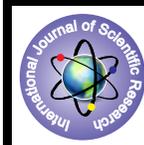


Histopathological Study of Endometrium in Dysfunctional Uterine Bleeding



Medical Science

KEYWORDS : Dysfunctional uterine bleeding (DUB), Endometrium, Endometrial hyperplasia.

DR. MEETA.G.NANAVATI	Professor & HOD (PATHOLOGY) GMERS medical college Gandhinagar
DR. JITENDRA. H. PARIKH	Associate professor (Department Of Medicine B. J. Medical College, Ahmedabad, Gujarat.)
DR. VARAD BISEN	M.D (Pathology) Department Of Pathology, B. J. Medical College, Ahmedabad, Gujarat.

ABSTRACT

BACKGROUND: Dysfunctional uterine bleeding is defined as excessively heavy, prolonged, or frequent bleeding of uterine origin that is not due to pregnancy or any recognizable pelvic or systemic cause. It is therefore a diagnosis of exclusion.¹ Dysfunctional uterine bleeding is a common problem in the women in the age 30- 50 years. The incidence increases as age advances till menopause.² It is also seen in perimenarchal and perimenopausal age group.

AIMS AND OBJECTIVES:

- 1) To study endometrial cytomorphology in cases of DUB.
- 2) To study correlation of different clinical presentation and different clinical finding with histopathology.
- 3) To correlate age and parity with histopathological finding.
- 4) To study incidence of various hyperplasia in women with DUB.

MATERIAL AND METHOD:

The study was done during periods of 2011- 2012, 410 cases were taken. In this study endometrium specimen were obtained by D& C and hysterectomy. Sections were stained by H&E stain and histopathological examination was done.

CONCLUSION :

Maximum numbers of DUB patients were in the age group 31-40 years. Most common presenting complaint was menorrhagia. Most of patients with DUB shows proliferative with hyperplastic pattern. Simple hyperplasia was more common than complex and atypical hyperplasia. As age advances incidence of hyperplasia was increased. Parity has no influence on endometrial pattern in DUB.

INTRODUCTION:

Dysfunctional uterine bleeding is defined as excessively heavy, prolonged, or frequent bleeding of uterine origin that is not due to pregnancy or any recognizable pelvic or systemic cause. It is therefore a diagnosis of exclusion.¹ Dysfunctional uterine bleeding is a common problem in the women in the age 30- 50 years. The incidence increases as age advances till menopause.² It is also seen at perimenarchal and perimenopausal age group.

The pathophysiology of DUB is not fully understood and it is complex.¹ The mechanisms for the abnormal bleeding and the site from which it arises are largely unknown. Menstruation is a very complex process involving oestrogen and progesterone and their receptors, endometrial vasculature, endometrial vasoactive substances, processes of tissue breakdown and remodelling and endometrial repair regeneration.¹ It accounts for above one third of all gynaecological consultations carried out for abnormal uterine bleeding.³ Menstrual disturbance is one of the commonest gynaecological problems for which curettage or hysterectomy specimen is received by pathologist. Over the past 10-15 years there has been a major escalation as the scientific efforts aimed at elucidating the underlying mechanism. It is now clear that these mechanisms are complex and involving a number of different molecular systems⁵. Being most common gynaecological problems, present study aimed to know full spectrum of DUB and its pathological aspects at this institute.

AIMS AND OBJECTIVES:

- 1) To study endometrial cytomorphology in cases of DUB.
- 2) To study correlation of different clinical presentation and different clinical finding with histopathology.
- 3) To correlate age and parity with histopathological finding.
- 4) To study incidence of various hyperplasia in women with DUB.

MATERIAL AND METHOD:

The study was done during periods of 2011- 2012 in B.J Medical

college ahmedabad. 410 cases were taken. In this study endometrium specimen were obtained by D& C and hysterectomy. Sections stained by H&E stain and histopathological examination was done.

RESULTS AND OBSERVATIONS

410 cases were studied in present series which were clinically diagnosed as DUB with exclusion of structural lesions by radiological investigations. Out of which 10 cases turned out to be some organic cause for the bleeding on histopathological examination. Those 10 cases (4 leiomyoma, 5 adenomyosis and 1 endometrial carcinoma) were excluded from present series and further study was carried out on 400 cases proven histopathologically also as DUB.

Table 1: Incidence of DUB in various age group in our study

AGE (YEARS)	NO OF CASES	PERCENTAGE(%)
≤ 20	2	0.5
21-30	64	16
31- 40	192	48
41-50	124	31
51- 60	18	4.5
TOTAL	400	100

Above table shows that largest no of patients were in 31-40 years(48%) with average age being 38.7 years. Next major group belonged to 41- 50 years (31%). The percentage of patients in age <30 years was 16.5%. Incidence of DUB was maximum in 31-40 years.

Table 2: Incidence of menstrual disorder in DUB

MENSTRUAL DISORDER	NUMBER OF CASES	PERCENTAGE (%)
Menorrhagia	206	76.5
Polymenorrhoea	32	8
Menorrhagia	54	13.5
Oligomenorrhea	8	2
Total	400	100

Above table shows that maximum incidence was of menorrhagia(76.5%) followed by Menorrhagia (13.5%)

TABLE 3: Endometrial pattern in DUB

ENDOMETRIAL PATTERN	NO OF CASES	PERCENTAGE(%)
Proliferative	159	39.75
Secretory	125	31.25
Hyperplasia	88	22
Atrophy	28	7
Total	400	100

Table 3 shows in present series proliferative phase was most common and found in 39.75% cases. Secretory phase was found in 31.25% and hyperplasia was seen in 22% cases.

Table 4: Incidence of types of hyperplasia associated with DUB

TYPES OF HYPERPLASIA	NO OF CASES	PERCENTAGE(%)
Simple	60	68.18
Complex	24	27.27
Atypical	4	4.55
Total	88	100

Above table shows that most common incidence of hyperplasia with DUB was simple hyperplasia followed by complex .Atypical hyperplasia was rare finding.

Table 5: Endometrial pattern in relation to Age:

ENDOMETRIAL PATTERN	AGE (YEARS)									
	≤20 YEARS		21-30 YEARS		31-40 YEARS		41-50 YEARS		51-60 YEARS	
	No	%	No	%	NO	%	NO	%	NO	%
Proliferative phase	2	100	32	50	80	41.66	44	35.48	1	0.555
Secretory phase	-	-	26	40.63	70	36.45	28	22.58	1	0.555
Simple hyperplasia	-	-	4	6.25	30	15.65	22	17.74	4	22.22
Complex hyperplasia	-	-	-	-	6	3.12	16	12.91	2	11.11
Atypical hyperplasia	-	-	-	-	2	1.04	-	-	2	11.11
Atrophic	-	-	2	3.12	4	2.08	14	11.29	8	44.44
Total	2	100	64	100	192	100	124	100	18	100

Above table shows proliferative phase was common in ≤ 30 years age group. Secretory phase was common in 21- 30 years age group. Hyperplasia was common in > 40 years age group. Atrophy was also common in > 40 years age group.

Table 6: Endometrial pattern in relation to parity

ENDOMETRIAL PATTERN	PARITY					
	≤2		3-4		≤5	
	NO	%	NO	%	NO	%
Proliferative phase	76	42.18	64	37.65	22	44
Secretory phase	68	37.74	50	20.41	10	20
Hyperplasia	28	15.64	46	27.06	14	28
Atrophic	8	04.44	10	05.88	4	08
Total	180	100	170	100	50	100

Table 7: Age incidence comparison with other series

AGE (YEARS)	OUR STUDY(%) 400 Cases	Muhammad et al. ³ 260 cases	Sutherland et al. ⁴ 848 cases
≤ 20	0.5	-	3.9
21-30	16	12.7	22.5
31-40	48	39.2	34.5
41-50	31	48.1	37.7
51-60	4.5	-	1.6

Above table shows that in our study maximum number of a pa-

tients were in 31-40 years age group (48%) and 4.5% patients belonged to > 50 years age group ,0.5% belonged to ≤ 20 years age which is comparable to other series. Low incidence was found in ≤ 20 years age group (0 .5%), however actual incidence of DUB is not low in this group. They are treated on conservative (hormonal) basis and most often they are unmarried and they do not undergo endometrial sampling.

Table no 8: Endometrial pattern comparison with other series.

Endometrial pattern	Our series		R.K.Narula et al study ¹⁵		Sanaullah et al study ¹⁶	
	NO	%	NO	%	NO	%
Proliferative phase	159	39.75	83	37.77	31	31
Secretory phase	125	31.25	79	35.95	43	43
Hyperplasia	88	22	46	20.90	11	11
Atrophic	28	7	12	5.46	-	-
Others	-	-	-	-	15	15
Total	400	100	220	100	100	100

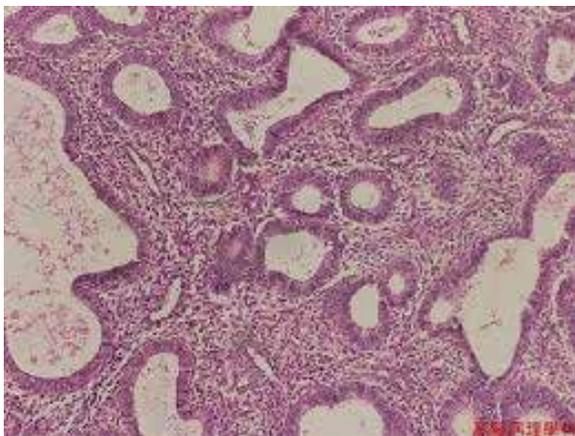
Proliferative pattern is commonest in our series (39.75%) which is nearly comparable with R.K. Narula et al study. It is evident

from above table that large proportions of patients showed proliferative or hyperplastic endometrial pattern. Incidence of hyperplasia is comparable in two studies.

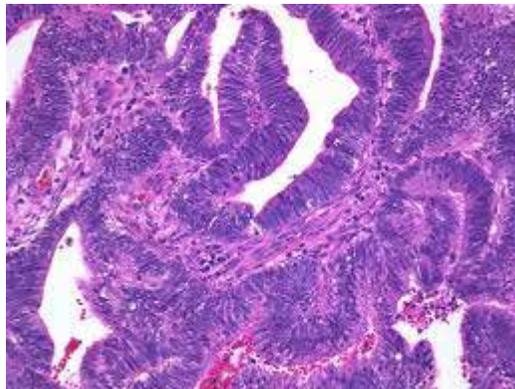
Table 9: Incidence of types of Hyperplasia comparison with other series:

TYPES OF HYPERPLASIA	OUR STUDY		Muhammad et al study ³		Sanaulla et al study ¹⁶	
	NO	%	NO	%	NO	%
	Simple	60	68.18	39	60.93	9
Complex	24	27.27	22	34.39	2	18.18
Atypical	4	4.55	3	4.68	-	--
Total	88	100	64	100	11	100

Above table shows that in our series maximum number of patients (68.18%) had simple hyperplasia of Endometrium which is comparable with other series. In our study atypical hyperplasia was less common (4.55%) which is also comparable with Muhammad et al study. Overall finding show that simple hyperplasia was more common than complex hyperplasia and incidence of atypical hyperplasia was very low.



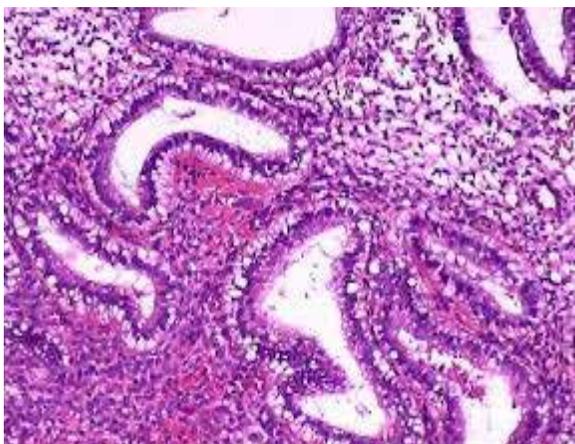
SIMPLE CYSTIC HYPERPLASIA



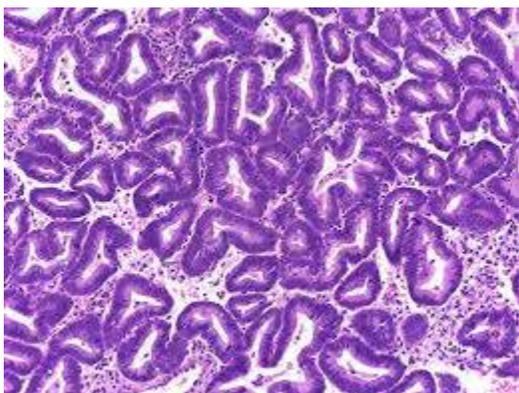
ENDOMETRIAL ADENOCARCINOMA

CONCLUSION :

Maximum numbers of DUB patients were in age group 31-40 years. Most common presenting complaint was menorrhagia. Most of patients with DUB shows proliferative with hyperplastic pattern. Simple hyperplasia was more common than complex and atypical hyperplasia. As age advances incidence of hyperplasia was increased. Parity has no influence on endometrial pattern in DUB.



SECRETORY PHASE



COMPLEX HYPERPLASIA

REFERENCE

1) Elizabeth Farell .Dysfunctional uterine bleeding ,Australian Family Physical ,2004 ;33(11):906-08 | 2) George A.Vilos ,Guylain Lefebvre Gillin R.Graves,Halifax ,N.S.Guidelines for the management of abnormal uterine bleeding SIOGC Clinical practice guideline :NO 106 aug 2001. | 3) Muhammad Muzaffer , Khalida Adeeb Khanam Akhter Shahima Yasmin ,Mahmood –ur –Rehman ,Wasim Iqbal : Menstrual irregularities with excessive blood loss :a clinic- pathological Correlation .JPMA .,2005:55-486. | 4) Divya Makhija ,Alka Mary Mathai ,Ramdos Naik, Suneet kumar, Sharda Rai, Mukhta P. Rai ,Poornima Baliga. Morphometric evolution of endometrial blood vessels. Indian Journal of pathology and Microbiology .2008:51(31):436-350. | 5) Mark Livingstone ,Jan S Fraser . Mechanism of abnormal uterine Bleeding . Human Reproduction Update, 2002 :8(1):60 -67. | 6) I an S. Fraser, Hilary O.D.Critchely ,Malcom G Munero .Abnormal uterine bleeding :getting our terminology straight. Curr Opin Obs. Gynecolo.2007:19:591-595. | 7) Juan Rosai ,Ackerman's surgical pathology ,9th edition chap 19 :1569-1583. | 8) Barbara young, James S Lowe Alans Stevens John W. Health. Whetear's Functional histology .5th edition cap 19:359- 391. | 9) D.C. Dutta Text Book Of Gyenecology , chap 15 :218-222 | 10) Arthur C. Guyton John . E. Hall. Text book of Medical Physiology .11TH EDITION ,chapter 81:1011-1025. | 11) D.C Dutta Text Book Of Gyenecology chap 1:1-11. | 12) Jamie E. Siegel Abnormalities of hemostasis and abnormal uterine bleeding clinical obstetrics and gynecology .2005:48(2):284-294. | 13) Robbins and cotran .Pathologic Basis of Disease ,7th edition chapter -22.1079-1086. | 14) Sternberg 's Diagnostics Surgical pathology .4th edition vol.3 Section X: Female reproductive system and peritoneum chap 53.2435- 2475. | 15) R. K. Narula. Endometrial histopathology in DUB. Journal of obstetrics & Gynecology of India.1967:17:614-18 | 16) Sanaullah Gazozai , Qurban Ali Bugti ,Aysha ,Siddique ,Naila Ehsan .Excessive Uterine haemorrhage – A histopathological study .Gomal Journal of medical Science Jan –june 2004:2(1): 13-15. |