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MORPHOLOGY OF THE PROSTATE IN 6-MONTH-OLD RATS AND ITS REACTIVE CHANGES IN CHRONIC ALCOHOLISM

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ABSTRACT

The article analyzes the results of a study on the histological relationship of glandular and non-glandular structures of the prostate gland of mature rats and its structural changes in chronic alcoholism.

In rats with chronic alcoholism, compared with control, a decrease in the volume fraction of the glandular parenchyma in the structure of the organ, an increase in the number of acini in the field of view due to a decrease in the diameter of their lumen, foci of epithelial stratification, cell proliferation, and desquamation of epithelial cells are observed.

Chronic exposure to alcohol leads to a pronounced degree of lymphocytic infiltration with lymphoid nodular formation, to a moderate form of connective tissue proliferation, in the interglandular stroma the number and diameter of vessels increase and their wall thickness decreases.

KEYWORDS

Rat prostate, gland, morphometry, chronic alcoholism.

INTRODUCTION

Due to the wide spread of prostate diseases, functional disorders that occur in the prostate under the influence of environmental factors, the action of which

the body experiences in the process of individual development, are of particular interest (7,9,10).

The morphostructure of the prostate gland can be influenced by various environmental factors, including chemical ones. Alcohol is one of the most toxic xenobiotics, which has a detrimental effect on almost all organ systems (1,3,4,6,8).

According to the World Health Organization, as a result of the harmful use of alcohol in the world, 3 million deaths are recorded every year, which is 5.3% of all deaths [2].

It should be noted that in a comparative aspect, the morphology of the prostate at puberty and under chronic alcohol exposure with a description of the morphometry of all glandular and non-glandular structures has not been practically studied in a comprehensive manner. In this regard, the study of the morphological features of an organ under conditions of chronic alcohol exposure is of undoubted interest for theoretical and practical medicine.

MATERIAL AND METHODS

The study was performed on 26 outbred white male rats at the age of 6 months. 2 experimental groups were formed: 1st - control (n=14); 2nd – experimental group (n=12).

In the experimental group, for modeling chronic alcoholism, forced alcoholization of animals using 40.0% ethanol solution was used (5). The solution was administered intragastrically using a metal probe 1 time

per day at a total dose of 7 g/kg of body weight for 1 month before the study age. Control animals received intragastrically equal volumes of 0.9% NaCl solution.

For histological examination, pieces of the prostate were fixed in 10% buffered formalin and embedded in paraffin according to the standard method. Histological sections obtained from paraffin blocks, 5-7 μ m thick, were stained with hematoxylin and eosin for review purposes, collagen fibers were detected by van Gieson staining.

With a microscope magnification of 70 times (7x10), the sections were determined:

- the shape of the lumen of the glands, the number of terminal sections of the glands in the field of view, the volume fraction of acini with and without secretion (in %), the number of acini with desquamated epithelial cells in the field of view, in the intralobular stroma, the number of intraorgan vessels in the field of view was counted;

In the preparations, at a magnification of 280 times (7x40), using an eyepiece micrometer, the diameter of the lumen of the glands, the height of the epithelium, the inner diameter and wall thickness of intraorganic vessels were measured. In addition, the thickness of collagen fibers and their distribution in the tissues of the gland were determined.

In the field of view (7x40), the presence and severity of lymphocytic infiltration in the tissues of the gland was assessed. When distributing lymphocytes by severity (cell density), the classification of the North American Chronic Prostatitis Collaborative Research Network and the International Prostatitis Collaborative Network was used:

- 1) mild degree - single lymphocytic cells separated by distinct intermediate zones;
- 2) moderate degree - confluent fields of lymphocytic cells without tissue destruction and / or lymphoid nodular / follicular formation;
- 3) severe degree - confluent fields of lymphocytic cells with tissue destruction and / or lymphoid nodular / follicular formation.

To assess the severity (fibrosis) of the proliferation of connective tissue using an eyepiece micrometer with a magnification of the objective x40, eyepiece x7 in the field of view, the thickness of the stroma layers between the glands was measured.

The degree of compaction of the connective tissue was determined by the appropriate method as follows: mild form (increase in the thickness of stromal septa up to 2 times in 2-4 fields of view out of 10); 2) moderate form (the thickness of the stromal septa is increased up to 2 times in more than 4 fields of view or a sharp thickening - more than 3 times and is present in single

(1-2) fields of view); 3) pronounced form (stromal septa are enlarged up to 3 times or more in 7-10 fields of view).

A study was made of the volume fractions of glandular and stromal elements (in %). To do this, using the morphometric grid G.G. Avtandilov (with the number of intersections 100) using an eyepiece x10, a lens x10 in each preparation of the prostate in 10 fields of view, the number of intersections falling on the stromal and glandular (including the lumen of the gland) elements was counted to determine their ratios.

RESULTS AND DISCUSSION

The study showed that in 6-month-old rats, the prostate has a normal structural plan, consists of numerous separate alveolar-tubular glands and muscular-elastic stroma in the form of loose fibrous connective tissue, bundles of smooth myocytes and vessels.

Plain microscopy reveals single epithelial-stromal outgrowths in the acini, the terminal sections are represented by highly prismatic epithelium with high columnar and basal cells resting on a clearly visible basement membrane. The height of the epithelial layer varies from 12.6 to 21.0 μm , on average, $17.8 \pm 0.38 \mu\text{m}$. Acini have oval and rounded shapes. The diameter of the lumen of the glands ranges from 210.0 to 441.0 microns, on average - 330.5 ± 9.7 microns. The number of acini in the field of view ranges from 18 to 28,

averaging 22.0 ± 0.5 . The volume fraction of acini with a secret is in the range of 90.0-100%, on average - 93.3 ± 0.5 . The proportion of acini without a secret is 0-10.0%, on average 6.7 ± 0.5 . Acini with desquamated epithelial cells were not found in the preparations.

In the periglandular stroma, single scattered lymphocytes are determined, separated by clear intervals. Their number in the field of view ranges from 8 to 12, on average 10.0 ± 0.22 . In the preparations, a thin stroma is determined, in most cases the acini are located back to back. The thickness of the stromal septa between the acini ranges from 12.6 to 33.6 μm , averaging $23.1 \pm 1.13 \mu\text{m}$.

The number of stromal vessels in the field of view is in the range of 4-9, averaging 7.0 ± 0.3 . The inner diameter of the venules is in the range from 21.0 to 29.4 microns, on average - 24.8 ± 0.42 microns. The thickness of their wall ranges from 4.2 to 8.4 microns, on average - 5.8 ± 0.21 microns. The diameter of the capillaries varies from 8.4 to 16.8 μm , on average $13.1 \pm 0.42 \mu\text{m}$. The wall thickness is in the range of 2.1-4.2 microns, on average - 4.12 ± 0.13 microns. The inner diameter of arterioles ranges from 12.6 to 16.8 μm , averaging $14.7 \pm 0.21 \mu\text{m}$. Their wall thickness varies from 4.2 to 8.4 μm , on average $7.69 \pm 0.21 \mu\text{m}$.

The volume fraction of glandular tissue is 77.0-90.0%, on average $82.5 \pm 0.7\%$. The proportion of stromal tissue ranges from 10.0-23.0%, averaging $17.5 \pm 0.7\%$.

Collagen fibers envelop the terminal sections of the glands, most of the fibers lie under the epithelium, and form a fine-loop network in the stroma. The thickness of collagen fiber bundles varies from 4.2 to 12.6 μm , averaging $10.0 \pm 0.21 \mu\text{m}$.

The study showed that in 6-month-old rats of the experimental group, acini in 70-80% of cases have a folded appearance, are represented by a cubic epithelium, in some places a highly prismatic epithelium is determined. The thickness of the epithelial lining varies from 4.2 to 12.6 μm , on average, $9.79 \pm 0.38 \mu\text{m}$. In some preparations, areas of cell proliferation are detected in the epithelium, which are distinguished by a dark color due to the high density of epitheliocytes. In places in the epithelium, foci of epithelial stratification are defined, known as prostatic intraepithelial neoplasia (PIN). At the same time, the rows of layers in the epithelium are disturbed, cell polymorphism and the presence of large and multiple nucleoli in the nuclei are noted, mitotic cells are visualized in places. The terminal sections of the glands are predominantly irregular in shape; acini of rounded and oval shapes are found. The lumens of the acini are convoluted. The diameter of the lumen of the glands ranges from 105.0 to 327.6 microns, on average - 204.1 ± 9.2 microns. The number of acini in the field of view ranges from 20 to 52, averaging 35.9 ± 1.7 . The lumens of the acini are filled with a homogeneous secret. The volume fraction of acini with a secret is in the range of

85-95%, on average - 87.8 ± 0.5 . The proportion of acini without a secret is 5-15%, on average 12.2 ± 0.5 . In the lumen of individual acini, there are many fragments of desquamated, desquamated cells, whole conglomerates are formed, which partially fill the lumen of the acinus. In the field of view, the number of acini with desquamated epithelium varies from 5 to 11, on average 9.1 ± 0.3 .

In the interacinar stroma, a large number of diffusely scattered lymphocytes are visualized. In most preparations, merging fields of lymphocytes are observed, which, in the form of chains, envelop the terminal sections of the glands. Epithelial-stromal outgrowths are completely infiltrated with lymphocytes. In addition, focal accumulations in the form of nodules are found inside the lumens of the acini, in some places around the vessels of the intralobular stroma, single lymphocytes infiltrating the walls of the vessels are detected. At the same time, in some places there is a violation of the integrity of the epithelial lining. The number of lymphocytes in the stroma (in the field of view) ranges from 30 to 70, on average 46.0 ± 2.2 . The thickness of the stromal septa between the acini is sharply increased, especially in the subcapsular zone, it ranges from 75.6 to 201.6 μm , averaging $122.6 \pm 6.7 \mu\text{m}$.

In 6-month-old rats of the experimental group, the stroma is larger than the glandular tissue. Morphometry of the parenchymal-stromal ratio

showed that the relative area of its parenchyma varies within 37-55%, averaging $43.6 \pm 0.1\%$. The proportion of stromal tissue ranges from 45-63%, averaging $56.4 \pm 0.1\%$.

In the interglandular stroma, a large number of venules, capillaries and arterioles are determined. The number of stromal vessels in the field of view is in the range of 7-12, averaging 9.9 ± 0.3 . The inner diameter of the venules ranges from 25.2 to 33.6 μm , on average $29.8 \pm 0.42 \mu\text{m}$. The thickness of their wall ranges from 4.2 to 8.4 microns, on average - 4.54 ± 0.21 microns. The diameter of the capillaries varies from 12.6 to 16.8 μm , on average $15.75 \pm 0.21 \mu\text{m}$. The thickness of their wall is in the range of 2.1-4.2 microns, on average - 3.7 ± 0.13 microns. The inner diameter of arterioles ranges from 12.6 to 21.0 μm , averaging $17.85 \pm 0.42 \mu\text{m}$. Their wall thickness varies from 4.2 to 8.4 μm , on average $6.13 \pm 0.21 \mu\text{m}$.

In the experiment, numerous collagen fibers occupy all interepithelial regions of the stroma. They are found around the acini and ducts of the gland, where they densely braid the smooth myocytes of the stromal layer. In places, a coarse network of collagen fibers is formed in the interacinar stroma. The thickness of collagen fiber bundles varies from 4.2 to 8.4 μm , averaging $7.52 \pm 0.21 \mu\text{m}$.

CONCLUSION

The prostate of 6-month-old rats shows morphological features typical of an organ with complete differentiation of the development of glandular-stromal elements.

In animals with chronic alcoholism, there is an increase in the number of acini in the field of view due to a decrease in their diameter, a decrease in the volume fraction of the glandular parenchyma in the structure of the organ, and focal desquamation of varying severity. Acini are detected with foci of epithelial stratification, which can subsequently cause a cancerous process.

The impact of alcohol leads to a pronounced lymphocytic infiltration, to an increase in the number and diameter of blood vessels, to a moderate form of expansion of connective tissue layers, pathological growth of fibrous connective tissue and compaction of bundles of collagen fibers, which can be considered as a reaction of the body aimed at isolating the focus of damage from surrounding tissues.

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