

# DESCRIPTION OF IMMUNOGISTOCHEMICAL CHANGES IN ENDOMETRIOSIS

K.A. Karimjonov<sup>1</sup>  R.I. Israilov<sup>2</sup>, Mamataliev A.R.<sup>1</sup> 

1. Andijan State Medical Institute, Assistant, Andijan, Uzbekistan.

2. Republican Center of Pathological Anatomy, Director, Tashkent, Uzbekistan.

OPEN ACCESS  
*IJSP*

## Correspondence

K.A. Karimjonov, Andijan State Medical Institute, Andijan, Uzbekistan.

e-mail: [karimjonovhomidjon@gmail.com](mailto:karimjonovhomidjon@gmail.com)

Received: 07 March 2023

Revised: 13 March 2023

Accepted: 22 March 2023

Published: 31 March 2023

## Funding source for publication:

Andijan state medical institute and I-EDU GROUP LLC.

**Publisher's Note:** IJSP stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



**Copyright:** © 2022 by the authors. Licensee IJSP, Andijan, Uzbekistan. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY-NC-ND) license (<https://creativecommons.org/licenses/by-nc-nd/4.0/>).

**Abstract.** In the study of the etio-pathogenesis of endometriosis, the study of the molecular-biological specificity of eutopic and ectopic endometrium - the expression of estrogen and progesterone receptors, proliferation, apoptosis, adhesion, angiogenesis, and cell invasion markers is one of the promising directions. The purpose of the work: to study the molecular-biological characteristics of adenomyosis and ovarian endometriosis in the proliferative phase of the menstrual cycle in women of reproductive age without other gynecological pathologies. Materials and methods: In this study, biopsies of premenopausal (18-51 years 60 biopsies) and postmenopausal women (age 51 and older, 20 biopsies) who underwent hysterectomy between 2019 and 2022 were examined by immunohistochemical methods. In this study, in 2019-22 AVPAB examined 80 adenomyosis and ovarian endometriosis biopsies, immunohistochemical examination of estrogen and progesterone hormonal activity disorders showed that in 60-90% of cases, progesterone, and in 50-70% of cases, estrogen hormone leads to the development of uterine adenomyosis and glandular hyperplasia of the uterus. atypical development of the endometrium and transition to a malignant tumor disease was observed. Ki-67 and p53 were shown in 20% of postmenopausal patients with the transition to the initial state of a low-grade tumor of the uterus.

**Key words.** endometriosis, adenomyosis, estrogen, progesterone, K67, r53, immunohistochemical tests.

**Dolzarbliqi.** Endometriozda to'qimalarni geterotopiyasi, molekulyar genetik nuqsoni, apoptozni sekinlashuvi, biriktiruvchi to'qimali qobig'ini yo'qligi, maxalliy aseptik reaktsiyani mavjudligi va immunkompetent hujayralarni disfunktsiyasi kuzatiladi [1, 2, 5].

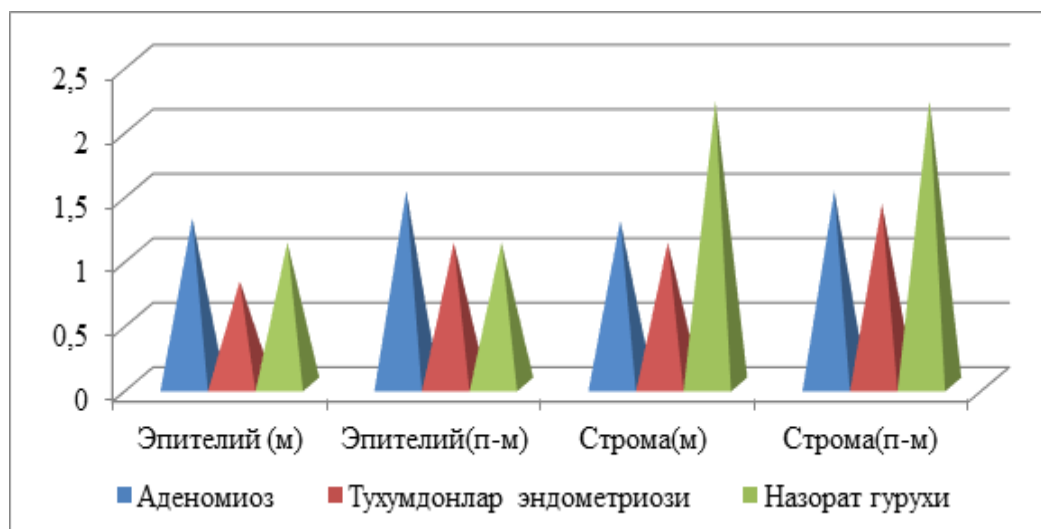
Endometriozni etio-patogenezi o'rganishda eutopik va ektopik endometriyaning molekulyar-biologik o'ziga xosligi-estrogen va progesteron restseptorlarini ekspressiyasi, proliferastiyasi, apoptozi, adgeziyasi, angiogenezi, hujayra invaziyasi markerlarini o'rganish perspektiv yo'nalishlardan hisoblanadi [1, 3,4, 6,7,8].

**Ishning maqsadi.** Reproduktiv yoshdagi ayollarda boshqa ginekologik patologiyalarsiz xayz davrining proliferativ bosqichidagi adenomioz va tухumdon endometriozidagi molekulyar-biologik xususiyatlarini o'rganish.

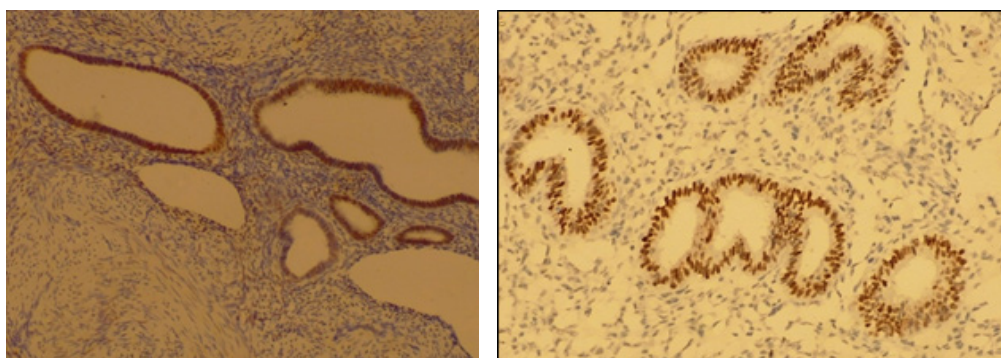
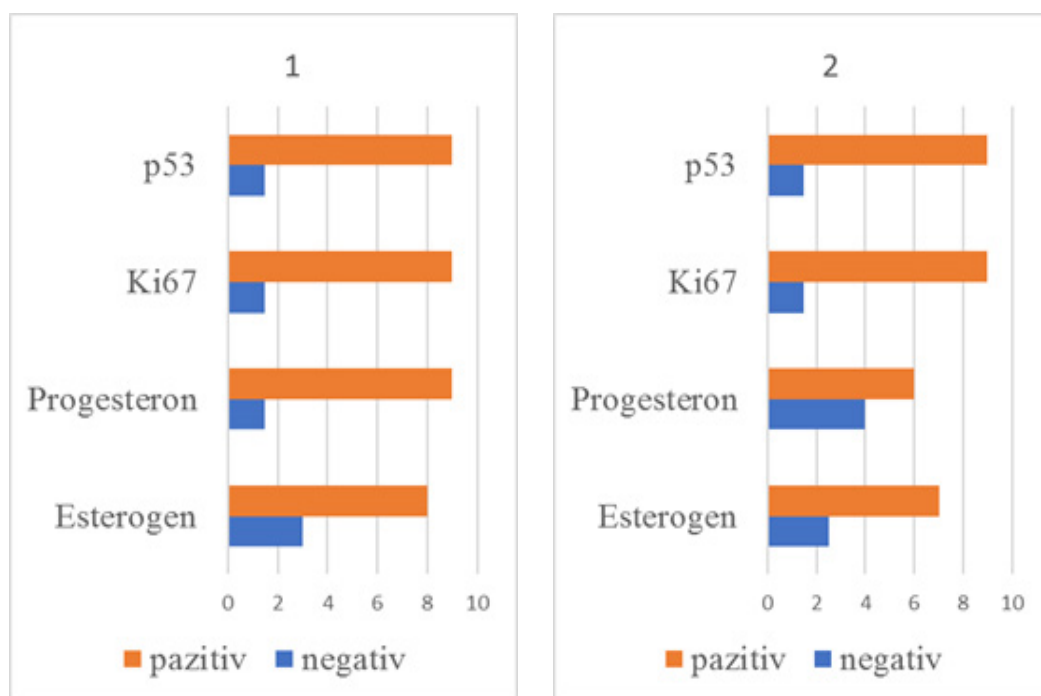
**Material va uslublar.** Ushbu tadqiqotda 2019-2022 yillar mobaynida gisterektomiya qilingan menopauzagacha (18-51yosh 60-ta bioptat) va postmenopauza davridagi ayollar bioptatlariga (51 va undan katta yoshdagilar, 20-ta bioptat) immunogistokimyoviy usullarda tekshirildi. Barcha bemorlar reproduktiv yoshda, buzilmagan xayz davrining proliferativ bosqichida bo'lgan. Immunogistokimyoviy tekshiruvga Bond Leica Australia (Avstraliya) immunogistoprostessordan foydalangan holda Ki67 va r53 ekspressiyasi, estrogen va progesteron gormonlari biomerklari ekspressiyasi o'rganildi, bunda musbat bo'yalgan hujayralar % bilan hisoblanib, proliferativ indeksi sifatida baholandi. Hujayralarni bo'yalish intensivligi (yoki ularni yadrolari - Ki - 67 oqsili, estrogen restseptorlari va progesteron restseptorlari uchun) vizual ravishda 0 dan 3 gacha (salbiy, zaif, o'rtacha bo'yalgan) ball bilan baholandi va ijobiy bo'yalgan hujayralar % har bir ko'rsatkichni intensivligi qiymatida hisoblandi.

**Natijalar.** Menopauzagacha bo'lgan davrdagi ayollarda estrogen restseptorlari ekspressiyasi o'rganilganda: adenomiozlarda - stroma hujayralarida, tухumdon endometriozlarida esa epiteliy va stroma hujayralarida miqdorini nazorat guruhiga nisbatan kam bo'lishi qayd etildi. Menopauzagacha bo'lgan davrdagi 10ta bemorlarni estrogen reagenti orqali olingan natijalar shuni ko'rsatdiki 8ta bemorlarda (80%) estrogen restseptori pozitiv reaktsiya jarayoni kuzatildi. 2 ta (20 %) bemorda negativ reaktsiya kuzatildi (diagramma-1,2).

**Diagramma-1**  
**Endometriyning proliferastiya fazasida adenomioz va tухumdonlar endometriozlarida esterogen resteptorlari ekspressiyalanishi**



**Diagramma-2**  
**Menopauzagacha (1) va menopauzadan keyingi (2) davrlarda bachadon adenomiozi va tухumdonlar endometriozlarida estrogen, progesteron, K67 reagentlarining proliferativ faollik darajasi**

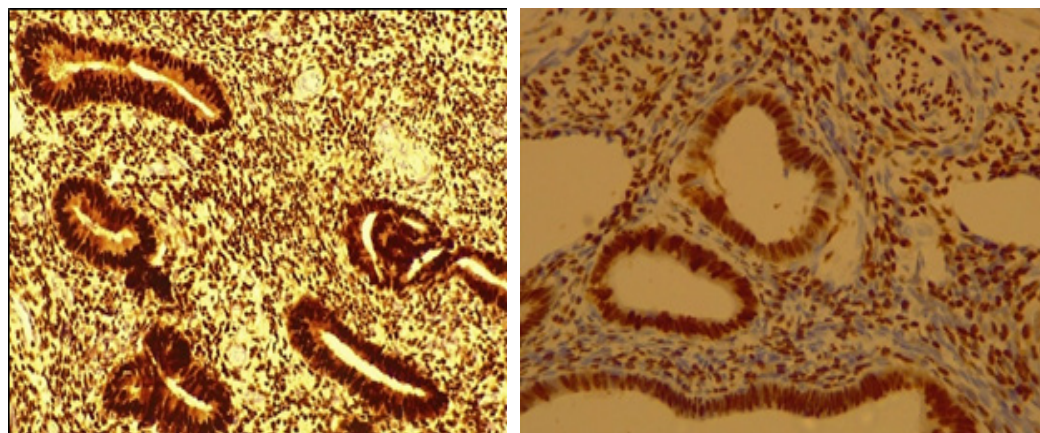


**Rasm-1.** Endometriyning proliferastiya fazasida adenomiozdagi epiteliy yadrosi va stromasida esterogen resteptorlarini ekspressiyasi. 2.Tухumdonlar endometriozlarida estrogen reagentining pozitiv reakstiyasi. IGX – Dab xromagen. Ob10xok40

Mikroskopik ko'rinishi bo'yicha miometriyadagi adenomioz (endometriya bezlarini proliferastiyasi), bezlar atrofida limfoid folikulalar proliferastiyasi va yallig'lanish o'choqlari mavjudligi bilan namoyon bo'ldi. Bezlar o'lchamlari turli ko'rinishda bo'lib, proliferastiyalangan, giperplaziyaga uchragan epiteliy, asosan bir qatorli bo'lib, immunogistokimyoviy tekshirishlarda yadrolari to'q jigarrang rangga bo'yaldi (rasm-1).

Menopauzagacha bo'lgan davrdagi bemorlarda adenomioz va tuhumdonlar endometriozi progesteron resteptorlari ekspressiyasi nazorat guruhida epiteliyda nisbatan yuqori, stromasida esa nazorat guruhi bilan bir xil ko'rsatkichda namoyon bo'ldi, natijalar 9ta bemorlarda (90%) pozitiv reakstiya, 1 ta (10%) bemorda negativ reakstiya kuzatildi (diagramma 2,3).

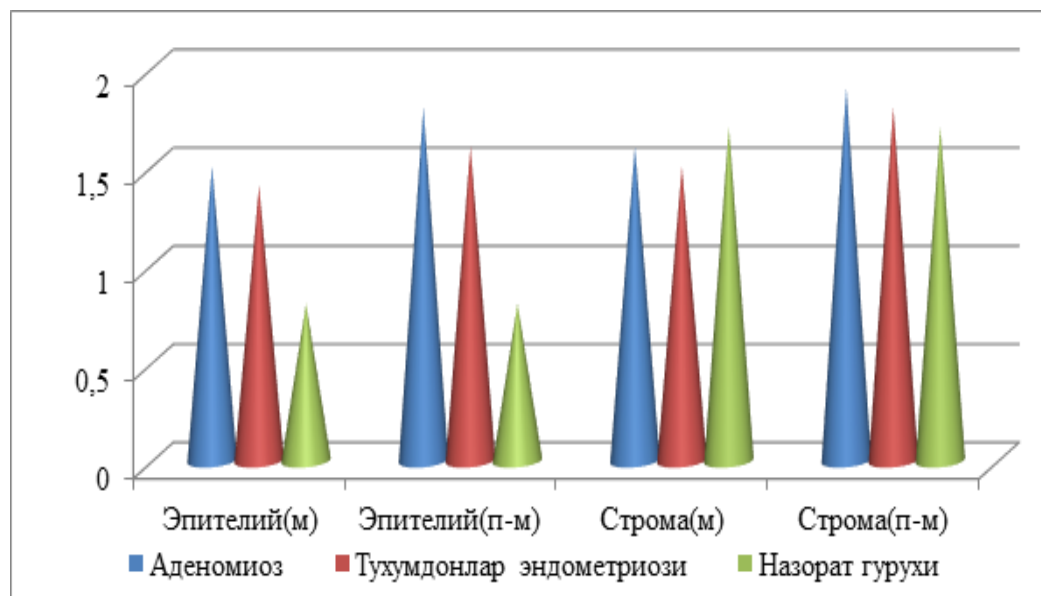
Immunogistokimyoviy ko'rinishi bo'yicha esterogen resteptorlari ekspressiyasi epiteliy hujayralariga nisbatan, progesteron resteptorlari ekspressiyasi esa stromada ekspressiyalanishi pasayishi bilan namoyon bo'lib, adenomiozda miometriyda endometriya bezlarini proliferastiyasi, giperplaziyasi, bezlar atrofida limfoid hujayralarini proliferastiyasi va yallig'lanish o'choqlari mavjudligi bilan birga kuzatildi. Bezlar o'lchamlari turli ko'rinishda bo'lib, immunogistokimyoviy tekshirishda endometriya bezlari giperplaziyalanib, asosan bir qatorli va yadrolari to'q jigarrang rangga bo'yalganligi aniqlandi (rasm-3).



**Rasm-2.** 1.Bachadon adenomiozida miometriyda endometriyaning bezlarini proliferastiyasi, giperplaziyasida progesteron resteptori pozitiv reakstiyasi. 2.Proliferastiya fazasida tuhumdonlar endometriozi endometrioid o'chog'larida epiteliy yadrosi va stromasida progesteron resteptorlari ekspresssiyasi: epiteliyda ekspressiyani ustunlik sohalari. IGX – Dab xromagen. Ob10x. ok40.

**Diagramma-3**

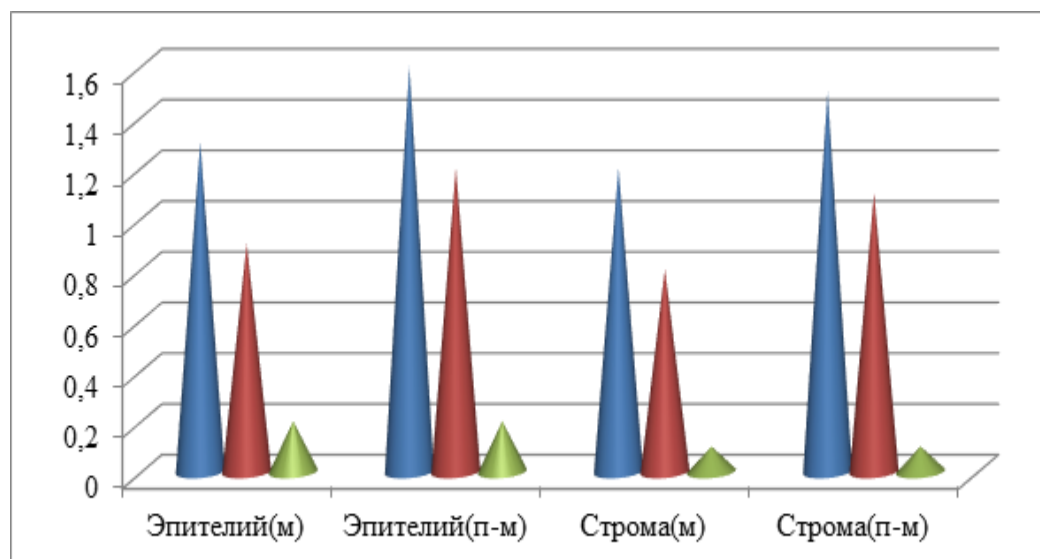
**Endometriyning proliferastiya fazasida adenomioz va tuhumdonlar endometriozi endometriozi progesteron resteptorlari ekspressiyalanishi**



**Izox:** m-menopauzagacha bo'lgan davr, p-m postmenopauzadan keyingi davr.

Diagramma-4

Endometriyning proliferastiya fazasida adenomioz va tuhumdonlar endometriozlarida Ki – 67 ekspressiyasi koeffitsienti

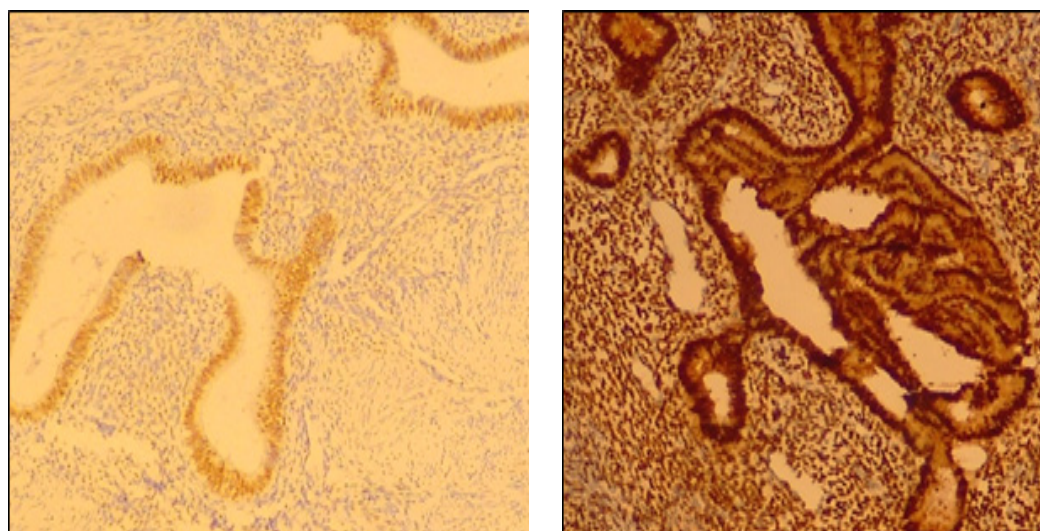


**Izox:** m-menopauzagacha bo'lgan davr, p-m postmenopauzadan keyingi davr.

Menopauzacha bo'lgan davrdagi ayollar adenomiozida Ki-67 ekspressiyasi koeffitsienti adenomioz epiteliysida nazorat guruhi bilan bir xil, stromasida esa past, tuhumdonlar endometriozida o'chog'ida epiteliy va stromasida nazorat guruhida ko'rsatkichidan past natijani namoyon qildi, natija 9tasida engil darajali (5-10 %), 1tasida (10-20%) o'rta faollik holati kuzatildi (diagramma-2,4).

Adenomioz va tuhumdonlar endometriozlari epiteliysi va stromasi nazorat guruhi ko'rsatkichlariga nisbatan kamroq bo'lishi qayd etildi.

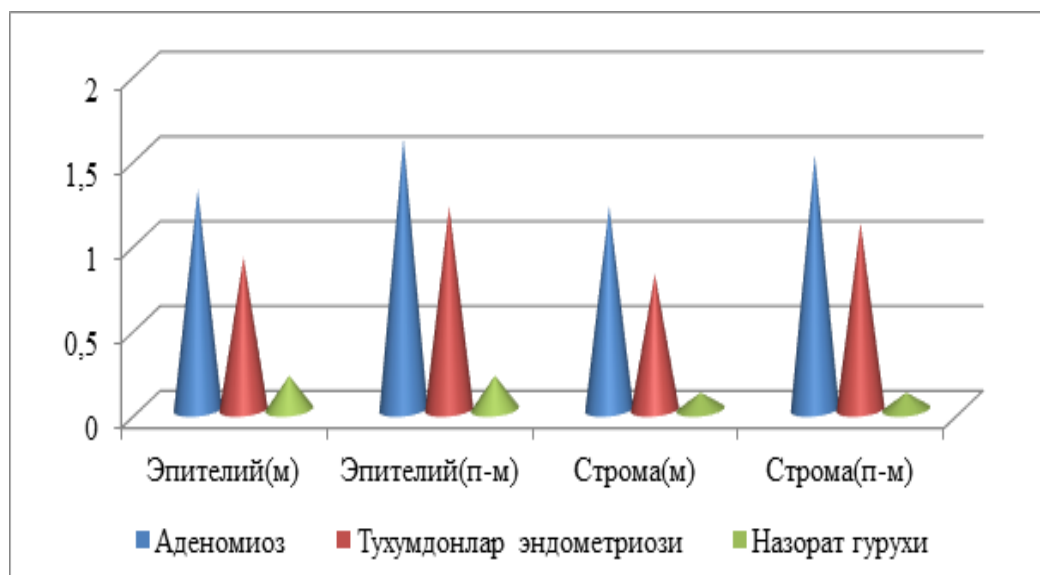
Immunogistokimyoviy ko'rinishi bo'yicha adenomiozda endometriya giperplaziyasi, bezlar atrofida limfoid folikulalar proliferastiyasi va yallig'lanish o'choqlari mavjudligi kuzatildi. Bezlari o'lchamlari turli ko'rinishda giperplaziyaga uchragan, asosan bir qatorli, immunogistokimyoviy ko'rinishda endometriya bezlari giperplaziyalanishi va yadrolari to'q jigarrang rangga bo'yalishi aniqlandi (rasm-3).



**Rasm-3.** 1.Bachadon adenomiozida endometriya bezli giperplaziyasining Ki – 67 ekspressiyasi koeffitsientini o'rta darajali pozitiv reakstiyasi. 2.Bachadon adenomiozida endometriyaning atipik giperplaziyasida r53 reagentning pozitiv reakstiyasi. Epiteliy va stromal hujayralarda apoptoz ingibitorlari ekspressiyasi. IGX – Dab xromagen. Ob10. Ok40.

Diagramma-5

**Endometriyning proliferastiya fazasida adenomioz va tuhumdonlar endometriozlarida apoptoz ingibitori ekspressiyasi**

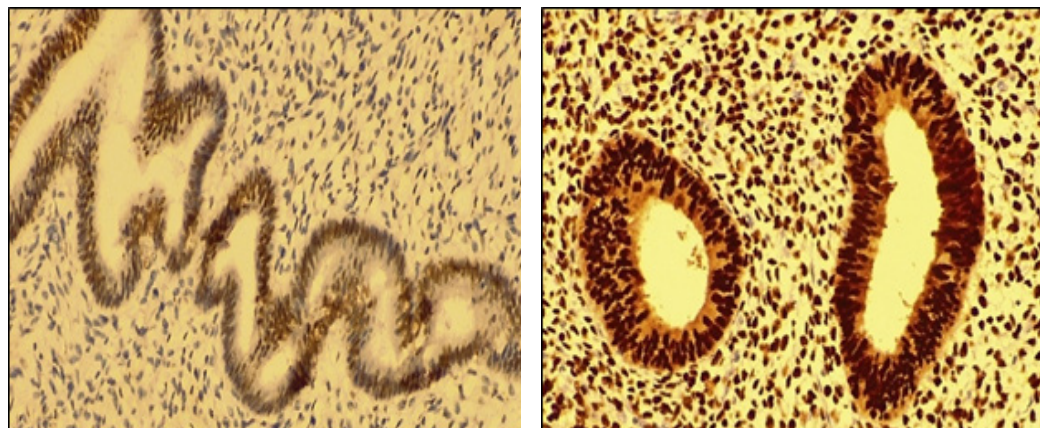


**Izox:** m-menopauzagacha bo'lgan davr, p-m postmenopauzadan keyingi davr.

**Menopauzagacha bo'lgan davrdagi** ayollar bachadon adenomiozi r53–o'sma oqsili ekspressiyasi natijasi 1tasida (10 %) pozitiv reakstiya, 9tasida (90%) negativ reakstiya holati kuzatildi ( 4 -rasm).

Adenomioz va tuhumdonlar endometriozlarida apoptoz ingibitorlari ekspressiyasi epiteliy va stromasida nazorat guruhiga nisbatan baland ko'rsatkichni namoyon qildi.

Immunogistokimyoviy ko'rinishi bo'yicha bachadon endometriya to'qimasining kistoz shakldagi polimorfizm ko'rinishdagi atipik hujayralarga ega giperplaziyasi, hujayra yadrolarini to'q jigarrang rangga bo'yalishi adenomiozda bachadon bezli giperplaziyasining xavfli o'sma kasalligiga o'tganligidan dalolat beradi.



**Rasm-4.** 1.Bachadon adenomiozida endometriyaning bezli giperplaziyada estrogen reagentining pozitiv reakstiyasi. 2.Bachadon adenomiozida endometriyaning bezli giperplaziyasida progesteron resteptorlarini pozitiv reakstiyasi. IGX – Dab xromagen. Ob10xok40.

Postmenopauza davrida estrogen reagenti orqali olingan natijalar 7 tasida (70%) pozitiv reakstiya, 3 tasida (30%) negativ reakstiya kuzatildi (diagramma-2, 5).

Mikroskopik ko'rinishi bo'yicha adenomiozli endometriya giperplaziyasida bezlari o'lchamlari turli ko'rinishda giperplaziyaga uchragan, asosan bir qatorli, immunogistokimyoviy ko'rinishda yadrolari to'q jigarrang rangga bo'yalganligi bilan namoyon bo'ldi (rasm-4).

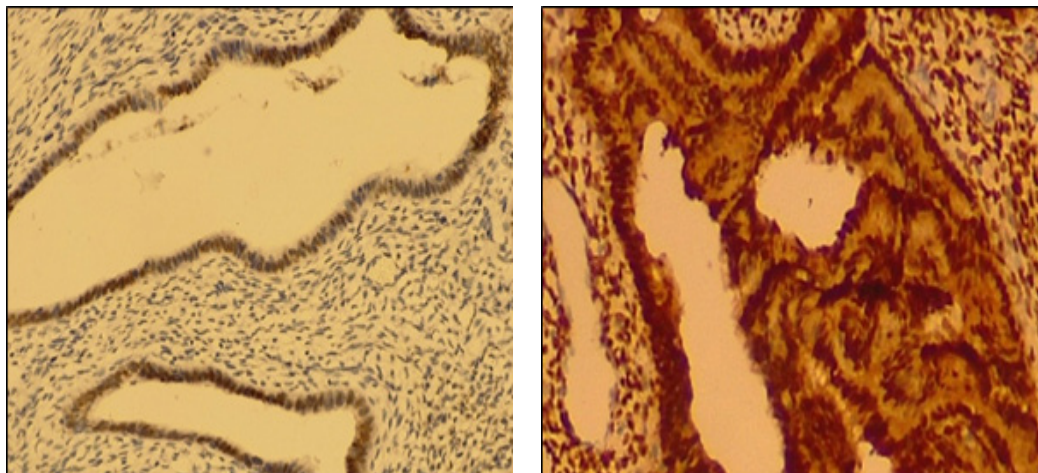
Postmenopauza davrida progesteron reagenti natijalar 6 ta bemorlarda (60%) pozitiv reakstiya, 4ta bemorda (40%) negativ reakstiya kuzatildi.

Immunogistokimyoviy ko'rinishi bo'yicha adenomiozda endometriy giperplaziyasi

bezlari o'lchamlari turli ko'rinishda giperplaziyaga uchragan, asosan bir qatorli, immunogistokimyoviy ko'rinishda endometriya bezlari giperplaziyalangan va yadrolari to'q jigarrang rangga bo'yalganligi kuzatildi.

Postmenopauza davrida Ki 67- hujayralar proliferativ faolligi natijasi 8 tasida engil darajali (5-10%), 2 tasida (10-20%) o'rta faollik holati kuzatildi (diagramma-5).

Immunogistokimyoviy ko'rinishi bo'yicha adenomiozda endometriy giperplaziyasi, bezlar atrofida limfoid folikulalar proliferastiyasi va yallig'lanish o'choqlari mavjudligi kuzatildi. Bezlarni o'lchamlari turli ko'rinishda giperplaziyaga uchraganligi, asosan bir qatorli, immunogistokimyoviy tekshirishlarda bezlar giperplaziyasi va yadrolarini to'q jigarrang rangga bo'yalishi qayd etildi (rasm-4).



**Rasm-4.** 1.Bachadon adenomiozida endometriy bezli giperplaziyasining Ki – 67 yuqori darajali pozitiv reakstiyasi. 2.Bachadon adenomiozida endometriyaning atipik giperplaziyasida r53 reagentning pozitiv reakstiyasi. IGX – Dab xromagen. Ob10. Ok40.

r53 – o'sma oqsili postmenopauza davrida gistologik va immunogistokimyoviy tekshiruv natijalarida 2ta bemorda adenomioz tashxisi bilan jarrohlik amaliyoti o'tagan bemorlar bachadonning boshlang'ich yomon sifatli o'sma holatiga o'tganligi namoyon bo'ldi, natijalar 2tasida (20%) pozitiv reakstiya, 8tasida (80%) negativ reakstiya holati kuzatildi (rasm-4).

**Hulosa:** Olingan natijalardan shuni ko'rsatadiki immunogistokimyoviy tekshiruv orqali ayollarda estrogen va progesteron gormonal faoliyatni buzilishi 60-90% xollarda progesteron, 50-70% hollarda estrogen gormoni bachadon adenomiozi rivojlanishda va bachadonning bezli giperplaziyasiga olib keladi, buni natijasida bemorlarda bachadon endometriysini atipik rivojlanishi va havfli o'sma kasalligiga o'tishi kuzatildi. Ki-67 va r53 postmenopauzada davridagi 2 ta bemorda bachadonning boshlang'ich yomon sifatli o'sma holatiga o'tganligi bilan namoyon bo'ldi.

#### LIST OF REFERENCES

- [1] Adamyan L.V. et al. The role of proliferation processes in the pathogenesis of endometriosis. *Reproduction problems. Technologies of the XXI century in gynecology.* Спец Выпуск-М 2008:82.
- [2] Zairatians O.V. et al. The role of apoptosis and proliferation in the pathogenesis of simple and proliferating uterine fibroids in combination with adenomyosis / In the book. *Uterine fibroids.* Edited by corresponding member of the Russian Academy of Sciences Sidorova I.S. M: "MIA" 2002:113–27.
- [3] Voloshchuk et al. Molecular biological aspects of the pathogenesis of adenomyosis. *Pathology Archive* 2007:56–61.
- [4] Sidorova I.S. et al. Molecular and biological features of endometriosis, pathogenetically justified therapy pathways. *Abstracts of the reports of the Russian Medical Forum-2006 "Fundamental science and Practice"-M* 2006:128.
- [5] Kogan E.A. et al. Clinical and morphological parallels and molecular aspects of adenomyosis morphogenesis. *Archive of pathology* 2008:8–12.
- [6] Matsumoto Y, Iwasaka T, Yamasaki F, Sugimori H. Apoptosis and Ki-67 expression in adenomyotic lesions and in the corresponding eutopic endometrium. *Obstet Gynecol* 1999;94:71–7. [https://doi.org/10.1016/s0029-7844\(99\)00279-3](https://doi.org/10.1016/s0029-7844(99)00279-3).

[7] Kitawaki J. et al. Expression of aromatase cytochrome P450 protein and messenger ribonucleic acid in human endometriotic and adenomyotic tissues but not in normal endometrium. *Biol Reprod* 1997;57:514–9.

[8] Schmidt C. Endometriosis: Pathogenesis and Treatment. *Fertil Steril* 1990;53:407–10.