Histamine as a Growth Factor and Chemoattractant for Human Carcinoma and Melanoma Cells: Action through Ca\(^{2+}\)-Mobilizing H\(_1\) Receptors

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Abstract. Histamine receptors are present on the surface of various normal and tumor-derived cell types, where their biological function is incompletely understood. Here we report that histamine not only stimulates cell proliferation under serum-free conditions, but also is chemoattractive for human carcinoma (Hela and A431) and melanoma (A875) cells expressing H\(_1\) type receptors. Histamine was found to be a potent activator of phospholipase C, leading to polyphosphoinositide hydrolysis and subsequent intracellular Ca\(^{2+}\) mobilization. In addition, histamine also causes the protein kinase C-mediated activation of Na\(^+\)/H\(^+\) exchange, as evidenced by an amiloride-sensitive rise in cytoplasmic pH. All histamine-induced responses, including chemotaxis and DNA synthesis, are completely inhibited by the H\(_1\) receptor antagonist pyrilamine, but not by cimetidine, an inhibitor of histamine H\(_2\) type receptors. Our results suggest that histamine may have a previously unrecognized role in the migration and proliferation of cells expressing H\(_1\) receptors.

Histamine is a widely occurring chemical mediator that has long been known as a neurotransmitter, inflammatory factor, and a modulator of gastrointestinal functions (10, 24). It has also been suggested that histamine is involved in certain types of cell proliferation in vivo, such as wound healing, embryonic development, and tumor growth (for review, see reference 2). There are several reports that elevated levels of histamine and of its synthesizing enzyme are associated with rapid tissue growth (17), but the precise link, if any, between histamine and cell proliferation remains obscure.

Three types of histamine receptors (H\(_1\), H\(_2\), and H\(_3\)) have been identified that differ in their sensitivity to antagonists and mediate different actions (14). While activation of H\(_2\) type receptors leads to an increase in cAMP (12), the H\(_1\) type receptor is linked to phosphoinositide breakdown and subsequent intracellular Ca\(^{2+}\) mobilization (8, 9, 23) and, therefore, is an attractive candidate for mediating growth stimulation.

Here we report that exogenously added histamine not only stimulates DNA synthesis and cell division but also evokes a chemotactic response in human Hela and A431 carcinoma cells and A875 melanoma cells. We show that these novel actions of histamine are mediated by the H\(_1\) type receptor that triggers the hydrolysis of phosphoinositides with consequent formation of various second messengers. Our results suggest that histamine may have a novel role in the migration and proliferation of H\(_1\) receptor-bearing (tumor) cells.

Materials and Methods

Materials

BSA, cimetidine, DiC8, histamine, and pyrilamine were purchased from Sigma Chemical Co. (St. Louis, MO). Other reagents were from the following sources: Bis(carboxy-ethyl)carboxyfluorescein (BCECF) from Molecular Probes (Eugene, OR); epidermal growth factor (EGF), receptor grade, from Collaborative Research, Inc. (Waltham, MA); FCS from Hyclone Laboratories (Logan, UT); Hoechst 33258 from Boehringer Mannheim GmbH (Mannheim, FRG); indol-1 acetoxymethylxanthine from Molecular Probes (Eugene, OR) and polycarbonate filters from Nuclepore Corp. (Pleasanton, CA). Myo[2-3H]inositol (12.3 Ci/mmol) and cAMP assay kits were obtained from Amersham International (Amersham, UK).

Cell Culture

Human A431 and Hela carcinoma and A875 melanoma cells were routinely grown at 37°C in DME containing 7.5% (vol/vol) FCS at 5% CO\(_2\).

Cell Counting

Hela, A431, and A875 cells were seeded at a density of 10\(^{3}\)/cm\(^2\) (Hela and A431) or 5 · 10\(^{2}\)/cm\(^2\) (A875) in DME containing 7.5% (vol/vol) fetal calf serum and allowed to attach for 24 h. After shifting the cultures to serum-free conditions, the cells were harvested and counted using a Coulter counter.

Abbreviations used in this paper: BCECF, bis(carboxy-ethyl)carboxyfluorescein; EGF, epidermal growth factor; IMBX, isobutylmethylxanthine.

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DNA Content

Cells were seeded and allowed to attach in DME. 24 h before growth factor addition, the cultures were shifted to serum-free medium consisting of DME/Ham's F-12 (1:1; vol:vol), supplemented with 10 μg/ml transferrin. After 2 d of hormonal stimulation, a cell lysate was prepared and total DNA content was measured using the nuclear dye Hoechst 33258 (18).

Determination of Inositol Phosphates

Nearly confluent cultures were shifted to serum-free DME/Ham's F-12 medium, containing 10 μg/ml transferrin and 2 μCi [3H]inositol for 24 h. After stimulation, an inositol phosphate containing fraction was prepared and quantitated by anion exchange chromatography as described previously (27). HPLC elution profiles were obtained by analyzing inositol phosphates containing fractions on a Partisil SAX column, eluted with a gradient to 1.5 M ammonium formate/phosphoric acid (pH, 3.7). [3H]radioactivity was determined by on-line scintillation counting using a radioactivity monitor (Berthold LB506C; Betrow, Rotterdam, the Netherlands).

cAMP Measurements

Nearly confluent cells in 6-well tissue culture dishes were preincubated for 2 h in serum-free DME containing 20 mM Hepes (pH, 7.6). Thereafter, cells were exposed to 1 μM isobutylnethylnitrate (IBN) for 10 min, then to agonists. The reactions were stopped by adding 10% (wt/vol) ice-cold TCA. After centrifugation, the supernatants were extracted with diethylether to remove TCA and neutralized with Tris-base. Cellular content of cAMP was determined using the [3H]cAMP assay kit from Amersham International according to the instructions of the manufacturer.

Ionic Responses

Nearly confluent cultures, attached to glass coverslips and maintained in serum-free DME/Ham's F-12 (1:1; vol:vol) medium containing 10 μg/ml transferrin for 24 h, were loaded with the fluorescent indicators indo-1 (for Ca2+) or BCECF (for pH). Fluorescence was recorded and calibrated as described previously (21).

Chemotactic Assay

Chemotactic assays were carried out in modified Boyden chambers (11), equipped with gelatin-coated filters from Nuclepore Corp. (8 μm pore size). Cells, resuspended to a density of 10⁶ cells/ml, were seeded into the upper chamber, while a histamine-containing agar solution (5%) was put in the lower chamber. All incubations were performed in DME/Ham's F-12 (1:1, vol:vol) medium containing 10 μg/ml transferrin and 1 mg/ml BSA. The results are expressed as number of cells migrated through the membrane per microscopic field (400×) during a 12-h period.

Results

Mitogenesis

In an initial screening, we found Ca2+-mobilizing H1 receptors to be present on various human tumor cell lines, including Hela, A431, epidermoid carcinoma, HT-29, colon carcinoma, and A875, melanoma cells, but not on MCF-7 breast carcinoma cells. No functional H2 receptors were found on mouse NIH and Swiss 3T3 cells or on rat-l fibroblasts. The Hela carcinoma cell line constitutes a convenient model for studying potential effects of histamine on cell proliferation, since these cells can be grown under serum-free conditions while they remain growth factor-responsive (15). When kept in a serum-free medium containing transferrin, Hela cells remain viable with a mean population doubling time of 72 h.

Addition of histamine to such serum-free cultures was found to stimulate DNA synthesis and cell division in a dose-dependent manner, half-maximal effects being observed at 10-15 μM and a saturating response at 5 x 10⁻⁴ M. As shown in Table I and Fig. 1 histamine is somewhat less potent as EGF in stimulating Hela cell growth, with histamine (10⁻⁴ M) decreasing generation time from 72 to 31 h and EGF (50 ng/ml) to 19 h. Insulin (5 μg/ml) was equally potent as histamine (Table I), whereas mitogenic peptides such as bombesin, bradykinin, and substance P, had no effect on Hela cell proliferation (our unpublished observations). When added together, histamine and insulin evoke an additive rather than a synergistic proliferative response (not shown). The mitogenic activity of histamine is not restricted to Hela cells. Also in A431 cells and A875 melanoma cells histamine exerts a marked mitogenic effect (Table I).

The growth stimulatory action of histamine appears to be mediated by the H1 type receptor, since the H2 antagonist pyrilamine completely blocks histamine-induced DNA synthesis, whereas the H2 antagonist cimetidine, even at millimolar concentrations, has no detectable effect (Fig. 1). Importantly, these antagonists affect neither the basal rate nor the EGF- and/or insulin-stimulated rate of DNA synthesis (Fig. 1).

Involvement of Phospholipase C

As illustrated in Fig. 2, addition of histamine to Hela cells evokes the phospholipase C-mediated breakdown of polyphosphoinositides which result in rapid formation of inositol

Table I. Growth Stimulation of Human Carcinoma and Melanoma Cells by Histamine

|                   | Hela | A431 | A875 | Hela | A431 | A875 |
|-------------------|------|------|------|------|------|------|
| Control           | 16 ± 1 | 19 ± 1 | 10 ± 1 | 72 | 50 | 56 |
| Histamine (100 μM) | 26 ± 2 | 30 ± 2 | 16 ± 1 | 31 | 29 | 36 |
| Insulin (5 μg/ml)  | 26 ± 2 | 28 ± 1 | 16 ± 1 | 31 | 30 | 36 |
| EGF (50 ng/ml)    | 51 ± 2 | ND | 11 ± 1 | 19 | ND | 48 |

Cells were seeded and allowed to attach for 24 h before growth stimulation. 24-48 h after addition of the various growth factors, cells were resuspended and counted. Data are expressed as means ± SEM for triplicate experiments.

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Figure 2. Accumulation of inositol phosphates evoked by histamine in Hela cells. A, time course of histamine-induced inositol phosphate accumulation in Li+-treated cultures. [3H]-inositol prelabeled cultures were incubated with histamine (100 μM) for the indicated times in the presence of 10 mM LiCl. Thereafter, an inositol phosphate containing fraction was prepared and analyzed as described under Materials and Methods. Inset shows the effects of pyrilamine (1 μM, hatched bars and cimetidine; 1 mM, dotted bars) on the histamine-induced inositol phosphate production (30-min incubations). Data are expressed as means ± SEM for triplicate incubations. B, HPLC profiles of [3H]inositol phosphates obtained from control and histamine-stimulated cultures (100 μM during 10 s).

Analysis of the various inositol phosphates formed (Fig. 2 B), reveals a rapid, several-fold increase in Ins(1,4,5)P3, the second messenger for Ca2+ mobilization (5), and other inositol polyphosphates. In addition to generating inositol phosphates, histamine transiently stimulates the formation of diacylglycerol in [2-3H]glycerol-labeled cells, as revealed by using conventional TLC (Tilly, B. C., L. G. J. Tertoolen, A. C. Lambrechts, R. Remorie, S. W. de Laat, and W. H. Moolenaar, manuscript submitted for publication), which binds to and directly activates protein kinase C.

Ionic Responses

The activation of phospholipase C by histamine is expected to lead to increases in cytoplasmic free Ca2+ ([Ca2+]i) and pH (pH), because of Ins(1,4,5)P3-mediated Ca2+ release (5) and protein kinase C–activated Na+/H+ exchange (20), respectively. Fig. 3 shows typical ionic responses to histamine. Histamine elicits a rapid, but transient, biphasic rise in [Ca2+]i, which approaches micromolar Ca2+ levels at 10–20 s after hormone addition and which is completely blocked by pyrilamine (Fig. 3 A). The initial rise in [Ca2+]i rise is largely, if not entirely, because of release of intracellular stored Ca2+, while the second phase requires Ca2+ influx across the plasma membrane (Tilly, B. C., L. G. J. Tertoolen, A. C. Lambrechts, R. Remorie, S. W. de Laat, and W. H. Moolenaar, manuscript in preparation).

Histamine also causes the protein kinase C–mediated activation of Na+/H+ exchange, as shown by a rise in pH, of ~0.20 U, that is abolished in the presence of the Na+/H+ exchange inhibitor amiloride (0.5 mM), while subsequent addition of cell permeable diacylglycerol (diC8) does not cause an additional elevation of pH, (Fig. 3 B). Similar ionic responses to histamine were observed in A431 and A875 cells.

Independence of Adenylate Cyclase

Although the mitogenic effects of Ca2+ mobilizing hormones are thought to proceed via the phospholipase C signal transduction pathway, recent evidence suggests that inhibition of adenylate cyclase (through a receptor-linked inhibitory G protein; Gi) is actually more important for mitogenesis (26). We determined the effects of histamine on
intracellular levels of adenosine 3',5'-monophosphate (cAMP) following stimulation of adenylate cyclase by isoproterenol (10 μM); the phosphodiesterase inhibitor IMBX (0.5 mM) was included to ensure that changes in cAMP levels were not mediated by phosphodiesterase action. Under conditions where exogenous phosphatidic acid (cf. reference 22) inhibits cAMP accumulation up to 60% within 15 min (from 1.23 to 0.30 ± 0.04 pmol CAMP/10^6 cells), histamine completely failed to attenuate isoproterenol-induced cAMP accumulation in Hela cells. Thus, mitogenic signaling through the H1 type receptor does not appear to involve the G protein that inhibits adenylate cyclase.

Chemotaxis

Several agents, including certain growth regulators, promote the chemotactic migration of their target cells (4, 25, 29). This activity is important in wound healing, but chemotaxis may also be involved in tumor cell metastasis (19). We examined whether histamine can function as a chemoattractant for Hela cells using the modified Boyden chamber assay (11), in which cells are allowed to migrate through membrane filters. Fig. 4 shows that histamine evokes a potent chemotactic response in a dose-dependent manner, with a half-maximal effect at ~5 × 10^-6 M. Histamine-induced chemotaxis is completely blocked by pyrilamine (Fig. 4), indicating again the involvement of H1 receptors. A chemotactic response (an approximately twofold increase relative to the control) is also observed after stimulating A875 melanoma cells with histamine (100 μM), although the chemotaxis assay was technically more difficult than with Hela cells because of the tendency of melanoma cells to aggregate when resuspended in a Boyden chamber.

Discussion

The present data clearly demonstrate that histamine, a ubiquitously occurring local hormone, can function as a growth factor and chemoattractant for cells types that express the Ca^{2+}-mobilizing H1 type receptor. We found no evidence that mitogenic signaling through H2 type receptors may involve alterations in cAMP levels. Our results raise the intriguing possibility of a role for histamine in promoting cell growth and migration in vivo. The present findings also suggest that histamine-secreting mast cells may have a more important role in modulating cell proliferation than has been assured to date. Indeed, the finding that mast cells are frequently abundant in the contiguous tissue of metastatic tumors supports this possibility (7).

Interestingly, histamine is not only stored in mast cells but is also actively synthesized in various other cell types. In particular, high levels of histamine and its synthesizing enzyme occur in many tissues undergoing rapid cell growth or tissue repair, including regenerating liver, bone marrow, embryonic tissues, and experimental tumors (1, 10, 17, 6). Although it remains to be seen whether histamine produced by these cells is released into the extracellular space, it is tempting to speculate that newly synthesized histamine, acting via H2 receptors, may contribute to cell proliferation and development by serving as an autocrine growth and motility factor.

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