

To Study And Evaluate the Benign, Pre Malignant And Malignant Endometrial Etiology Of Post Menopausal Bleeding Per Vaginum

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Submitted: 01-04-2022

Accepted: 11-04-2022

I. INTRODUCTION

Postmenopausal bleeding is defined as any bleeding from the genital tract after 1 year of menopause. However, vaginal bleeding occurring anytime after 6 months of amenorrhoea in menopausal age should be considered as postmenopausal bleeding and investigated ¹. It is a common gynaecological problem seen in about 5 - 10 % of all gynaecological patients.

Endometrial atrophy is the most common cause of vaginal bleeding among postmenopausal women, whereas endometrial hyperplasia and polyps are other common causes².

The dictum is "postmenopausal bleeding indicates malignancy until proved otherwise". About 90-95% of postmenopausal women with endometrial cancer experience a vaginal bleeding, whereas about 10% of symptomatic postmenopausal women reveal an intrauterine malignancy². Thickenedendometrium may indicate the presence of endometrial pathology. The investigations of PMB are pelvicultrasound and tissue biopsy. TVS is considered as an acceptable non invasive initial investigation in women with PMB. Hysteroscopy is considered the gold standard for the diagnosis of cervical and intrauterine pathogenesis, and also has a therapeutic role in different benign lesions³. The current study was carried out to evaluate various endometrial causes for Postmenopausal bleeding.

AIMS & OBJECTIVES:

1.To analyse various endometrial causes for postmenopausal bleeding.

2. To differentiate benign and malignant endometrial causes for postmenopausal bleeding histopathologically.

II. MATERIALS & METHODS:

This is a hospital based prospective observational study conducted in the department of Obstetrics & Gynaecology, ACSR govt medical college and hospital, Nellore over a period of one year from january 2019 to january 2020. 408 post menopausal women with bleeding per vaginum were taken for study.

Inclusion criteria:

- 1. Postmenopausal women ≥ 40 yrs of age.
- 2. Amenorrhoea of one year.

3. Postmenopausal women presenting with bleeding per vaginum.

Exclusion criteria:

- 1. Bleeding disorders.
- 2. Genital tract and vulval injuries.

3. Vulval, vaginal and cervical lesions (both benign and malignant).

An informed consent was taken and complete medical history was obtained. Physical examination was done. Ultrasonographyand endometrial biopsy were taken from women with no obvious cervicalgrowth. Data were collected and tabulated as shown in results. Statistical analysis was done in Microsoft Excel.Frequency and percentage of each parameter was calculated and analysed.

III. RESULTS :

• Post menopausal bleeding – Incidence:

A total of 5013 women with various gynaecological disorders were seen in both outpatient and inpatient departments during thestudy period, of which 408 women were presented with post menopausalbleeding, giving an incidence of 8.13%.





Table no.1: PMB – Endometrial thickness

Endometrial thickness (mm)	n/t	%
< 4 mm	138/408	34
4 –10mm	119/408	29
>10mm	151/408	37

Table 2. I Mib Instopationogy of Endometrian	Table 2 : PMB –	Histopathology of Endometrium	
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Table 2 : FMB – Histopathology of Endometrium					
Endometrial pathology	n/t	%			
Proliferative phase	126/408	31			
Atrophic endometrium	140/408	34			
Chronic endometritis	20/408	5			
Endometrial polyp	53/408	13			
Simple hyperplasia without atypia	19/408	4.6			
Simple hyperplasia with atypia	11/408	2.6			
Complex hyperplasia without atypia	25/408	6.1			
Complex hyperplasia with atypia	02/408	0.4			



International Journal Dental and Medical Sciences Research Volume 4, Issue 2, Mar-Apr 2022 pp 411-414 www.ijdmsrjournal.com ISSN: 2582-6018

Adenocarcinoma	of	12/408	3	
endometrium				

IV. DISCUSSION :

The incidence of post menopausal bleeding in our study is 8.13% and is correlating with the study done by Hyen Chul Jo etal $(10\%)^4$.

ENDOMETRIAL THICKNESS

In the present study, 66% of the women withpostmenopausal bleeding had endometrial thickness ≥ 4 mm. Endometrial thickness < 4mm correlates with lowrisk of endometrial pathology. If there is diffuse increase in endometrium more than20 mm, it indicates endometrial pathology. If there is focal increase in thickness, itindicates that there is a possibility of endometrial polyp which can be diagnosedmore accurately by sonohysterography.

BENIGN ENDOMETRIAL LESION:

ENDOMETRIAL POLYP: In the present study, 13% of the women with PMB had endometrial polyp, which is correlating with the study done by Ghoubara A etal $(14\%)^4$ and Min Kyoung Kim etal $(11\%)^8$.

ATROPHIC ENDOMETRIUM

In the present study, 34% of the women with PMB had atrophic endometrium, which is correlating with the studydone by Yi Liang Lee etal $(42\%)^6$.

In the benign endometrial lesions, most commonis atrophic endometrium (34%), followed by proliferative endometrium (31%). The sclerotic degeneration of endometrial vessels / local abnormal hemostaticmechanism in the uterus may cause bleeding from the atrophic endometrium.

PROLIFERATIVE ENDOMETRIUM: In the present study, 31% of the women with PMB had proliferative endometrium, which is correlating with thestudy done by Jasmina B etal (35%)². In some women, endometrium shows abnormal response tocirculating low level of estrogen leading to proliferative endometrium and presents aspost menopausal bleeding.

PREMALIGNANT ENDOMETRIAL LESION

In the present study, 14% of the women with PMB had pre malignant endometrial lesion, which is correlating withthe study done by Ghoubara etal (10%)⁴.Endometrial hyperplasia is a premalignant condition which carries both clinical and pathological significance and predisposes toendometrial carcinoma. The risk of endometrial carcinoma is high with atypicalchanges which requires definitive treatment.

ENDOMETRIAL CARCINOMA

In the present study, 3% of the women with PMB had endometrial carcinoma, which is correlating with the studydone by Maria BB etal $(4.66\%)^{7}$. Hyen Chul Jo etal $(5.1\%)^{5}$ and Min Kyoung Kim etal $(5.7\%)^{8}$. The risk of endometrial carcinoma increases with age. Approximately 1% of the cases are seen at the age of 50 years whichincreases to 25% at the age of 80 years. In the present study, 83% of theendometrial carcinoma were seen in > 70 years of age.

V. CONCLUSION

PMB is seen in about 5 - 10 % of gynaecological patients. It is important to rule out the underlying malignancy as a cause of post menopausal bleeding . With the increasing life expectancy aftermenopause, more and more number of endometrial cancers are to be expected in thefuture. There is no effective screening programme for endometrial cancer. Women with high risk factors like obesity, Diabetes, women on Tamoxifene or on HRT will require periodic assessment of endometrial thickness and if required, endometrial aspiration cytology is to be done to exclude endometrial pathology.To reduce the incidence of genital tract malignancy in post menopausal women, regular screening with pap smear upto the age of 70 years and USG assessment of endometrial thickness upto the age of 80 years isrecommended.

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