Histopathological Pattern of Endometrium in Abnormal Uterine Bleeding

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Abstract: Abnormal uterine bleeding (AUB) is one of the commonest complaints in women and when it occurs without organic lesions like tumor, inflammation, it is called as dysfunctional uterine bleeding. The aim of this study was to find the histopathological pattern of endometrium in abnormal uterine bleeding in all age groups. This is a prospective cohort study of 221 patients admitted over the period 2012 and May 2014 to the Obstetric Gynecologic University Hospital"Queen Geraldine", in Tirana, Albania.Cases of AUB with a probable endometrial cause were included in the study. The most common histological pattern of endometrium includes Secretory (29%) followed by Proliferative endometrium(25%), disordered proliferative (13%) and endometrial hyperplasia(7%).All patients having abnormal menstrual bleeding should be subjected to dilatation and curettage to rule out endometrial pathology.

Keywords: abnormal uterine bleeding, endometrium, histological pattern

1. Introduction

Abnormal uterine bleeding (AUB) is one of the commonestcomplaints leading to endometrial sampling byendometrial biopsy or curettage. Examination ofendometrial biopsy is challenge а to practicingpathologists, largely due to the wide range ofmorphologic patterns resulting from both normal andabnormal changes, exogenous hormones, infectionsand intrauterine tumor (1). Abnormal uterine bleedingmay be defined as bleeding pattern that differs infrequency, duration and amount from a patternobserved during a normal menstrual cycle or aftermenopause. It is a common problem having a long listof causes in different age groups (2). It interferessignificantly with the quality of life in an otherwisehealthy woman (3). In order to evaluate endometrialsamples information regarding age and menstrualhistory with clinical examination are a prerequisite (4). Histological examination of the submitted endometrial tissue remains the standard diagnostic procedure for the assessment of abnormal uterine bleeding. In addition, accurate histopathological diagnosis facilitates the implementation of optimal treatment strategies 4. Endometrium is a dynamic, hormonally sensitiveand responsive tissue which constantly andrhythmically undergoes changes in the activereproductive life. Abnormal uterine bleeding may be defined as a bleeding pattern thatdiffers in frequency, duration and amount from apattern observed during a normal menstrual cycleor after menopause (5). AUB is one of the mostcommon problems in women of all agesespecially those in the peri-menopausal agegroup. The abnormal bleeding can be caused by awide variety of disorders and it is one of the commonest complaints leading to endometrialsampling. It may represent a normal physiological state, and observation alone may be warranted. Alternatively, the bleeding can be a sign of underlying condition necessitatingaggressive aserious treatment. Dilatation and curettage is auseful and cost effective method of detectingintrauterine pathologies and very few lesionsescape detection (6). Wide range of morphologicpatterns resulting from both normal and abnormalchanges offer a diagnostic challenge topracticing pathologists.Histopathological diagnosis varies according to the age with endometrial hyperplasia and cancer are higher in peri and postmenopausal women while in younger age groups, changes related to hormonal effects seems to be more common. Histological characteristics of endometrial biopsy material as assessed by light microscopy remain the diagnostic standard for the clinical diagnosis of endometrial pathology. Indeed, the initial diagnosis is made by endometrial biopsy or by curettage, which in itself may be therapeutic. Conversely, the biopsy or curettage may not sample the entire endometrium, and the areas of greatest histological or cytological severity may thus escape histological identification. Management of AUB is not complete without tissue diagnosis especially in perimenopause and post menopause. AUB may be thesymptom of endometrial carcinoma in 8-50% of cases (7). The aim of this study was to find the histopathological pattern of endometrium in abnormal uterine bleeding in all age groups.

2. Materials and Methods

This is a prospective cohort study of 221 patients admitted over the period 2012 and May 2014 to the Obstetric Gynecologic University Hospital"Queen Geraldine", in Tirana, Albania. Patients with a gestational cause, hemostatic disorders, isolated cervical or vaginal pathology, and leiomyoma excluded. Relevant clinical data regarding age, pattern and duration of abnormal bleeding, menstrual history, obstetric history, use of exogenous hormones, physical and gynecological examination findings, lab investigation results, and sonological and hysteroscopic findings were collected. Detailed clinical history like age, menstrual status including pattern, period & regularity of cycle were obtained relevant findings of general, systemic examination were recorded. Patients were categorized intoreproductive (<40 yrs), perimenopausal (40-50yrs) andpostmenopausal (>50yrs) age groups. Histopathologicaldiagnosis was made, recorded and furthercategorization was done for all cases. All the specimens were fixed in 10% formalin, processed and embedded in paraffin, and 3-4 μ thick sections were made. Sections were stained with hematoxylin and eosin stain.

3. Results and Discussion

A total of 221 endometrial specimens submitted witha clinical diagnosis of AUB were studied.Patients' age ranged from 18-59 years and most of themwere seen in the age group of <40 years, followed by40-50 years. The commonest complaint was menorrhagia in98 patients (44%).Twenty four (11%) of patients were nulliparous, 71 (32%) were primiparous and 126 (57%) multiparous. In our study, 64%% of the patients were of normalweight, 20%% patients were overweight, and 16% wereobese. The most frequent pathology observed in the study wasSecretory endometrium in 64 (29%) of patients followed by proliferativeproliferative endometrium 55 in (25%)patients. Abnormal vaginal bleeding is defined as the appearance of blood at the vaginal introitus exclusive of normal menstruation and could present as menorrhagia, metrorrhagia, polymenorrhea, metromenorrhagia, peri and postmenopausal bleeding. Abnormal uterine bleeding can be caused by a wide variety of disorders, It might be part of normal physiological state such as adolescence, perimenopausal, lactation and pregnancy or it may be caused by a pathological process that is not directly related to the uterus such as hyper androgenic anovulation in patients with polycystic ovaries, hypothalamic dysfunction, hyperprolactinemia, hypothyroidism, pituitary disease, premature ovarian failure and iatrogenic causes such as irradiation or chemotherapy. The bleeding could be a sign of an underlying localized condition including benign tumors, malignancy and infection. Endometrial cancer and premalignant atypical hyperplasia are likely causes of abnormal bleeding in peri and postmenopausal bleeding. In the present study, the two most common endometrial histopathological patterns in all three age groups were secretory and proliferative endometrium. The bleeding in secretory phase is due to ovulatory dysfunctional uterine bleeding and is characterized by regular episodes of heavy menstrual blood loss. The main defect is in the control of processes regulating the volume of blood lost during the menstrual breakdown of endometrium (8). This pattern was commonly observed in the late reproductive and perimenopausal women in our study and other studies and may be due to the hormonal imbalance in this group leading to intermittent anovulatory cycles. The maximum incidence of AUB was in the <40 years age range 121 (55%) patients, followedby 40-50 years age group 83 (38%) patients. Our study and other studies have found a maximum incidence of AUB in the perimenopausal age group (9, 10). Perimenopause is defined by the as the 2-8 years preceding menopause and the 1 year after the final menses (11). As women approach menopause, cycles shorten and often become intermittently anovulatory due to a decline in the number of ovarian follicles and fluctuations in the estradiol level leading to various patterns of abnormal bleeding (12)Our study and other studies found menorrhagia as the most common complaint (13). Most of our patients were in parity the low category.Disordered proliferative endometrium is an exaggeration of the normal proliferative phase without significant increase in the overall ratio of glands to stromaand is due to persistent oestrogen stimulation (14). This pattern is particularly seen in perimenopausal women. This pattern was seen in 19 (12.2%) of our cases.

Endometrial hyperplasia was the most common histological pattern observed in our study and was seen in 55 (25%) cases. The incidence of endometrial hyperplasia without and with atypia peaks in the early 50s and early 60s respectively (15). In the present study, the maximum incidence of hyperplasia was noted in the 40-50 year age group and was seen in 9 (60%) of patients. This finding is similar to other studies (16). Among women undergoing endometrial biopsy or hysterectomy, the prevalence of endometrial polyps is 10-24%; the incidence rises with increasing age, peaks in the fifth decade of life and gradually declines after menopause (17). The present study showed a progressively increasing detection pattern of endometrial polyps in older age, 5.6% in the age group of <40 years, 10.2% in the age group of 40-50 years old and 13.1% in age group >50 years, our result is comparable to other studies (18). There is no direct evidence for a greater propensity of polypoid endometrium to undergo malignant change compared to the adjacent normal endometrium.. The present study shows that the detection rate of endometrial carcinoma increase with increasing age, 0.3% in reproductive age group, 1.5% in perimenopausal age group and 3.3% in post menopausal age group. This finding is similar to other studies (19). Endometrial hyperplasia is a precursor of endometrial cancer. Abnormal uterine bleeding in postmenopausal women requires further evaluation to exclude malignancies (6. 5). In this study complex hyperplasia with atypia was seen in the age groups <40 years 5 (33%0, 40-50 years 9 (60%) and >50 years. These figures are similar to some studies but different from others (20). Anovulatory and exogenous hormonal effect were found in 11 (5%) and 22 (10%) patients.

4. Conclusion

The present study revealed that secretory and proliferative endometrium are the most common endometrial histopathological patterns in endometrial samples obtained for abnormal uterine bleeding in our region. All patients having abnormal menstrual bleeding should be subjected to dilatation and curettage to rule out endometrial pathology. Accurate analysis of endometrial sample is the key to effective therapy and rational approach to treatment of women with abnormal uterine bleeding. Thus, histopathological evaluation of endometrium is especially indicated in women over the age forty years to rule out preneoplastic lesions and malignancies.

References

- [1] Goldstein SR. Menorrhagia and abnormal bleeding before themenopause. Best Pract Res ClinObstetGynaecol 2004;18:59-69.
- [2] Cornitescu FI, Tãnase F, Simionescu C, Iliescu D. Clinical, histopathological and therapeutic considerations in nonneoplastic abnormal uterine bleeding in menopause transition. Rom J MorpholEmbryol 2011;52:759-65.
- [3] McCluggage WG. Benign Diseases of the Endometrium. In: Kurman RJ, Ellenson LH, Ronnett eds. Blaustein's Pathology of the Female Genital Tract. 6th Ed. New York: Springer Verlag, 2011: 305-58.

International Journal of Science and Research (IJSR) ISSN (Online): 2319-7064 Index Copernicus Value (2013): 6.14 | Impact Factor (2014): 5.611

- [4] Ara S, Roohi M. abnormal Uterine Bleeding: Histopathological diagnosis by conventional dilatation and curettage. Prof Med J 2011; 18(4): 587-91.
- [5] Munro MG, Critchley HO, Fraser IS, FIGO Menstrual Disorders Working Group. The FIGO classification of causes of abnormal uterine bleeding in the reproductive years. FertilSteril 2011;95:2204-8.
- [6] Goldstein SR. Menorrhagia and abnormal bleeding before the menopause. Best Pract Res ClinObstetGynaecol 2004;18:59-69.
- [7] Bhosle A, Fonseca M. Evaluation and histopathological correlation of abnormal uterine bleeding in perimenopausal women. Bombay Hosp J 2010;52:69-72.
- [8] Mutter GL. Diagnosis of premalignant endometrial disease. J ClinPathol 2002;55:326-31
- [9] Baak JP, Mutter GL. EIN and WHO94. J ClinPathol 2005;58:1-6.
- [10] Perveen S, Perveen S. Endometrium histology in abnormal uterine bleeding. MC 2011;17:68-70.
- [11]Breijer M, Timemermans A, van Doorn HC, Mol BW, Opmeer BC. Diagnostic strategies for postmenopausal bleeding. ObstetGynecolInt 2010;2010:850812.
- [12] Lasmar RB, Dias R, Barrozo PR, Oliveira MA, Coutinho Eda S, da Rosa DB. Prevalence of hysteroscopic findings and histologic diagnoses in patients with abnormal uterine bleeding. FertilSteril 2008;89:1803-7.
- [13] AlHilli MM, Hopkins MR, Famuyide AO. Endometrial cancer after endometrial ablation: Systematic review of medical literature. J Minim Invasive Gynecol 2011;18:393-400.
- [14] Damle, R. P., Dravid, N. V., Suryawanshi, K. H., Gadre, A. S., Bagale, P. S. and Ahire, N. 2013. Clinicopathological spectrum of endometrial changes in perimenopausal and postmenopausal Abnormal uterine bleeding: A 2 year study. J ClinDiagn Res., 7 (12):2774-6.
- [15] Jetley, S., Rana, S., Jairajpuri, Z. S. 2013. Morphological spectrum of endometrial pathology in middle aged women with atypical uterine bleeding: A study of 219 cases. J Midlife Health., 4(4):216-20
- [16] Lacey JV Jr, Chia VM. Endometrial hyperplasia and the risk of progression to carcinoma.*Maturitas*. 2009;63(1):39–44.
- [17] ACOG Committee on Gynecologic Practice. Committee Opinion: number 263, December 2001. vonWillebrand's disease in gynecologic practice. Obstet Gynecol. 2009;98(6):1185–1186.
- [18] S.. Lama, S., KCS. 2013 Histopathological pattern of abnormal uterine bleeding in endometrial biopsies. Nepal Med Coll J., 15(1):74-7
- [19] Cho, H. J., Lee, E. S., Lee, J. Y., Hong, S. N., JIYI, Kim, H. Y. and Kim A. 2013. Investigations for postmenopausal uterine bleeding: Special consideration for endometrial volume. Arch Iran Med., 16(11):665-70
- [20] Koss LG. Proliferative disorders and carcinoma of endometrium. In: Koss LG, Melamed MR, editors. Koss' Diagnostic Cytology and its Histopathologic Basis. 5th ed.Philadelphia:LipincottWilliams& Wilkins;2006. p. 422-65

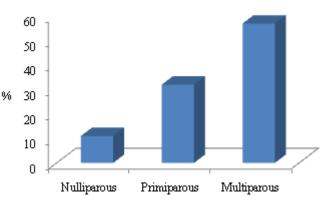


Figure 1: Distribution of patients according to parity

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<40	40-50	>50	Total
8 (12)	56 (88)		64 (29)
28 (51)	23 (42)	4 (7)	55 (25)
	5 (42)	6 (58)	11 (5)
2 (46)	3 (54	6	11 (5)
5 (33)	9 (60)	1 (7)	15 (7)
		2 (100)	2(1)
12 (41)	16 (55)	1 (3)	29 (13)
3 (27)	8 (73)		11 (5)
3 (14)	11 (50)	8 (36)	22 (10)
61 (28)	131 (59)	28 (13)	221 (100)
	28 (51) 2 (46) 5 (33) 12 (41) 3 (27) 3 (14)	<40 40-50 8 (12) 56 (88) 28 (51) 23 (42) 5 (42) 5 (42) 2 (46) 3 (54 5 (33) 9 (60) 12 (41) 16 (55) 3 (27) 8 (73) 3 (14) 11 (50)	$\begin{array}{c c c c c c c c c c c c c c c c c c c $