipranolol released 79% and 88%, respectively, of the total histamine content from the basophils. At the same concentration timolol released only 18% of the histamine and carteolol did not release any (Table 1).

<table>
<thead>
<tr>
<th>β blocker</th>
<th>Mean (SD) histamine release (%)</th>
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<tbody>
<tr>
<td>Betaxolol</td>
<td>88.5 (5.1)</td>
</tr>
<tr>
<td>Metipranolol</td>
<td>78.8 (6.9)</td>
</tr>
<tr>
<td>Timolol</td>
<td>18.5 (11.5)</td>
</tr>
<tr>
<td>Carteolol</td>
<td>0 (0)</td>
</tr>
</tbody>
</table>

Table 1: Histamine release from human leucocyte suspensions of healthy volunteers (n=14) after 1 hour incubation with β blockers (0.01 M) used in treatment of glaucoma.

Directly after starting the reaction, extinction was measured with a photospectrometer (iEMS Reader MF, Labsystems, Helsinki, Finland) at 340 nm every minute for 10 minutes to follow clearance of NADH from the reaction mixture. It was assumed that the change in the level of absorption was proportional to the amount of LDH activity in the samples. The amount of LDH was expressed as a percentage of the total, corrected for spontaneous release which never exceeded 10%. Total LDH was obtained by sonification of one aliquot of leucocytes. Sonification took place with a 30 kHz microprobe (Soniprep 150, MSE, Loughborough, UK) for 15 seconds and was repeated seven times during which the suspension was kept on ice. After sonification the suspension was centrifuged at 500 g for 15 minutes. Spontaneous release was determined by measuring the LDH content of the control supernatant.

### Agents

Betaxolol, carteolol, metipranolol, and timolol were provided by Tramedico BV, Weesp, Netherlands. They were dissolved in phosphate buffered saline without preservatives.

### Histamine Assay

Perchloric acid to a final concentration of 2% was added to 250 µl of the supernatant of incubated cell suspensions. The supernatants were then stored at 4°C until analysis (maximum 60 days). The histamine content was determined at the department of allergy at the central laboratory of blood transfusion (Amsterdam) by an automated fluorophotometric o-phthalaldehyde assay. Briefly, this method consists of extracting histamine and coupling it to o-phthalaldehyde. The fluorescence of this complex is then measured by a spectrofluorometer using an excitation wavelength of 355 nm and an emission wavelength of 450 nm. It is assumed that the measured fluorescence was proportional to the concentration of histamine.

Because the number of basophils and the amount of histamine vary between individuals, the histamine content was expressed as a percentage of the total histamine. Total histamine was obtained by adding perchloric acid (final concentration 2 mg/ml) to one aliquot of leucocytes in each blood sample to cause lysis of the cells. This aliquot was then centrifuged and histamine was determined using the method described above. Percentages were corrected for spontaneous release which was defined as the histamine content of the supernatant of the control sample. The results were discarded if spontaneous release exceeded 15%.

### Lactate Dehydrogenase (LDH) Assay

Directly after the incubation experiments the supernatant of incubated cell suspensions was analysed for LDH content. Supernatant (5 µl) was mixed with 200 µl substrate buffer in a 96 well flat bottom plate. The substrate buffer contained 3.8 mM Na2PO4, 1.9 mM KH2PO4, 7.9 µg/ml pyruvic acid, and 15.9 µg/ml NADH, pH 7.0. In this mixture LDH converts NADH to NAD at a rate determined by the amount of LDH present. Experiments were performed at room temperature. Each 96 well plate contained total, control, and tested β blocker concentrations in duplicate.

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not consider that high concentrations of explain the histamine liberation because we do determine the cytotoxic e

cytotoxic blockers are less toxic to basophils than to leucocytes. This cytotoxic e

LDH in the supernatant indicated that the cytotoxicity caused by perturbation of the cell membrane. We feel that, because of their higher lipid solubility than betaxolol and metipranolol (Table 2). We feel that, because of their higher lipid solubility than betaxolol and metipranolol (Table 2).17 We feel that, because of their higher lipid solubility than betaxolol and metipranolol disturbances the cell membrane integrity and that this accounts for their higher cytotoxicity. However, it should be kept in mind that eye drops need to have a certain lipid solubility to penetrate the cornea. In a single drop study timolol was found to reduce exercise tachycardia in contrast to metipranolol. The authors explained this by the higher lipid solubility of metipranolol permitting rapid cornea penetration and leaving less metipranolol available for systemic absorption.

In general, patients find timolol eye drops more comfortable than metipranolol or betaxolol eye drops.20 Because the vehicles of the three eye drops are comparable, the low cytotoxic effect of timolol may be partly responsible for its higher degree of comfort.

This research was supported by a grant of Dr Gerhard Mann GmbH, Germany, and Tramedico BV, Netherlands.

Table 2 Lipid solubility of the tested β blockers

<table>
<thead>
<tr>
<th>β blocker</th>
<th>Lipid solubility (log P)</th>
</tr>
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<tbody>
<tr>
<td>Betaxolol</td>
<td>2.81</td>
</tr>
<tr>
<td>Metipranolol</td>
<td>2.28</td>
</tr>
<tr>
<td>Timolol</td>
<td>1.16</td>
</tr>
<tr>
<td>Carteolol</td>
<td>0.82</td>
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