

RESEARCH ARTICLE

Open Access



# Histamine and migraine revisited: mechanisms and possible drug targets

Jacob Worm, Katrine Falkenberg and Jes Olesen\* 

## Abstract

**Objective:** To review the existing literature on histamine and migraine with a focus on the molecule, its receptors, its use in inducing migraine, and antihistamines in the treatment of migraine.

**Background:** Histamine has been known to cause a vascular type headache for almost a hundred years. Research has focused on antihistamines as a possible treatment and histamine as a migraine provoking agent but there has been little interest in this field for the last 25 years. In recent years two additional histamine (H<sub>3</sub> and H<sub>4</sub>) receptors have been discovered and a series of non-sedating antihistamines have been developed. It is therefore timely to review the field again.

**Methods:** For this review the PubMed/MEDLINE database was searched for eligible studies. We searched carefully for all articles on histamine, antihistamines and histamine receptors in relation to migraine and the nervous system. The following search terms were used: histamine, migraine disorders, migraine, headache, antihistamines, histamine antagonists, clinical trials, induced headache, histamine H<sub>3</sub> receptor, histamine H<sub>4</sub> receptor and pharmacology. Four hundred thirty-six titles were read, 135 abstracts were read, 112 articles were read in full and 53 articles were used in this review. Review process resulted in 12 articles added to a total of 65.

**Findings:** Early studies of H<sub>1</sub> and H<sub>2</sub> antihistamines lack scientific strength and show conflicting results. Most of the antihistaminic drugs used in these trials bind also to other receptors which makes it difficult to conclude on the antihistaminic effect. Histamine is an efficient inducer of migraine attacks in migraine patients by an H<sub>1</sub> mechanism most likely extracerebrally. These findings merit further investigation of antihistamines in clinical drug trials. The H<sub>3</sub> and H<sub>4</sub> receptors are found in primarily in CNS and immune tissues, respectively. H<sub>3</sub> is likely to be involved in antinociception and has been linked with cognitive, neurodegenerative and sleep disorders. The only marketed H<sub>3</sub> agent, pitolisant, is a brain penetrant H<sub>3</sub> antagonist/inverse agonist which increases central histamine and causes headache. The experimental H<sub>3</sub> agonist N<sup>q</sup>-methylhistamine has shown promising results as a migraine preventative in studies of uncertain quality. With the current limited knowledge of the H<sub>4</sub> receptor it is questionable whether or not the receptor is involved in migraine.

**Conclusion:** There is insufficient support for first generation antihistamines (both H<sub>1</sub> and H<sub>2</sub>) as preventive migraine medications and sedation and weight gain are unacceptable side effects. Non-sedating H<sub>1</sub> antihistamines need to be appropriately tested. Central H<sub>3</sub> receptors seem to have a role in migraine that merit further investigation. The histaminergic system may be a goal for novel migraine drugs.

**Keywords:** Migraine, Histamine, Antihistamines, Histamine receptors, Drug targets

\* Correspondence: [j.olesen@regionh.dk](mailto:j.olesen@regionh.dk)

Danish Headache Center and Department of Neurology N39, University of Copenhagen, Rigshospitalet Glostrup, DK-2600 Copenhagen, Denmark



## Introduction

Histamine is a biogenic monoamine synthesized from the amino acid L-histidine by the L-histidine decarboxylase and metabolized by the enzyme histamine N-methyltransferase, using S-adenosylmethionine as a methyl donor. Histamine is located throughout the entire organism with high concentrations in lungs, skin and gastrointestinal tract. It is synthesized and stored in mast cells and basophils. It plays a role in multiple mechanisms both immunological and physiological, stimulating gastric secretion, inflammation, smooth muscle contraction, vasodilatation, permeability and much more. Histamine also functions as a neurotransmitter. It is synthesized in histaminergic neurons located in the posterior hypothalamus. These neurons have axons extending throughout the brain. Histamine carries out its effects via 4 subtypes of 7-transmembrane G-protein coupled receptors; H<sub>1</sub>, H<sub>2</sub>, H<sub>3</sub> and H<sub>4</sub> receptors [1–3].

The relation between histamine and headache has been on the agenda for almost a hundred years. A multitude of studies have been published focusing on the ability of histamine to induce headache and on the effect of antihistamines in the treatment of headache. During the last two decades the issue has, however, received almost no interest apart from two reviews [2, 4]. One reason that we now take this issue up is that there has been a massive development in the basic understanding of histamine biology, including the growing knowledge of histamine H<sub>3</sub> and H<sub>4</sub> receptors. Another is that a series of new generation antihistaminic drugs with or without the ability to penetrate the blood brain barrier have become available. It is therefore timely to review the issue.

## Methods

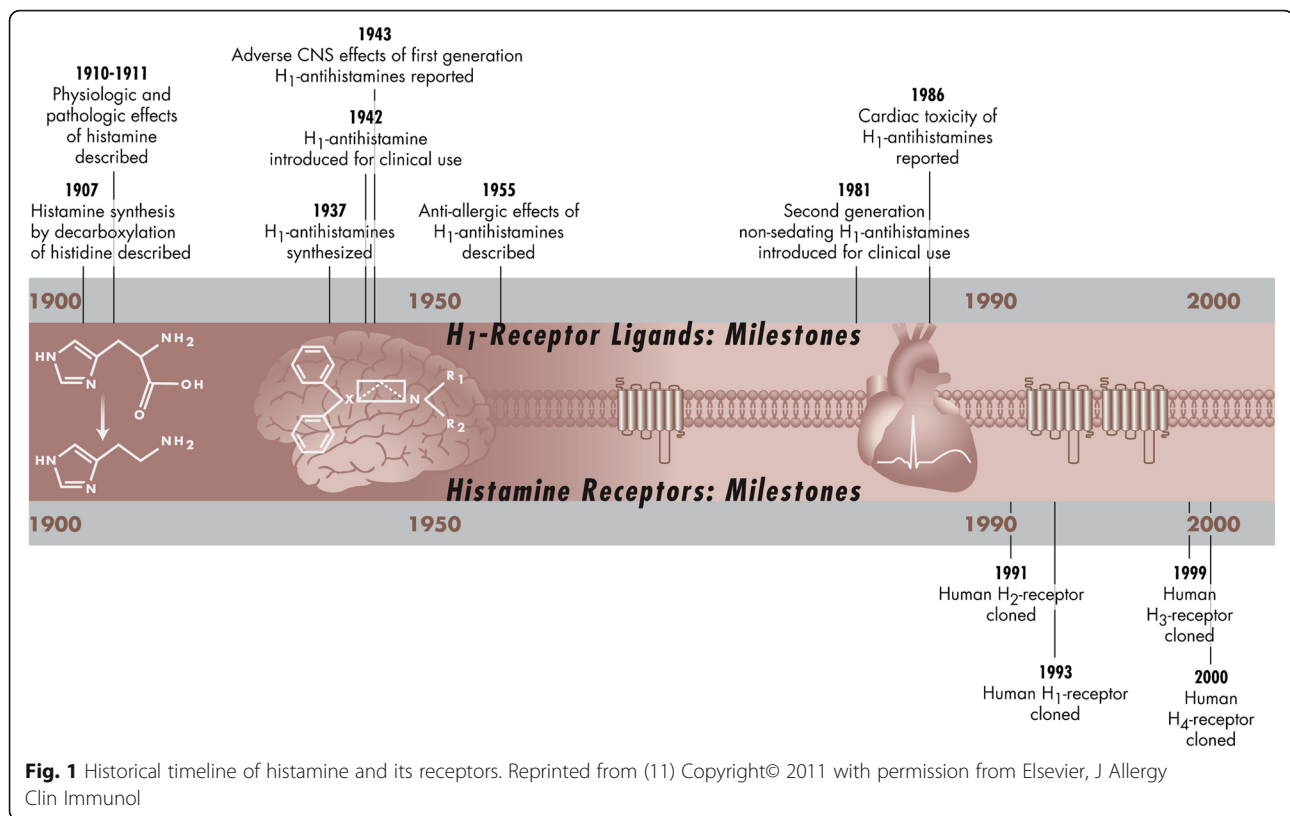
Literature used in this review was primarily searched in the PubMed/MEDLINE database and we limited our search to publications in English. References were selected by reading titles and abstracts judging relevance to the topic. We began our search with a screening to identify existing reviews of histamine and migraine. The search histamine [MeSH Terms] OR histamine [All Fields] AND migraine disorders [MeSH Terms] OR migraine [All Fields] AND disorders [All Fields] OR migraine disorders [All Fields] OR migraine [All Fields] with only articles from the last 5 years resulted in 41 articles of which 9 abstracts were read. We then searched for the very early literature of histamine and vascular headaches. This literature dates back almost a century and a systematic search is not possible. Therefore, bibliography from the PhD thesis by Lassen LH was examined to obtain references. Six articles were found eligible. To find articles regarding the studies of histamine antagonists in migraine prophylaxis we searched for the terms:

antihistamine AND (migraine OR headache) AND clinical trial from 1945 to 1985. This resulted in 68 articles of which 41 abstracts were read. Articles concerning migraine provocation studies with histamine were searched for using the terms: histamine AND induced AND headache from 1980 to present. One hundred twenty-three articles were found of which 24 abstracts were read. Articles of histamine H<sub>3</sub> and H<sub>4</sub> receptor pharmacology was searched for by the MeSH terms: Pharmacology AND Receptors, histamine with review as publication type. The search led to 44 articles of which 21 abstracts were read. Individual free text searches of histamine H<sub>3</sub> receptor OR histamine H<sub>4</sub> receptor AND pharmacology from 2000 until present was made and yielded 160 articles of which 40 abstracts were read. In addition to the search results we looked through the reference lists of the selected articles to find more relevant studies. The search combined resulted in 436 titles, of which 135 abstracts were read and 112 articles were read in full. Fifty-three were included in this review. Review process resulted in 12 articles added to a total of 65.

## Historical overview

The research of histamine began over a century ago [5], with the first antihistamines being introduced to clinical use around 70 years ago (Fig. 1) [1]. In 1926 Harmer and Harris were the first to describe headache after infusion of histamine [6]. In the 1930s Pickering did a series of careful experimental studies. Although none of them were placebo controlled, histamine was intelligently investigated for its ability to induce a vascular headache. The headache after infusion of histamine was thoroughly described and it was shown that histamine induced headache in a dose-dependent manner but that there was great variability between subjects. Pickering proposed that the origin of the headache was intracranial. He suggested stretching of large intracranial arteries to be a possible explanation of the perceived pain because headache severity was reduced by lowering blood pressure or raising intracranial pressure (i.e. less expansion of the vasculature). He concluded that it is in the dura mater, via its trigeminal nerve supply, that the headache arises [7].

In the following years several papers investigated the origin of histamine induced headache. Northfield injected histamine in the external and internal carotid artery and concluded that the pain arose from the internal carotid artery i.e. not from dura mater [8]. Another paper argued that the origin of pain in migraine headache is different from that of a histamine induced headache because of the absence of relief of migraine headache by increasing intracranial pressure while confirming a decrease of histamine induced headache [9]. Von Storch investigated the minimum dose of histamine required to induce headache.



Migraineurs seemed to be more susceptible to histamine induced headache than control subjects [10]. Histamine was also tested as a treatment of migraine [11, 12]. The idea was to induce histamine tolerance in migraineurs based on a theory that these patients had lowered tolerance thus susceptible to external triggers of histamine (i.e. foods, antigens etc.). A method found successful but difficult to practice. The older literature had thus shown beyond doubt that histamine could induce a vascular type headache, the characteristics of which were carefully described. But how close was it to migraine? This could not be determined because there were no internationally accepted diagnostic criteria for migraine.

Experimental in vitro and animal studies have been performed adjacent to the clinical studies investigating histamines vasodilatory effect in the meningeal blood vessels as part of the underlying mechanism of migraine [13]. The research began decades ago with multiple studies published around the millennium [14] and researchers continue to investigate the endogenous mediators including histamine for their ability to generate migraine pain [15]. Histamine has been extensively studied alongside cytokines, chemokines and vasoactive peptides (i.e. the inflammatory soup) to gain knowledge of the underlying cellular mechanisms producing and maintaining the pain of migraine. At present a link

between migraine and neurogenic inflammation comprising mast cell degranulation is widely accepted and this has been reviewed extensively most recently by Ramachandran [16].

#### Are H<sub>1</sub> and H<sub>2</sub> blockers effective in migraine treatment?

Because histamine induces a vascular headache with similarity to migraine, it was logical to test antihistamines in the treatment of migraine when the class of antihistamines became available. These old drugs were sedative and soon became classified as histamine receptor 1 and 2 antagonists as no other histamine receptors or antagonists were known at the time. The literature on clinical trials of the H<sub>1</sub> and H<sub>2</sub> receptor antagonists in the prevention of migraine is rather scarce. Few papers have been published. Even fewer are randomized controlled trials and mostly used drugs that did not bind exclusively to H<sub>1</sub> or H<sub>2</sub> receptors. The following section examines the existing literature on antihistamine trials performed in the years from around 1960 to 2014 (see Table 1 for overview). All the existing literature describes antihistamines as histamine receptor antagonists. Today most of these drugs are defined pharmacologically as inverse agonists i.e. inhibitors of the basal receptor activity instead of being neutral antagonists which

**Table 1** Antihistamines in migraine prophylaxis

Type (Ligand)	Participants	Daily dose	Duration	Main findings	First author (year)
H <sub>1</sub> antagonist (cyproheptadine) vs. 5HT antagonist (methysergide) vs. Bellergal <sup>b</sup> vs. placebo	453 <sup>a</sup>	12–24 mg 6 mg NA NA	6 months	Clinical improvement (Headache free or substantial improvement):c: methysergide 64%, cyproheptadine 46%, Bellergal 34%, placebo 20%.	Curran (1964) (12)
Antihistaminic drug (pizotifen)	32 <sup>a</sup>	1.5–2 mg	NA	Cessation or attenuation of attacks in 33%. Significant reduction in migraine days and increased response to acute medication in 40%. Increased resilience against induced attacks. AE <sup>d</sup> : weight gain and drowsiness.	Sicuteri (1967) (13)
Antihistaminic drug (pizotifen)	11 MA <sup>e</sup> 27 MO <sup>f</sup> 2 Cluster headache	1.5 mg (1–3 mg)	2–6 mo: 16 > 6 mo: 17	Significant (42.5%) clinical improvement compared to previous placebo reports (Curran 1964).	Selby (1970) (14)
H <sub>1</sub> antagonist (cyproheptadine) vs. antihistaminic drug (pizotifen) vs. antihistaminic drug (methdilazine)	165 <sup>a</sup>	12–24 mg 4.5–9 mg 16–32 mg	4 weeks	Clinical improvement: pizotifen 58%, methdilazine 41%, cyproheptadine 40% No placebo control.	Lance (1970) (15)
Antihistaminic drug (pizotifen) vs. 5HT antagonist (divascan) vs placebo	4 MA <sup>e</sup> 26 MO <sup>f</sup>	3 mg 15 mg	8 weeks	Significant lowered attack rate with pizotifen. AE <sup>d</sup> : weight gain and drowsiness.	Osterman (1977) (16)
Antihistaminic drug (pizotifen) vs placebo	28 <sup>a</sup>	3 mg	12 weeks	Complete resolution in 6 cases, reduced frequency and severity in 6 and no improvement in 2 cases. No improvement in the placebo group. AE <sup>d</sup> : weight gain and dizziness.	Lawrence (1977) (17)
H <sub>2</sub> (cimetidine) vs. H <sub>2</sub> and H <sub>1</sub> combined (cimetidine and chlorpheniramine) vs. placebo	24 MO 1 basilar migraine	200 mg cimetidine 4 mg chlor pheniramine	1 week	No significant improvement over placebo with H <sub>2</sub> antagonist alone or in combination with H <sub>1</sub> antagonist. AE <sup>d</sup> : weight gain and drowsiness.	Anthony (1978) (18)
H <sub>2</sub> (cimetidine) vs. H <sub>2</sub> and H <sub>1</sub> combined (cimetidine and chlorpheniramine) vs. placebo	6 MA 28 MO	200 mg cimetidine 4 mg chlor- pheniramine	12 weeks	No significant improvement over placebo with H <sub>2</sub> antagonist alone or in combination with H <sub>1</sub> antagonist. AE <sup>d</sup> : weight gain and drowsiness.	Nanda (1980) (20)
H <sub>1</sub> (cinnarizine)	11MA <sup>e</sup> 69 MO <sup>f</sup>	75 mg	14 weeks	Significant reduction in migraine days and use of acute medication. AE <sup>d</sup> : weight gain, drowsiness, mild reversible depression and dyspepsia. Open label. Dropout rate 3.75%.	Rossi (2003) (22)
H <sub>1</sub> (cinnarizine)	60 <sup>g</sup>	Up to 75 mg	12 weeks	Significant reduction in migraine days. AE <sup>d</sup> : Palpitations and dizziness. Open label. Dropout rate 5%.	Togha (2006) (21)
H <sub>1</sub> (cinnarizine) vs valproate	86 MO <sup>f</sup> 18 MA <sup>e</sup>	50 mg Cinnarizine 400 mg valproate	12 weeks	Valproate more effective than cinnarizine. AE <sup>d</sup> : Dry mouth, fatigue and somnolens as the most frequent. No placebo control. Dropout rate: 23% cinnarizine, 19.4% valproate.	Bostani (2013) (23)
H <sub>1</sub> (cinnarizine) vs placebo	68 children (5–17y) <sup>a</sup>	50 mg or 1.5 mg/kg (> 30 kg)	12 weeks	Significant better than placebo to reduce headache frequency by at least 50% reduction. AE <sup>d</sup> : weight gain and drowsiness. Dropout rate: 12% cinnarizine, 5% placebo.	Ashrafi (2014) (24)

<sup>a</sup>No subtype stated<sup>b</sup>Mixture of phenobarbitone 20 mg, ergotamine tartrate 0.3 mg and belladonna alkaloids 0.1 mg<sup>c</sup>Not placebo corrected<sup>d</sup>Adverse effects<sup>e</sup>Migraine with aura<sup>f</sup>Migraine without aura<sup>g</sup>Subtype unknown

solely block the agonist response [3, 17]. For ease, we will describe all antihistamines as compounds with a histamine antagonistic effect.

In a placebo controlled, but not double blinded or randomized study, Curran and Lance investigated cyproheptadine, an antiserotonin agent with H<sub>1</sub> receptor antagonism, methysergide, a serotonin antagonist and

bellergal (mixture of phenobarbitone 20 mg, ergotamine tartrate 0.3 mg and belladonna alkaloids 0.1 mg). Their ability to reduce attack frequency was compared and clinical improvement was defined as either headache free or substantially improved. Ninety-two patients received 12–24 mg/day of cyproheptadine, 137 patients received 6 mg/day of methysergide, 174 patients received the

bellergal mixture and 50 were treated with placebo. Clinical improvement was 34% with bellergal, 64% with methysergide, 46% with cyproheptadine and 20% with placebo. No statistical analysis was made comparing cyproheptadine with placebo [18]. Thus, the result with cyproheptadine is only suggestive of a possible effect of antihistamines in migraine treatment.

Sicuteri et al. treated 32 outpatients suffering from migraine with the antihistaminic substance named BC-105, later known as pizotifen. BC-105 also possesses a strong antiserotonin effect. Patients were treated with a maintenance dose of 1.5–2 mg/day (up to 4–5 mg). No predetermined mandatory duration was mentioned and the study was not blinded or placebo controlled. Response to treatment was usually experienced after 10–20 days of treatment. A third of the patients reported absence or significant attenuation of attacks and 40% reported a lower frequency or increased response to acute therapy with ergotamine. Interestingly, BC-105 increased the resilience against migraine attacks induced with trinitroglycerin and reserpine. Side effects were well documented in this study, with 90% of the patients developing an increased appetite. Patients gained 2–5 kg during the first month of treatment. Fatigue and drowsiness was a side effect in 70%. The drug was administered with a starting dose of 0.5 mg twice a day (b.i.d.) increasing to 1 mg b.i.d. after 4–5 days. This was to minimize the initial drowsiness typically experienced during the first period of treatment [19].

Selby et al. studied BC-105 in 40 patients, 11 with migraine with aura, 27 with migraine without aura and 2 with cluster headache. Thirty-five were treated with a dose of 1.5 mg (3 tablets) daily. Two patients received 1 mg and three patients received 2–3 mg/day. The design was uncontrolled without placebo. 42.5% (17 patients) clinically improved with 2 becoming headache free and 15 with a reduction of attack frequency between 25 and 50%. This study reported fewer side effects which the author interpreted as a result of the lesser dose [20].

Another comparative trial investigated three drugs relevant to histamine. Cyproheptadine, BC-105 (pizotifen) and methdilazine, a drug with a four times greater antihistaminic action than promethazine. The patients recruited were not randomly allocated and furthermore had the possibility to shift to another group after 1 month if the current treatment failed. Because of the free movement between the groups, only data for the first month were statistically analyzed. The placebo group of patients was drawn from two other studies conducted simultaneously. Clinical improvement after 1 month was defined as headache free or a reduction in attack frequency of at least 50%. Sixty-one patients received cyproheptadine 4–8 mg three times a day (t.i.d.) of whom 40% improved. Fifty-three patients received

1.5–3 mg of BC-105 t.i.d. with 58% improving and 51 patients received methdilazine 8–16 mg b.i.d. of which 41% improved. All three resulted in a lesser improvement than methysergide, but the compound BC-105 was also significantly better than placebo. Cyproheptadine and methdilazine showed only a positive trend [21].

In a double-blind placebo controlled crossover study, Osterman investigated the effect of Sandomigran® (pizotifen or BC-105), and Divascan®, a serotonin antagonist in migraine prophylaxis. Four patients with migraine with aura and 26 patients with migraine without aura were included. All patients experienced at least 2–3 attacks per month and had a confirmed diagnosis based on then accepted criteria (Ad hoc committee on classification of headache 1962). Three patients discontinued due to adverse events, lack of compliance or unpleasant side effects. Twenty-seven patients were divided into three groups with different treatment sequences of divascan, pizotifen and placebo. Each treatment was given for 8 weeks in a total period of 24 weeks and all three compounds were given in gradually increasing doses using the same amount of capsules. The dose of pizotifen started at 0.5 mg (1 capsule) increasing to a daily maintenance dose of 3 mg (2 capsules, t.i.d.) at day 11. Divascan started at 2.5 mg (1 capsule) increasing to a daily maintenance dose of 15 mg (2 capsules t.i.d.) at day 11. Side effects caused some patients to discontinue the maintenance doses of both drugs. Four patients had only 1.5–2.5 mg/day and 2 patients 10–12.5 mg/day of pizotifen and divascan respectively. The patients kept a headache diary during the trial, with attack frequency, duration and severity as well as the use of attack medication. Severity was rated using an arbitrary 3 point scale and only data from the last 6 weeks of each treatment were analyzed as the first 2 weeks served as wash out period. Treatment with pizotifen gave a significantly lower attack frequency than both placebo and divascan. It also significantly reduced the mean headache index. No significant difference in the duration of attacks was observed. Consumption of ergotamines was significantly higher in the placebo group versus pizotifen and divascan respectively, but not when comparing pizotifen and divascan. In overall evaluation of the treatment divascan did significantly better than placebo but pizotifen had better effect than both placebo and divascan with 70% reporting a good or very good effect. However, when looking at the side effects the authors found that the treatment with pizotifen resulted in significantly more side effects. Most frequent were drowsiness and weight gain (> 1.5 kg) [22].

A multicenter double-blind placebo controlled study in 36 patients in general practice also investigated the effect of pizotifen. All patients had a history of migraine with at least 4 attacks per month. Twenty-four of the



patients had failed previous prophylactic therapy. The dose of pizotifen was increased gradually to a daily dose of 1 mg three times a day at day 24 and onwards. The treatment period was 12 weeks and each patient kept a record of frequency and severity of migraine attacks. Twenty-eight patients completed the trial, 14 on pizotifen and 14 on placebo. No statistical analysis was made, but 6 patients reported complete resolution, 6 reported reduced frequency and severity and only 2 patients did not improve clinically. All fourteen patients given placebo failed to show any improvement [23].

Anthony and Lance performed a controlled double blind trial of the H<sub>2</sub> receptor antagonist cimetidine alone or in combination with the H<sub>1</sub> receptor antagonist chlorpheniramine as migraine prophylaxis. Twenty-four patients with migraine without aura, and 1 with basilar symptoms received 3 different treatments, each for 1 month during 3 consecutive months. The treatments consisted of 1 month of 200 mg (2 tablets four times a day (q.i.d.)) cimetidine and chlorpheniramine placebo (1 capsule q.i.d.), 1 month of 200 mg (two tablets q.i.d.) cimetidine and chlorpheniramine maleate 4 mg (1 capsule q.i.d.) and 1 month of cimetidine placebo (2 tablets q.i.d.) and chlorpheniramine placebo (1 capsule q.i.d.). Patients were randomly allocated to the treatment sequence and kept a record of headache intensity and frequency as well as any side effects. Nineteen patients completed the trial and results were that neither cimetidine in combination with chlorpheniramine nor cimetidine alone proved better than placebo. The authors interpreted the outcome as a result of the intracellular histamine formation by decarboxylation of histidine in multiple tissues, generating headache via other pathways or mechanisms than H<sub>1</sub> or H<sub>2</sub> receptors [24]. Another explanation may be a short receptor residence time at the H<sub>1</sub> receptor by chlorpheniramine (K<sub>i</sub> 5–30 nM) [4]. Furthermore, by today's standards only 1 month of preventative medication is not sufficient to precisely estimate results in migraine prophylaxis. A minimum of 3 months for a phase II and up to 6 months for a phase III study is recommended [25]. Nanda et al. investigated the same drug-placebo combination for 3 months in each group over 9 months combined. Twenty-eight patients with migraine without aura and 6 patients with migraine with aura were included in the trial of which 22 patients began all three periods, but only 16 completed the full 9 months. These patients were allowed to move forward to another treatment group before the 3 months were completed if their headache frequency intensified. The authors failed to show a positive effect on migraine even with a 3 months treatment period. They found the same side effects with drowsiness related to chlorpheniramine and dizziness related to cimetidine [26]. These two studies were blind and placebo controlled

and so provide the best evidence. However, using a H<sub>2</sub> receptor antagonist alone or in combination with H<sub>1</sub> receptor antagonist showed no significant relief of migraine when used prophylactically.

Cinnarizine, a calcium channel antagonist with H<sub>1</sub> antihistaminic action and a drug used in Meniere's disease was tested in migraine prophylaxis. Two open label trials, one with 80 participants and another with 60 tried out cinnarizine in doses up to 75 mg daily in 12 and 14 weeks respectively. Safety and efficacy was measured and both studies generally consider cinnarizine well-tolerated in their participants. Although both groups report a significant reduction of headache frequency the open trial design limits the results to only a hypothetical positive prophylactic effect of cinnarizine [27, 28]. A double blind parallel group study investigated cinnarizine vs. sodium valproate. One hundred four participants were randomized to a daily treatment with either 50 mg cinnarizine (50 participants) or 400 mg valproate (54 participants) for 12 weeks. Dropout rate was ~ 20% for both arms and mainly because of side effects. Valproate proved more effective than cinnarizine but since there was no placebo control efficacy results may be overestimated [29].

A double blind placebo controlled parallel group study of cinnarizine in the prophylaxis of migraine in children was conducted by Ashrafi et al. [30]. Sixty-eight participants aged 5–17 years with migraine with and without aura according to the 2004 IHS criteria for migraine [31] were randomized into 2 groups after 4 weeks of washout. The groups consisted of a treatment group with 50 mg cinnarizine daily (1.5 mg/kg/day for children weighing less than 30 kg) and a placebo group. During a 12 week treatment period participants kept a headache diary and primary outcome measures were frequency, intensity and duration of migraine. Adverse events were too recorded with weight gain and extra pyramidal signs monitored during the trial. Dropout rate was ~ 12% and ~ 5% for the treatment and placebo group, respectively. Cinnarizine significantly reduced headache frequency to < 50% in 60% of the patients compared to only 31.3% in the placebo group.

In conclusion, low level of evidence and conflicting results make it difficult to determine whether or not H<sub>1</sub> or H<sub>2</sub> receptor antagonists are effective in migraine treatment. There is a reasonable indication that pizotifen, cinnarizine and also amitriptyline have effect but they have many other effects in addition to being antihistamines. More pure antihistamines such as chlorpheniramine have not shown convincing efficacy. It would be interesting to study the efficacy of pure antihistamines using modern principles of drug trials [25]. The antihistamines used in the trials have primarily an H<sub>1</sub> antagonistic effect, but also exert antagonism on other receptors, mainly serotonin receptors. The serotonergic effects in

migraine is well described elsewhere [32]. Although the complete sites of action remains to be fully described. Serotonin and histamine are both secreted from the mast cells in dura mater [13, 33] and it is likely that the anti-serotonin and antihistamine effects potentiate each other, making the results difficult to interpret. Finally the old antihistamines could not be used in high doses because of sedation and weight gain. It might be worth to try high doses of non-sedating modern antihistamines.

### Histamine induction of migraine

The advent of diagnostic criteria for migraine, first unofficial [34] and then internationally accepted ICHD-1 [35] made it possible to further study the relation between histamine induced headache and migraine. Although histamine induction of headache dates back to the 1920s [6] much remained to be elucidated when the modern provocation studies began. The initial hypothesis was that H<sub>1</sub> and – to a lesser extend – H<sub>2</sub> receptors on intracranial blood vessels was involved, but the clinical studies [36], experimental studies on cerebral blood flow in human [37] and isolated intra- and extracranial blood vessels [38] found that it was in fact the vasodilatation of the extracerebral vessels that caused the pain.

Krabbe and Olesen investigated a 30 min intravenous infusion of histamine with increasing doses (0.16–0.33–0.66 µg/kg/min.) in 48 participants, 13 healthy volunteers, 10 patients with chronic tension type headache then called chronic muscle contraction headache and 25 patients with migraine without aura, then called common migraine. The objective was to evaluate the sensitivity to histamine in normal subjects and in patients with these headaches and to develop an experimental model of vascular headache. All 48 participants received a continuous infusion of histamine chloride for 30 min. 24 of 25 migraineurs developed a throbbing headache, while participants from the two other groups only experienced, at worst, a moderate, pressing type of headache. Adverse events from the histamine infusion consisted mainly of a heating sensation in the face and palpitations. Objectively, all participants developed a flushing of the face, lowering of blood pressure and increased heart rate. No serious adverse events occurred during the study.

In 18 patients who developed headache during the histamine infusion, 0.5 mg/kg of the H<sub>1</sub> receptor antagonist mepyramine was administered intravenously for the last 2 min of the histamine infusion. Mepyramine greatly reduced or abolished the headache in 15 of 18 migraineurs. In 10 of the patients suffering from migraine who developed a severe throbbing headache, another single blinded crossover study was performed with cimetidine, an H<sub>2</sub> receptor antagonist. A pretreatment with cimetidine was administered intravenously with a bolus of

3.3 mg/kg, followed by a continuous infusion of 1.66 mg/kg/hour for 45 min finished by the infusion of histamine. The pretreatment with cimetidine slightly but significantly decreased the headache. The only side effect was a metallic taste in the mouth in the majority of the patients [36]. This study proved that histamine induces more or stronger headache in migraine patients than in normal individuals. H<sub>1</sub> antagonism was effective in aborting the headache and pretreatment with an H<sub>2</sub> antagonism had some effect.

In a double blind placebo-controlled cross-over study, Lassen et al. investigated whether pretreatment with the H<sub>1</sub> receptor antagonist mepyramine had an effect on histamine induced headache. In 20 patients with migraine without aura histamine was administered intravenously as an infusion of 0.5 µg/kg/min for 20 min. Prior to this a pretreatment with 0.5 mg/kg mepyramine or placebo (0.9% NaCl) was infused intravenously for 10 min. Data from the first 3 h were documented at the study site and then data were recorded by the patients after discharge for 8 h. Histamine induced an immediate and a delayed headache. Immediate headache was defined as the headache occurring during the first 40 min after start of infusion and the delayed headache was defined as occurring from 40 min to 12 h after start of infusion.

The study was first of its kind describing that a histamine infusion can provoke a delayed genuine migraine attack in migraineurs, as 5 of 10 patients developed an attack fulfilling IHS criteria for migraine without aura. Pretreatment with mepyramine prevented development of both the immediate and the delayed headache [39]. Glyceryl trinitrate (GTN), a nitric oxide donor, is also a potent headache inducer [40] and similar to histamine capable of inducing an immediate headache followed hours later by a delayed migraine-like headache. The headache caused by both substances is biphasic in migraine patients suggesting a shared pathway. However, GTN induced migraine is resistant to H<sub>1</sub> receptor blockade with mepyramine [41] and histamine could therefore not be the common mediator. The question of nitric oxide being a common mediator of histamine and GTN induced migraine was investigated in two double-blinded and placebo controlled studies, using L-NMMA, a non-selective nitric oxide synthase inhibitor to prevent histamine induced headache in 9 healthy male volunteers [42] and in 12 patients with migraine without aura [43]. Although nitric oxide synthase (NOS) inhibition is effective as a treatment of spontaneous attacks [44, 45], both studies showed no effect on the immediate or delayed headache induced by histamine after pre-treatment with L-NMMA. This underlines the need of further investigation of histamine's site of action. Because histamine is unable to cross the blood-brain barrier and H<sub>1</sub> receptors are present in the endothelium of cerebral arteries, migraine attacks

could have an origin in the walls of intracerebral arteries and not in the brain itself [41].

Histamine inhalation was investigated by Lassen et al. in 12 patients with migraine without aura, 2 patients with migraine with aura and 1 patient with migraine both with and without aura. They were matched with 15 healthy volunteers in a double blind study. All participants underwent 2 min inhalation periods with 2-fold increase in dose of histamine for every period. The objectives were to evaluate whether histamine inhalation could provoke a headache, whether it showed dose-dependency, whether induced attacks were genuine migraine attacks and whether attacks were induced with the same frequency in both groups. Immediate and delayed headache were reported in both groups with no significant difference, but only migraineurs (6 of 15) developed a headache fulfilling the IHS criteria for migraine without aura. Development of migraine after inhalation had a positive predictive value of 1.0, but the sensitivity was poor. The clinical signs and symptoms other than headache were flushing, heat sensation, palpitation, coughing, dizziness and dysphonia, none of which were severe [46].

Histamine plasma levels in migraine patients have been reported elevated compared with controls both ictally and interictally [47] while another study reported no change during attacks but significant difference between migraineurs and controls between attacks [48]. Latest unpublished results from a biomarker study in patients with migraine without aura using mass spectrometry found that histamine levels both during and outside of attacks were below detection limit and we will not review this further.

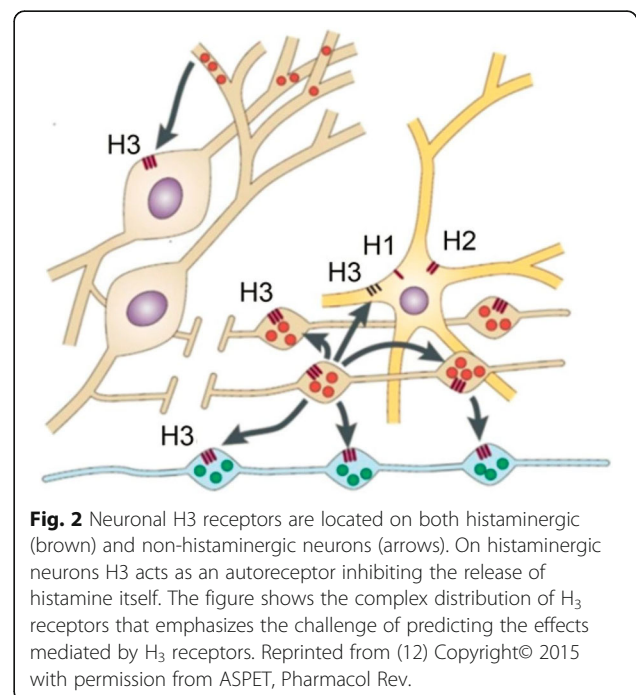
### H<sub>3</sub> and H<sub>4</sub> receptor pharmacology and possible involvement in migraine

A third member of the histamine receptor family was discovered pharmacologically by Arrang in 1983 when he showed that histamine inhibited its own release from depolarized slices of rat cerebral cortex. The receptor responsible was identified as a presynaptic inhibitory receptor – an autoreceptor – independent of other neurotransmitter antagonists and with a different ligand profile. He concluded that a novel class of histamine receptors was discovered and proposed the name H<sub>3</sub> [49]. The histamine H<sub>3</sub> receptor cDNA was cloned by Lovenberg and colleagues in 1999 and they described an extensive expression in the rat CNS especially in the caudate nucleus, thalamus and cortex [50]. Apart from the receptor's full length polypeptide approximately 20 human isoforms have been described [3]. Studies of rat, guinea pig and human brains contribute to the knowledge of the H<sub>3</sub> receptor's expression in mammalian CNS. The H<sub>3</sub> receptor has been mapped by various methods such as autoradiography and later by in situ hybridization and reverse

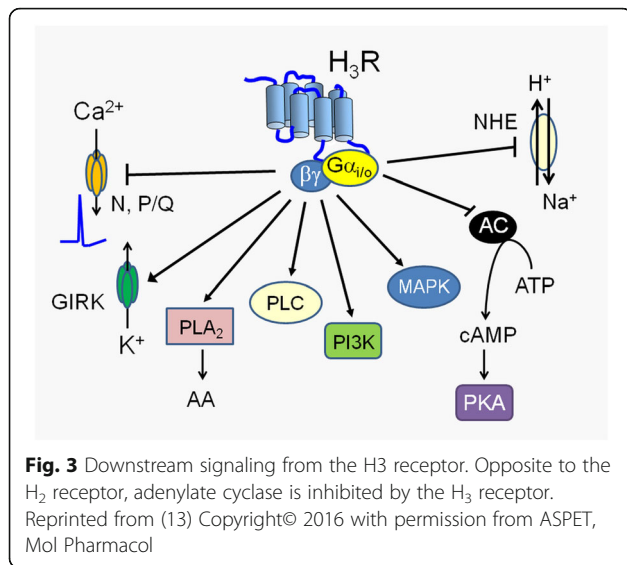
transcription polymerase chain reaction of H<sub>3</sub> receptor mRNA. Countless studies report expression of the H<sub>3</sub> receptor in the brain of rodents, guinea pigs, monkeys and humans. Thus there is good evidence that histamine H<sub>3</sub> receptor is present in many important brain structures e.g. cortex, cerebellum, dorsal thalamus, hypothalamus, amygdala, hippocampus, striatum and basal ganglia. The receptor is primarily located as a presynaptic autoreceptor on histaminergic neurons (on dendrites, axons and somata), which are localized in the tuberomammillary nucleus in the posterior hypothalamus of rodents and humans, or as a heteroreceptor in non-histaminergic neurons. There is some evidence for a postsynaptic H<sub>3</sub> receptor in the CNS as well as a location in the PNS (Fig. 2) [3, 51, 52].

The H<sub>3</sub> receptor alone has been reviewed recently [52] and alongside the H<sub>1</sub>, H<sub>2</sub> and H<sub>4</sub> receptors [3]. The histamine H<sub>3</sub> receptor has multiple signal transduction mechanisms. The receptor couples to G<sub>i/o</sub> protein leading to inhibition of adenylate cyclase (AC) and a reduction of cytosolic cAMP (the opposite action of the H<sub>2</sub> receptor which stimulates AC) is one of the main signaling mechanisms (Fig. 3). The H<sub>3</sub> receptor also signals by activation of mitogen-activated protein kinase (MAPK), by the Akt/PKB signaling pathway and via activation of phospholipase C (PLC) leading to an increase in cytosolic calcium with the latter leading to an inhibition of histamine synthesis and release, hence the auto-receptor function [3].

The numerous functions of the histamine H<sub>3</sub> receptor reflect its widespread localization in the central and peripheral nervous systems as well as interactions with - and

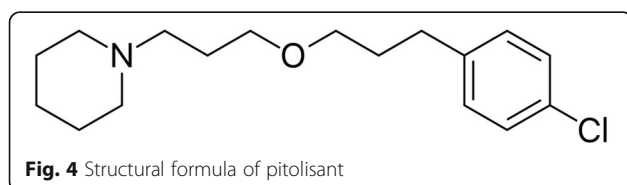






influence by - other neurotransmitters and neuropeptides. Many functions are still hypothetical or controversial, particularly the effects *in vivo*. In general, the receptor mediates the function of histamine itself in the nervous system. The autoreceptor inhibits the release of histamine from histaminergic neurons and this receptor has a significant constitutive activity which refers to the ability of signaling independent of agonist action. The presynaptically located H<sub>3</sub> receptor regulates the release of histamine and other neurotransmitters (serotonin, GABA, glutamate, noradrenaline and possibly acetylcholine and dopamine) and neuropeptides in the CNS and PNS (Fig. 4). This links the histamine H<sub>3</sub> receptor to several neurological and psychiatric disorders with H<sub>3</sub> agonists being potential treatments in Alzheimer's and Parkinson's disease, schizophrenia, sleep disorders and addiction [52]. H<sub>3</sub> agonists might also perhaps be effective in migraine.

The role of histamine in neurogenic inflammation has been reviewed by Rosa and Fantozzi who described an interaction of mast cells and afferent nerve fibers. Mast cell degranulation is stimulated by release of neuropeptides (e.g. CGRP, substance P and VIP) from the nerve endings [16, 53] and is inhibited by histamine via the H<sub>3</sub> receptor [54]. The degranulation of mast cells promotes an inflammatory response including neurogenic inflammation. This leads to a release of inflammatory mediators by afferent neurons which in turn stimulate other mast



cells to degranulate and thereby sustain and prolong the inflammation. There is evidence that mast cell degranulation activates meningeal nociceptors and it is possible that mast cell degranulation has a role in migraine [13, 33, 53, 55]. A potential therapeutic target would be a receptor with a stabilizing function on mast cells preventing or minimizing the degranulation and secondly the neurogenic inflammation. Another possible target is a receptor located on the nerve terminals inhibiting the release of neuropeptides. The H<sub>3</sub> receptor agonist R<sup>α</sup>-methylhistamine inhibited plasma protein extravasation following capsaicin or electrical stimulation of the trigeminal nerve [56]. This suggests that neuropeptide release from sensory C fibers in the rat meninges is at least partially controlled by H<sub>3</sub> receptors. Better knowledge of these mechanisms in neurogenic inflammation could create a possibility of intervening with receptor specific drugs.

Histamine reduces nociceptive transmission when injected into the cerebrospinal fluid or microinjected in brain structures. Central H<sub>1</sub> and H<sub>2</sub> receptors are thought to mediate this effect, but theoretically H<sub>3</sub> receptors could also be involved. An antagonist effect on the H<sub>3</sub> receptor leads to an increase of brain histamine and several basic studies of acute and chronic pain tests have investigated H<sub>3</sub> antagonism as a modifier of nociceptive thresholds. Investigation of thioperamide, an H<sub>3</sub> receptor inverse agonist, reduces nociception in rodents, but the effect is limited and shows a biphasic response with decreasing effects at highest doses [57].

A single research group investigated doses of 1–10 ng histamine injected subcutaneously twice a week as migraine treatment [58] and subsequently compared it with topiramate [59] in patients with refractory migraine with positive results. Their hypothesis was that histamine acts as an agonist on H<sub>3</sub> receptors on mast cells thereby reducing the neurogenic inflammation. The group continued with an investigation of the H<sub>3</sub> agonist N<sup>α</sup>-methylhistamine in migraine prophylaxis. They published their results from phase I and II studies with 30 healthy volunteers and 17 patients with migraine without aura and 1 patient with migraine with aura. Doses of 1–3 ng subcutaneous twice a week proved effective in migraine patients with a reduction of frequency, intensity and duration of the attacks. Adverse effects were intense headache with doses above 10 ng and 4 ng in healthy volunteers and migraine patients respectively. The authors attributed this to an increased H<sub>1</sub> agonistic effect [60]. A double blind placebo controlled phase III study was performed and results were published in 2006. Sixty patients with migraine (50 migraine without aura and 10 migraine with aura) were included. Two from the placebo group and 3 patients from the N<sup>α</sup> group left the study and 55 patients completed the 12 week period. The study confirmed the results seen in the phase II trial

along with lower intake of rescue analgesics in the  $N^{\alpha}$ -group vs. the placebo group and without any adverse events. The hypothesis in the study was that low doses of a  $H_3$  agonist modulates the interaction between mast cells and C-fiber endings and by reducing histamine release lowering the neurogenic inflammation [61]. The doses used in these studies are unusually low making it difficult to understand the mechanism of action or to even accept the results of such minute doses. Furthermore,  $N^{\alpha}$ -methylhistamine does not cross the blood brain barrier. Apparently, no filing for registration of  $N^{\alpha}$ -methylhistamine has been done, so industry is not convinced of these results.

The only  $H_3$  receptor agent approved for medical use is pitolisant, an  $H_3$  inverse agonist. Pitolisant has recently been discovered to be an efficient treatment of narcolepsy and marketed by Bioprojet Pharma as Wakix. The mechanism behind is increased histamine in the brain leading to an increased wakefulness. One of the pivotal studies described headache as an adverse event in 35% of the participants [62]. In a review by Schwartz of the first clinical trials with pitolisant in narcolepsy and Parkinson's disease, he also describes headache as an adverse event of pitolisant. Overall in these studies 9.7% developed headache on pitolisant compared to 2.9% on placebo [63]. Pitolisant readily crosses the blood brain barrier [3] but histamine is known to lack this ability. It remains unclear if this headache observed in the pitolisant trials is due to central or peripheral mechanisms but it is interesting that a drug which promotes histamine release is linked to an increased susceptibility of headache.

The discovery of the histamine  $H_4$  receptor was reported by several independent research groups around the turn of the millennium. The  $H_4$  receptor has ~40% homology with the  $H_3$  receptor. Two additional inactive isoforms have been identified [3]. The  $H_4$  receptor is a coupled to  $G_i/G_o$  protein and uses intracellular  $Ca^{2+}$  as the primary second messenger. In recombinant systems cAMP increase or decrease has been coupled to an agonist response and inverse agonist response, respectively, indicating mediation by adenylate cyclase. There are probably additional signaling mechanisms and supporting evidence is emerging. The localization of the  $H_4$  receptor is complicated by deficiency of validated  $H_4$  selective antibodies to identify the  $H_4$  receptor expression. Consensus is that  $H_4$  receptors are present in the bone marrow and hematopoietic cells.  $H_4$  receptor expression was also shown in sensory neurons innervating skin in mice but there is no evidence for its presence in human sensory neurons yet. Pharmacological methods have shown that the  $H_4$  receptor is expressed on mast cells [3] and there is some evidence that microglial cells express  $H_4$  receptors [64].

The literature describes an immune modulating function of the  $H_4$  receptor with an involvement in inflammatory responses. It mediates chemotaxis and is involved in cytokine release from various immune cells. Coruzzi et al. suggested anti-inflammatory and antinociceptive effects of  $H_4$  antagonists in a rat model [65], a characteristic shared with the  $H_3$  receptor [3]. The  $H_4$  receptor has not yet been associated with migraine and further description of its localization and function is needed.

### Conclusions and future perspectives

Histamine receptors  $H_1$ ,  $H_2$ ,  $H_3$  and  $H_4$  are widespread throughout the body but there is limited knowledge about the  $H_4$  receptor.  $H_3$  is an autoreceptor, inhibiting release of histamine where  $H_1$  and  $H_2$  have pure agonistic effects and have effect on multiple sites. Modern provocation studies with histamine have increased the interest in the histaminergic system in relation to migraine because histamine proved to be an effective migraine provoking agent. Studies of the effect of antihistamines in migraine are limited and of poor quality. Thus, there is little support for  $H_1$  or  $H_2$  antihistamines as preventive migraine medication. In addition, there are undesirable adverse effects. Still, non-sedating  $H_1$  antihistamines need to be appropriately tested. This has not been done and it might provide valuable information of histamine's site of action and its relation to migraine. There are positive reports about the  $H_3$  receptor agonist  $N^{\alpha}$ -methylhistamine as migraine treatment, but the use of almost homeopathic doses makes the results less credible. Additional exploration of the histaminergic system, peripheral and central, could reveal further goals for migraine therapy.

### Abbreviations

AC: Adenylate cyclase; B.i.d.: Two times a day; CGRP: Calcitonin gene-related peptide; GTN: Glyceryl trinitrate; MAPK: Mitogen-activated protein kinase; NOS: Nitric oxide synthase; PLC: Phospholipase C; Q.i.d.: Four times a day; T.i.d.: Three times a day; VIP: Vasoactive intestinal peptide

### Acknowledgements

Not applicable.

### Funding

The authors received no financial support for the research, authorship, and/or publication of this article.

### Availability of data and materials

Not applicable.

### Authors' contributions

JW and JO conceived and designed the review. JW conducted the literature search and drafted the manuscript. JO and KF contributed to the manuscript and all authors revised the manuscript for intellectual content. All authors read and approved the final manuscript.

### Ethics approval and consent to participate

Not applicable.

### Consent for publication

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

**Publisher's Note**

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Received: 20 January 2019 Accepted: 18 March 2019

Published online: 25 March 2019

**References**

1. Simons FER, Simons KJ (2011) Histamine and H1-antihistamines: celebrating a century of progress. *J Allergy Clin Immunol* 128:1139–1150.e4
2. Alstadhaug KB (2014) Histamine in migraine and brain. *Headache J Head Face Pain* 54:246–259
3. Panula P, Chazot PL, Cowart M, Gutzmer R, Leurs R, Liu WLS, Stark H, Thurmond RL, Haas HL (2015) International Union of Basic and Clinical Pharmacology. XC VIII. Histamine receptors. *Pharmacol Rev* 67:601–655
4. Yuan H, Silberstein SD (2017) Histamine and migraine. *Headache J Head Face Pain*. <https://doi.org/10.1111/head.13164>
5. Dale HH, Laidlaw PP (1910) The physiological action of  $\beta$ -iminazolyethylamine. *J Physiol* 41:318–344
6. Harmer JM, Harris KE (1926) Observations on the vascular reactions in man in response to histamine. *Heart* 13:381
7. Pickering GW, Hess W (1933) Observations on the mechanism of headache produced by histamine. *Clin Sci* 1:77–101
8. Northfield DWC (1948) Some observations on headache. *Brain* 61:133–162
9. Schumacher GA, Wolff HG (1941) Experimental studies on headache: A. Contrast of histamine headache with the headache of migraine and that associated with hypertension B. Contrast of vascular mechanisms in preheadache and in headache phenomena of migraine. *Arch Neurol Psychiatr* 45:199–214
10. Storch TJC (1940) Relation of experimental histamine headache to migraine and nonmigraine headache. *Arch Neurol Psychiatr* 44:316–322
11. Macy D, Horton BT (1948) TREATMENT OF MIGRAINE WITH HISTAMINE: review of one hundred and forty-four cases. *J Am Med Assoc* 137:1110–1114
12. Thomas WA, Butler S (1946) Treatment of migraine by intravenous histamine. *Am J Med* 1:39–44
13. Schwenger N, Dux M, de Col R, Carr R, Messlinger K (2007) Interaction of calcitonin gene-related peptide, nitric oxide and histamine release in neurogenic blood flow and afferent activation in the rat cranial dura mater. *Cephalalgia* 27:481–491
14. Akerman S, Williamson DJ, Kaube H, Goadsby PJ (2002) The role of histamine in dural vessel dilation. *Brain Res* 956:96–102
15. De Logu F, Landini L, Janal MN, Li Puma S, De Cesaris F, Geppetti P, Nassini R (2019) Migraine-provoking substances evoke periorbital allodynia in mice. *J Headache Pain*. <https://doi.org/10.1186/s10194-019-0968-1>
16. Ramachandran R (2018) Neurogenic inflammation and its role in migraine. *Semin Immunopathol* 40:301–314
17. Katzung BG, Masters SB, Trevor AJ (eds) (2012) Basic & clinical pharmacology, 12th edn. McGraw-Hill Medical, New York
18. Curran DA, Lance JW (1964) Clinical trial of methysergide and other preparations in the management of migraine. *J Neurol Neurosurg Psychiatry* 27:463–469
19. Sicuteri F, Franchi G, Bianco PLD (1967) An Antaminic drug, BC 105, in the prophylaxis of migraine. *Int Arch Allergy Immunol* 31:78–93
20. Selby G (1970) A clinical trial of an antiserotonin drug, bc-105 in the prophylaxis of migraine. *Proc Aust Assoc Neurol* 7:37–43
21. Lance JW, Anthony M, Somerville B (1970) Comparative trial of serotonin antagonists in the Management of Migraine. *Br Med J* 2:327–330
22. Osterman PO (1977) A comparison between placebo, Pizotifen and 1-Isopropyl-3-Hydroxy-5-Semicarbazono-6-Oxo-2,3,5,6-Tetrahydroindol (divascan®) in migraine prophylaxis. *Acta Neurol Scand* 56:17–28
23. Lawrence ER, Hossain M, Littlestone W (1977) Sanomigran for migraine prophylaxis; controlled multicenter trial in general practice. *Headache J Head Face Pain* 17:109–112
24. Anthony M, GDA L, Lance JW (1978) Controlled trials of cimetidine in migraine and cluster headache. *Headache J Head Face Pain* 18:261–264
25. International Headache Society Clinical Trials Subcommittee members, Tfelt-Hansen P, Pascual J et al (2012) Guidelines for controlled trials of drugs in migraine: third edition. A guide for investigators. *Cephalalgia* 32:6–38
26. Nanda RN, Arthur GP, Johnson RH, Lambie DG (1980) Cimetidine in the prophylaxis of migraine. *Acta Neurol Scand* 62:90–95
27. Togha M, Ashrafi H, Tajik P (2006) Open-label trial of Cinnarizine in migraine prophylaxis. *Headache J Head Face Pain* 46:498–502
28. Rossi P, Fiermonte G, Pierelli F (2003) Cinnarizine in migraine prophylaxis: efficacy, tolerability and predictive factors for therapeutic responsiveness. An open-label pilot trial. *Funct Neurol* 18:155–160
29. Bostani A, Rajabi A, Moradian N, Razavian N, Rezaei M (2013) The effects of cinnarizine versus sodium valproate in migraine prophylaxis. *Int J Neurosci* 123:487–493
30. Ashrafi MR, Salehi S, Malamiri RA, Heidari M, Hosseini SA, Samiei M, Tavasoli AR, Togha M (2014) Efficacy and safety of Cinnarizine in the prophylaxis of migraine in children: a double-blind placebo-controlled randomized trial. *Pediatr Neurol* 51:503–508
31. Lipton RB, Bigal ME, Steiner TJ, Silberstein SD, Olesen J (2004) Classification of primary headaches. *Neurology* 63:427–435
32. Dussor G (2014) Serotonin, 5HT1 agonists, and migraine: new data, but old questions still not answered. *Curr Opin Support Palliat Care* 8:137–142
33. Kilinc E, Guerrero-Toro C, Zakharov A et al (2017) Serotonergic mechanisms of trigeminal meningeal nociception: implications for migraine pain. *Neuropharmacology* 116:160–173
34. Olesen J (1978) Some clinical features of the acute migraine attack. An analysis of 750 patients. *Headache J Head Face Pain* 18:268–271
35. Classification and diagnostic criteria for headache disorders, cranial neuralgias and facial pain (1988). Headache Classification Committee of the International Headache Society. *Cephalalgia Int J Headache* 8 Suppl 7:1–96
36. Krabbe AA, Olesen J (1980) Headache provocation by continuous intravenous infusion of histamine. Clinical results and receptor mechanisms. *Pain* 8:253–259
37. Krabbe AÆ, Olesen J (1982) Effect of histamine on regional cerebral blood flow in man. *Cephalalgia* 2:15–18
38. Edvinsson L, Owman C (1975) A pharmacologic comparison of histamine receptors in isolated extracranial and intracranial arteries in vitro. *Neurology* 25:271–276
39. Lassen LH, Thomsen LL, Olesen J (1995) Histamine induces migraine via the H1-receptor. Support for the NO hypothesis of migraine. *Neuroreport* 6:1475–1479
40. Thomsen LL, Kruuse C, Iversen HK, Olesen J (1994) A nitric oxide donor (nitroglycerin) triggers genuine migraine attacks. *Eur J Neurol* 1:73–80
41. Lassen LH, Thomsen LL, Kruuse C, Iversen HK, Olesen J (1996) Histamine-1 receptor blockade does not prevent nitroglycerin induced migraine. *Eur J Clin Pharmacol* 49:335–339
42. Schmetterer L, Wolzt M, Grasel U, Findl O, Strenn K, Simak S, Kastner J, Eichler H-G, Singer EA (1997) Nitric oxide synthase inhibition in the histamine headache model. *Cephalalgia* 17:175–182
43. Lassen L, Christiansen I, Iversen H, Jansen-Olesen I, Olesen J (2003) The effect of nitric oxide synthase inhibition on histamine induced headache and arterial dilatation in Migraineurs. *Cephalalgia* 23:877–886
44. Lassen LH, Ashina M, Christiansen I, Ulrich V, Olesen J (1997) Nitric oxide synthase inhibition in migraine. *Lancet* 349:401–402
45. Lassen L, Ashina M, Christiansen I, Ulrich V, Grover R, Donaldson J, Olesen J (1998) Nitric oxide synthase inhibition: a new principle in the treatment of migraine attacks. *Cephalalgia* 18:27–32
46. Lassen LH, Heinig JH, Oestergaard S, Olesen J (1996) Histamine inhalation is a specific but insensitive laboratory test for migraine. *Cephalalgia* 16:550–553
47. Haimart M, Pradalier A, Launay J, Dreux C, Dry J (1987) Whole blood and plasma histamine in common migraine. *Cephalalgia* 7:39–42
48. Heatley RV, Denburg JA, Bayer N, Bienenstock J (1982) Increased plasma histamine levels in migraine patients. *Clin Allergy J Br Allergy Soc* 12:145–149
49. Arrang J-M, Garbarg M, Schwartz J-C (1983) Auto-inhibition of brain histamine release mediated by a novel class (H3) of histamine receptor. *Nature* 302:832–837
50. Lovenberg TW, Roland BL, Wilson SJ, Jiang X, Pyati J, Huvar A, Jackson MR, Erlander MG (1999) Cloning and functional expression of the human histamine H3 receptor. *Mol Pharmacol* 55:1101–1107
51. Berlin M, Boyce CW, de Lera RM (2010) Histamine H3 receptor as a drug discovery target. *J Med Chem* 54:26–53
52. Nieto-Alamillo G, Márquez-Gómez R, García-Gálvez A-M, Morales-Figueroa G-E, Arias-Montañón J-A (2016) The histamine H3 receptor: structure, pharmacology, and function. *Mol Pharmacol* 90:649–673
53. Rosa AC, Fantozzi R (2013) The role of histamine in neurogenic inflammation. *Br J Pharmacol* 170:38–45

54. Rozniecki JJ, Letourneau R, Sugiultzoglu M, Spanos C, Gorbach J, Theoharides TC (1999) Differential effect of histamine 3 receptor-active agents on brain, but not peritoneal, Mast Cell Activation. *J Pharmacol Exp Ther* 290:1427–1435
55. Levy D (2009) Migraine pain, meningeal inflammation, and mast cells. *Curr Pain Headache Rep* 13:237–240
56. Toshio M, Moskowitz MA, Zhihong H (1992) UK-14,304, R (-)- $\alpha$ -methyl-histamine and SMS 201-995 block plasma protein leakage within dura mater by prejunctional mechanisms. *Eur J Pharmacol* 224:145–150
57. Hough LB, Rice FL (2011) H3 receptors and pain modulation: peripheral, spinal, and brain interactions. *J Pharmacol Exp Ther* 336:30–37
58. Millán-Guerrero RO, Cárdenas MAI, Ocampo AA, Pacheco MF (1999) Histamine as a therapeutic alternative in migraine prophylaxis: a randomized, placebo-controlled, double-blind study. *Headache J Head Face Pain* 39:576–580
59. Millán-Guerrero RO, Isais-Millán R, Barreto-Vizcaíno S, Gutiérrez I, Rivera-Castaño L, Trujillo-Hernández B, Baltazar LM (2008) Subcutaneous histamine versus Topiramate in migraine prophylaxis: a double-blind study. *Eur Neurol* 59:237–242
60. Millán-Guerrero RO, Pineda-Lucatero AG, Hernández-Benjamín T, Tene CE, Pacheco MF (2003) N $\alpha$ -methylhistamine safety and efficacy in migraine prophylaxis: phase I and phase II studies. *Headache J Head Face Pain* 43:389–394
61. Millán-Guerrero RO, Isais-Millán R, Benjamín T-H, Tene CE (2006) N $\alpha$ -methyl histamine safety and efficacy in migraine prophylaxis: phase III study. *Can J Neurol Sci J Can Sci Neurol* 33:195–199
62. Dauvilliers Y, Bassetti C, Lammers GJ, Arnulf I, Mayer G, Rodenbeck A, Lehert P, Ding C-L, Lecomte J-M, Schwartz J-C (2013) Pitolisant versus placebo or modafinil in patients with narcolepsy: a double-blind, randomised trial. *Lancet Neurol* 12:1068–1075
63. Schwartz J-C (2011) The histamine H3 receptor: from discovery to clinical trials with pitolisant. *Br J Pharmacol* 163:713–721
64. Schneider EH, Seifert R (2016) The histamine H4-receptor and the central and peripheral nervous system: a critical analysis of the literature. *Neuropharmacology* 106:116–128
65. Coruzzi G, Adami M, Guaita E, de Esch IJP, Leurs R (2007) Antiinflammatory and antinociceptive effects of the selective histamine H4-receptor antagonists JNJ777120 and VUF6002 in a rat model of carrageenan-induced acute inflammation. *Eur J Pharmacol* 563:240–244

**Ready to submit your research? Choose BMC and benefit from:**

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

**At BMC, research is always in progress.**

Learn more [biomedcentral.com/submissions](https://biomedcentral.com/submissions)

