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THE CENTRAL EFFECT OF BIOLOGICAL AMINES ON IMMUNOSUPPRESSIVE EFFECT OF RESTRAINT STRESS IN RAT

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Abstract - The effects of some histaminergic agents were evaluated on stress-induced immunosuppression in immunized male rats. In rats immunized with sheep red blood cells (SRBC's), restraint stress (RS) prevented the booster-induced rise in anti-SRBC antibody titre and cell immunity response. Intracerebroventricular (ICV) injection of histamine (150 μg/kg) induced a similar effect with RS. Pretreatment with cromolyn sodium (50 μg/kg) reduced the inhibitory effect of RS on immune function. Also histamine could inhibit the effect of cromolyn sodium when injected simultaneously. Pretreatment with ranitidine (10 μg/kg) had not a significant effect. Serotonin (5 μg/kg) and dopamine (6.2 μg/kg) could reverse the effect of cromolyn sodium when injected with cromolyn sodium (100 μg/kg). Epinephrine (6.2 μg/kg) had not a significant effect. The results indicated that histamine mediates the immunosuppression of restrained stress by influencing the histaminergic H2 receptor in the brain and this effect of histamine may be mediated by serotonergic and dopaminergic system. Acta Medica Iranica 38 (3): 182-186, 2000

Key Words: Restraint stress, histamine, immunity

INTRODUCTION

Psychoneuroimmunology is a relatively new discipline which deals with CNS immune system interaction. Two pathways link the brain and the immune system: The autonomic nervous system and neuroendocrine output via the pituitary (1). Different studies have been conducted to understand the clinical implication and mechanisms of the effects of stress on the immune system.

The effect of stress on immune function is related to different factors such as type and duration of particular stressor being used, the specific immune tested, and the time of day that the stressor was applied (2,3,4). The restraint stress for 24h can reduce the humoral and cell mediated immunity (5,6). The role of dopaminergic (6,7,8), sympathetic nervous system (9,9) and different hormones such as growth hormone (21), prolactin (10), corticosteroids and endogenous opioids (11,12,13), and other agents such as substance P and adenosine (14,15) have been studied in immunosuppression related to stress. For example, the release of prolactin and growth hormone increases following restraint stress and induce immunosuppressor effect, but corticosteroids and opioids reduce immune system activity.

Histamine has direct effect on immune cells and reduces activity of this cells via H2 receptors (14,16,17). Also histamine releases during stress in the brain and mediate the activity of different systems (18).

Histaminergic neurons mediate restraint stress-induced activation of 5-hydroxytryptaminergic and dopaminergic neurons (19,20) and the activity of noradrenergic neurons projecting to the hypothalamus (21) in the rats. Restraint stress-induced immunosuppression may be mediated by different mechanisms via some of hormones or nervous system (5,6,9,13,22). The role of histamine receptors in the effect of stress has not been studied.

MATERIALS AND METHODS

Subjects: 200 Male Wistar rats (200-250g) were used. They were housed in standard laboratory conditions of light (12h light-dark schedule) and temperature (22±2°C) and had free access to food and water.

Drugs: The drugs used were histamine, cromolyn sodium, serotonin, dopamine and epinephrine (all from Sigma) and ketamine for anesthesia (Park Davis). Drugs were dissolved diluted in distilled water and were injected intracerebroventricularly (5 μl/ rat) 15 minute prior to restraint stress.

Stress procedure (6) restraint stress (RS) was applied in plexiglas restrainers (9 x 7 x 15 cm)
RESULTS

In rats immunized with SRBCs, a booster dose of antigen clearly augmented the humoral immune response to antigen. RS clearly attenuated the booster (SRBC)-induced rise in the secondary antibody titre (P < 0.01, Fig. 1). Similar changes (suppression) were also seen when histamine (150 µg/rat) was given intracerebroventricularly (I.C.V.) in nonstressed rats as well (Fig. 1).

![Fig. 1. The central effect of histamine on humoral immunity following restraint stress (RS) in rat. The effect of RS, histamine (150 µg/rat) and control (non-stressed) groups were compared on antibody titre in immunized rat with SRBC. Each column shows mean ± SE of antibody titre related to six rats. ** P < 0.01.](www.SID.ir)

Pre-treatment with chlorpheniramine (50 µg/rat) reduced the immunosuppression effect of stress (P < 0.01, Fig. 2).

![Fig. 2. The role of histamine receptors on the reduction of humoral immunity following restraint stress (RS) in rat. Centre: The effect of pre-treatment with chlorpheniramine (chl, 30 µg/rat), chlorpheniramine (chl, 30 µg/rat), and histamine (chl, 150 µg/rat) was compared on antibody titre. Each column shows mean ± SE of antibody titre related to six rats. ** P < 0.01 (compared with control). ** P < 0.01 (compared with group received chlorpheniramine).](www.SID.ir)
Histamine (150 μg/rat) when injected with chlorpheniramine (50 μg/rat) simultaneously inhibited the effect of chlorpheniramine (Fig. 2). Ranitidine (10 μg/rat) did not reduce the inhibitory effect of RS on immune function (Fig. 2). Serotonin (3 μg/rat), dopamine (0.2 μg/rat) and epinephrine (0.2 μg/rat) were injected with chlorpheniramine in different groups of stressed rats.

Fig. 5. The effect of serotonin, dopamine, and epinephrine on increment of antibody titre by chlorpheniramine following restraint stress (RS) in rat. Histamine (chl, 50 μg/rat), chl + serotonin (ser, 3 μg/rat), chl + dopamine (dop, 0.2 μg/rat), chl + epinephrine (epi, 0.2 μg/rat) were compared on antibody titre in stressed rats. Each column shows mean ± SE of antibody titre related to six rats. * P<0.05

Proctreatment with chlorpheniramine (50 μg/rat) reduced the inhibitory effect of RS (P<0.01). Ranitidine (10 μg/rat) did not show a significant response (Fig. 5). Also histamine (150 μg/rat) could inhibit the effect of chlorpheniramine when injected simultaneously (Fig. 5). Chlorpheniramine when injected with serotonin (3 μg/rat) and dopamine (0.2 μg/rat) caused a significant reduction in inhibitory effect of chlorpheniramine on RS-induced immunosuppression (P<0.05 Fig. 6). But epinephrine (0.2 μg/rat) had not a significant effect (Fig. 6).

Fig. 6. The central role of histamine receptors on the reduction of cell immune response following restraint stress (RS) in rat. The effects of pretreatment with ranitidine (ran, 10 μg/rat), chlorpheniramine (chl, 50 μg/rat) and chl + histamine (chl, 150 μg/rat) were compared on difference in paw volume (DIPV) in stressed rat. Each column shows mean ± SE of DIPV related to six rats. ** P<0.01 (compared with control), T P<0.01 (compared with group received chlorpheniramine).

Serotonin and dopamine inhibited the chlorpheniramine effect on RS-induced immunosuppression (P<0.05), but epinephrine had not a significant effect (Fig. 5). In rats immunized with SRBCs, and then challenged with SRBCs into the left paw (right paw received saline), there was a marked enhancement in the paw volume as measured by using the volume differential meter.

In the RS treated rats, there was significant reduction in paw volume difference compared to the control (P<0.01 Fig. 4). Similar effects on paw volume changes were also seen after histamine (150 μg/rat) injection in non-stressed rats (P<0.01 Fig. 5).

Fig. 4. The central effect of histamine on the cell immune response following restraint stress (RS) in rat. The effect of RS, histamine (hist, 150 μg/rat) and control (non-stressed) groups were compared on difference in paw volume (DIPV) in immunized rat. Each column shows mean ± SE of DIPV related to six rats. ** P<0.01


DISCUSSION

Complex neurochemical mechanisms are involved in the organism's biological response to noxious stimuli like stress. Stress can induce the immunosuppression effect in a specific condition.

It has been shown that the restraint stress (RS) for 2h reduces the humoral immune response (5/6) and RS for 1h daily for five days reduces the cell-mediated immune response (5) . In this study, RS induced a significant reduction in humoral and cellular immune function. The release of histamine and mediation of different nervous system activity following RS has been recognized (19,19,20,21). Similar immune response (immunosuppression) has been seen with histamine injection in the absence of RS. The similarity between histamine and RS effects on immune function can be a reason for the role of histamine in immunosuppression effect of RS. Pretreatment with ranitidine had no any effect on RS - induced immunosuppression but chlorpromazine reduced the RS effect on immune function. This result shows that histamine affects the immune function via action on H1 receptor in the brain. Simultaneous injection of histamine and chlorpromazine inhibited the effect of H1 receptor antagonist, so it can be concluded that the effect of chlorpromazine is related to inhibition of H1 receptor and the other effects of drugs on different sites are not important.

Following RS, histamine increases the serotonergic, dopaminergic and adrenergic nervous system via H1 receptor in the brain (19,20,21). Serotonin and dopamine could reverse inhibitory effect of chlorpromazine on RS - induced immunosuppression, but epinephrine could not induce significant effect. It can be concluded that the immunosuppression effect of RS via histamine release can be mediated by serotonergic and dopaminergic nervous system in the brain.

In summary, the RS induced immunosuppression by release of histamine and changing in dopaminergic and serotonergic nervous system activity via H1 receptor affects stimulation by histamine.

REFERENCES


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