

Lectin receptor interactions in rat lungs structural components on the background of experimental hypothyroidism



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INTRODUCTION

Hypothyroidism is one of the most common endocrine diseases, which develops due to deficiency of thyroid hormones. Statistics show an increase in the number of diseases of the endocrine system among the working age population in the world and in Ukraine in particular. The prevalence of hypothyroidism in the world is variously estimated as 1.5-6% and subclinical – 4.3-15% [5, 10, 11]. Hypothyroidism is a syndrome, or complex of changes in majority of organs and systems. Combined thyroid and bronchopulmonary pathology are often found, which significantly complicates diagnostics and often causes inadequate and ineffective treatment [1, 2, 14]. It is known, that the lack of hormones is common natural phenomenon that occupies a special place among the world's problems by the number of medical and social consequences for individuals and society. Analysis of the literature confirms the steady growth of bronchopulmonary lesions and frequency of endocrine dysfunction in older age groups [3, 4, 7, 13]. Alveolar hypoventilation of lungs is observed because of muscle discoordination and disorders of the central regulation. Vital lung capacity is slightly decreased due to the weakness of the intercostal muscles or depression of the respiratory center. The lungs are actively involved in the metabolism of glucocorticoids and biogenic amines, who regulate bronchial tonus [9, 12]. In the scientific literature available to us there are no data concerning the role of glycoconjugates in the secretory processes of lung structural components in norm and on the background of thyroid hypofunction.

THE AIM

On the basis of lectin - receptor interaction to investigate glycoconjugates of the lung structural components in norm and on the background of experimental hypothyroidism.

MATERIALS AND METHODS

Research was performed on 20 Wistar rats with mass 180-120g, which were divided into two groups: 1st – control (10), 2nd – experimental (10). Animals were kept in standard vivarium conditions in compliance with hygiene standards and food ration, manipulations were performed according to the positions of "General ethical principles of the experiments on animals", approved by I National Congress of Bioethics (Kyiv, 2001).

Experimental hypothyroidism was caused by daily administration with food the drug mercapozolium at a dose of 5 mg / kg body weight. Function control of thyroid gland was performed by determining T3 and T4 hormones in blood serum using radiological method by means of standard sets in radioisotope laboratory of regional clinical hospital and by measuring of thyrocytes' height, also colloid condition has been visually estimated.

Animals were scored by decapitation after ether narcosis overdose. Lung pieces were taken from normal and experimental animals and fixed in 4% neutral formalin. Sections 5-7 μm thick were stained by hematoxylin and eosin for getting panoramic slides. Carbohydrate determinants were examined by the following lectins: concanavalin A (Con A, specific to αDMan, αDGlC), peanut lectin (PNA, specific to βDGal (β1-3) DGalNAs), soybeans (SBA, specific to DGalNac), wheat germ (WGA, specific to DGlcNac, NeuNAs), elderberry bark (SNA, specific to Neu5As (α2-6) Gal / DGalNac) crust of golden rain (LABA, specific to αLFuc), labeled by horseradish peroxidase. Control of lectin-binding reaction was done by exclusion of the lectin from protocol, and also by exclusion of inhibition reaction of endogenous peroxidase with methanol. Visualization was conducted by 3'3'-diaminobenzidine tetrahydrochloride ("Sigma", USA) in H₂O₂ presence, as previously described [Lutsyk et al. 1989]. Slides were analyzed by means of Carl Zeiss Jena Ng microscope, for photographing digital camera Canon IXUS 700 and photo-system Olympus on the base of BX-41 microscope were used.

RESULTS AND DISCUSSION

In animals with experimental hypothyroidism decreased T3 and T4 hormone levels were observed in comparison with animal control group (Figure 1).

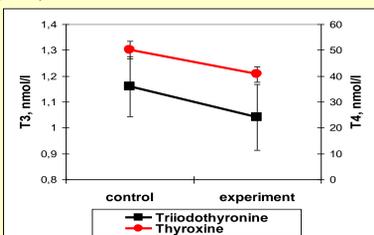


Figure 1. Thyroid hormones level in rat blood serum; p<0.01

Overview slides of rat lungs in the control group stained with hematoxylin and eosin showed, that the investigated organ has a typical structure, medium and small bronchi, terminal bronchioles, alveolar ducts and alveolar sacs are observed. Near bronchi lymphoid tissue in the form of follicles, and blood vessels filled with blood cells are located (Figure 2 A, B). Between alveoli in interstitial areas there are small capillaries and cellular elements characteristic for connective tissue.

In the lungs of the animal experimental group it has been noticed swelling of the endothelium, perivascular interstitial edema and vasodilatation filled with blood cells, among which red blood cells dominate. Amount of lymphoid tissue around blood vessels and bronchial tree increases (Figure 2 C, D). Probably in hypothyroidism interstitial edema is caused by processes of products accumulation of protein exchange, glycosaminoglycans, protein derivatives of glucuronic and chondroitinsulfuric acids.

Slides processing of animal lungs by lectins of different carbohydrate specificity in control and experimental group showed specificity of lectins binding in lungs structural components (Table 1).

So, DMan- specific lectin ConA in the animal lungs of the control group showed homogeneous binding with components of the respiratory part and conducting portion and interstitium including BALT (Bronchus Associated Lymphoid Tissue) components. In animal lungs of research group ConA lectin binding did not differ from the animals' control group.

WGA lectin revealed high affinity with luminal surface of epithelial cells of small and medium bronchi, clearly outlined vascular endothelium and formed elements in their lumen, and also apical surface of alveolar cells (Figure 4, A, B). In interstitium near alveoli it has been detected cell groups with eccentrically located nucleus and cytoplasmic granularity, that had high expression of that lectin receptors (Figure 4, C, D). The most intensive expression of WGA lectin receptors in animal experimental group was on the apical surface of epithelial cells of the small bronchi and type I alveolar cells (surfactant component), and vascular endothelium (Figure 5, A, B, C, D). The appearance of the glycoconjugates' NAcDGlC determinants, WGA lectin receptors as a part of surfactant indicates the peculiarity of its chemical composition, which can affect the permeability of air-blood barrier. Similar changes were observed by [8] as part of surfactant in preterm infants. In our case exhibiting of carbohydrate determinants NAcDGlC, suggests about metabolic disorders on the background of experimental hypothyroidism.

SBA lectin receptors in animals in the control group are associated with the apical surface of alveoli epithelial cells and apical surface of the cells of bronchial tree (Fig. 3 A, B), while in the experimental group lectin showed relatively high affinity to alveolar cells and alveolar macrophages (Fig. 3, C, D), whose number increased significantly. Increase in the number of alveolar macrophages can be considered as a factor of immune processes stimulation in response to the mercapozolium introduction.

Homogenous binding of SNA lectin was observed in animal lungs of control group with lungs structural components, while in experimental group that lectin revealed high affinity to goblet cells of bronchial tree. The rest of structural components demonstrated homogeneity with that lectin binding. Appearance of sialo-specific SNA lectin as part of goblet cells indicates, to our mind, the change in chemical composition of mucociliary barrier and their secretory activity and is one of manifestations of defence mechanisms.

Fucospecific LABA lectin in lungs of animal control group revealed affinity to the cells, which were located near alveoli or in alveolar lumen and which contained cytoplasmic granularity affine to that lectin. Besides abovementioned, LABA lectin receptors were identified on the apical surfaces of epithelial cells of some alveoli. In experimental group LABA lectin receptors were revealed in lymphoid follicles near small bronchi, probably, in dendritic cells. Also, expression of those lectin receptors was declared in perinuclear zone of small bronchi epithelial cells, and in secretory alveolar cells. Figdor CG et al. [15] detected, that C-type lectin receptors and lectin-like receptors are located on the surface of dendritic cells.

Currently it is known, that endogenous lectins recognize antigens and regulate the migration of dendritic cells and their interaction with lymphocytes. The appearance of LABA lectin receptors in some cells of lymphoid follicles, whose number is increasing on the background of hypothyroidism, confirms the version of [15] about activation of these receptors in the processes of interaction of lymphoid follicles cellular elements.

Homogenous binding of PNA lectin was observed in animal lungs of control and experimental group with alveolar cells, in some alveoli their apical surface was outlined.

Our studies have shown, that on the background of experimental hypothyroidism the modification of glycoconjugates in dendritic cells of lymphoid follicles, secretory alveolar cells and goblet cells of the bronchial tree, the endothelial cells of blood vessels are observed.

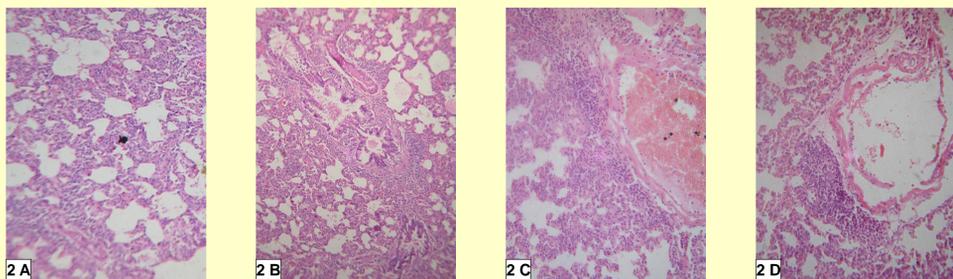


Fig. 2. Overview slides of rat lung. x400. H&E.

A, B – control; C, D – experimental hypothyroidism (expansion of the vascular lumen and perivascular edema)

RESULTS AND DISCUSSION

Table 1
Lectin receptors of rat lung structural components in the norm and in experimental hypothyroidism

Lectin name, and its carbohydrate specificity	Animal control group				Animal experimental group			
	Bronchi epithelium	Lymphatic nodules	Alveoli	Interstitial cells	Bronchi epithelium	Lymphatic nodules	Alveoli	Interstitial cells
Wheat ties lectin WGA (DGlcNac, NeuNac)	l.s. +++	h.b.	h.b.	+++	g.c. +++	h.b.	+++	+/-
Elderberry bark lectin SNA (Neu5Ac(α2-6)Gal / DGalNac)	h.b.	h.b.	h.b.	h.b.	g.c. +++	h.b.	h.b.	h.b.
Lectin crust golden rain LABA (αL-Fuc)	h.b.	h.b.	s.a. +++	+/-	s.b. +++	d.c. +++	s.a. +++	-
Concanavalin A lectin ConA (αDMan, αDGl)	h.b.	h.b.	h.b.	h.b.	h.b.	h.b.	h.b.	h.b.
Soybean lectin SBA (DGalNac)	l.s. +++	h.b.	+	h.b.	h.b.	h.b.	a.m. +++	h.b.
Peanut lectin PNA (βDGal(β1-3)DGalNac)	h.b.	h.b.	h.b.	h.b.	h.b.	h.b.	h.b.	h.b.

Note: - absence of binding; + slight reaction; l.s. - luminal surface; +++intensive; h.b.- homogenous binding; g.c. – goblet cells; d.c. – dendritic cells; s.a. – secretory alveolar cells; s.b. – small bronchi; a.m. – alveolar macrophages.

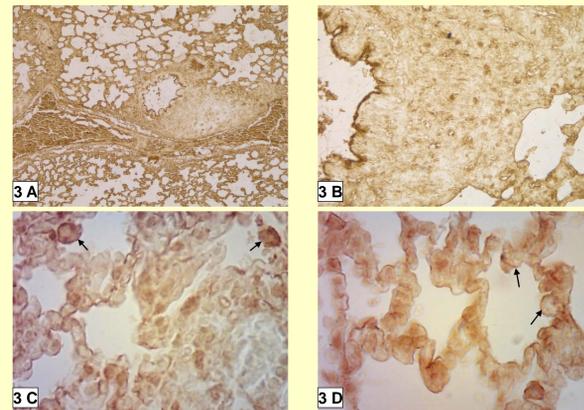


Fig. 3. A, B – localization of SBA lectin receptors on the luminal surface of epithelial cells of bronchial tree in rat control group; C, D – SBA lectin receptors on the surface of secretory alveolar cells and alveolar macrophages (arrows); x120 (A); x 300 (B); x 600 (C, D).

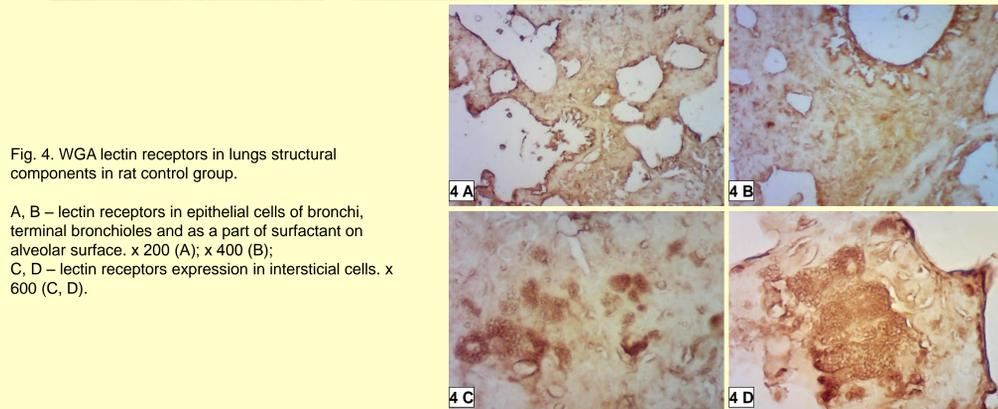


Fig. 4. WGA lectin receptors in lungs structural components in rat control group.

A, B – lectin receptors in epithelial cells of bronchi, terminal bronchioles and as a part of surfactant on alveolar surface. x 200 (A); x 400 (B); C, D – lectin receptors expression in interstitial cells. x 600 (C, D).

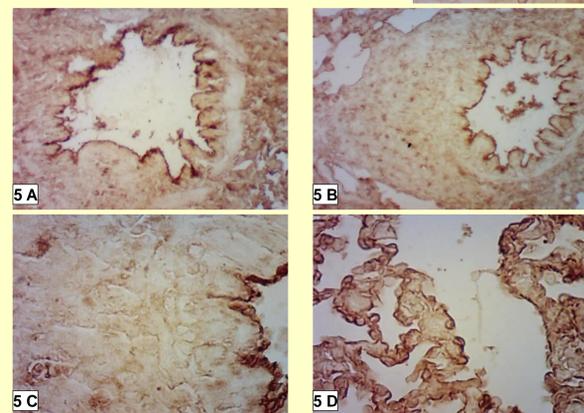


Fig. 5. Amplification of WGA lectin receptors in epithelial cells of bronchial mucosa (A, B, C), and on the surface of secretory alveolar cells (D) in rats on the background of hypothyroidism. x 600.

CONCLUSION

Experimental hypothyroidism, caused by introduction of mercapozolium at a dose of 5 mg / kg body weight is accompanied by slight perivascular interstitial edema and modification of lectin receptors in the structural components of the lungs, stimulates the activation of immune processes, that manifest the increase in number of SBA - positive alveolar macrophages and LABA - positive dendritic cells in lymphoid follicles associated with the bronchi. Lectin SBA (specific for a DGalNac) may be considered as one of markers of alveolar macrophages, lectin LABA (specific for αLFuc) as a marker of BALT-associated dendritic cells.

Prospects for further developments in this direction. There are plans to expand the range of lectins panel to determine the optimal set of lectins to study the lung structural components in norm and on the background thyroid dysfunction.

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