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# Correlation of Pap Smear with Histopathological Findings in Malignant and non Malignant Lesions of Cervix Dr. Vijay Kumar Bodal<sup>1</sup> <sup>1</sup> Department of Pathology, Government Medical College Patiala, (Punjab) India *Received: 12 December 2013 Accepted: 2 January 2014 Published: 15 January 2014*

#### 7 Abstract

<sup>8</sup> Background: Conventional cervical cytology is the most widely used cervical cancer screening

<sup>9</sup> test in the world. Squamous intraepithelial neoplasia (SIL) and cervical cancer remain

<sup>10</sup> important health problems for women worldwide. Aim and Objective: To study various types

of cervical lesions with relevant factors such as age, parity, to classify cervical lesions into

<sup>12</sup> malignant benign groups and to correlate the cytological with histopathological

13 findings.Materials and Methods: This study was conducted on 200 cases of Pap smears and

<sup>14</sup> cervical biopsies, along with resected specimens. After fixation and staining, smears and

<sup>15</sup> cervical biopsies were processed and examined under microscope.

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17 Index terms— malignant, cervical cancer, pap smear, cervical biopsy.

# Correlation of Pap Smear with Histopathological Findings in Malignant and non Malignant Lesions of Cervix

Introduction apanicolaou (Pap) smear is a simple, safe, noninvasive and effective method for detection of 20 precancerous, cancerous and non-cancerous changes in the cervix. [1] Conventional cervical cytology is the most 21 widely used cervical cancer screening test in the world and cytology screening programmes in several developed 22 countries have been associated with impressive reduction in cervical cancer burden. [2] Squamous intraepithelial 23 lesions are viewed as precancerous lesions exhibiting many of the morpho-Author ?: Department of Pathology, 24 Government Medical College Patiala, (Punjab) India. e-mail: vijay\_bodal@yahoo.com logical characteristics 25 of invasive carcinomas. Identification of these entities is the focus of cervical screening programs that aim to 26 discover them and commence their treatment in order to prevent invasive disease. [3] Though data from the 20 27 populations based cancer registries in India indicate a steady decline in cervical cancer incidence rates over the last 28 two decades, it still occupies second position and the risk of disease is still high. [3] Cervical carcinoma documents 29 the remarkable effects of screening, early diagnosis, and curative therapy on the mortality rate. Death rate has 30 declined for which the credit goes to Pap test and accessibility of cervix to colposcopy and biopsy. Though, the 31 Pap smear is an effective screening test, yet confirmation of the diagnosis of cervical cancer or pre invasive lesions 32 of cancer requires a biopsy of the cervix. 33

#### 34 **2** II.

#### 35 3 Aims and Objectives

The aims of this study were to study the changes in cervical cytology with relation to age, parity and other presenting features, to classify cervical lesions into malignant and benign groups on cytological and histopathological basis and to correlate the changes observed in cervical cytology with cervical biopsy.

#### <sup>39</sup> **4 III.**

#### 40 5 Materials and Methods

This study was done on 200 cases of Pap smears and cervical biopsies (including hysterectomy specimens). Most 41 of the patients with symptoms suggestive of cervical disease were selected. However, some having gynaecological 42 symptoms other than cervical disease were also included. Few cases reporting for routine screening were also 43 included. A detailed clinical history especially age, duration of symptoms, parity, menstrual pattern and vaginal 44 discharge were noted. The patients in whom both Pap smear and biopsy was available, were included in the 45 study. The fixed cervical smears were subjected to staining according to Papanicolaou's method. The cytological 46 interpretation of the smears was made according to the New 2001 Bethesda system. After grossing and processing, 47 cervical biopsies were subjected to histopathological examination. 48

#### 49 6 Results

Age wise maximum number of patients were in fourth decade (54.50%), followed by fifth decade (Table-1). 50 Duration of symptoms varied from few months to many years. Some patients presented within 1 year (79%), 51 but few mainly cases with discharge and history of prolapse presented late (Table ??2). In 200 cases, various 52 symptoms were seen, some patients showed multiple symptoms. Majority of patients (58%) presented with 53 vaginal discharge followed by irregular bleeding (47%). Menstrual changes were also seen in large number of 54 patients. There was seen low usage of oral contraceptive pills in our study group (10.50%). Duration of OCP 55 usage varied from few months to years, but long term usage was not seen in any case. On cytology, 59% were 56 inflammatory smears and frank malignancy was reported in 10% cases, LSIL and HSIL was reported in 9% and 57

58 8.50% respectively (Table ??

#### <sup>59</sup> 7 Discussion

60 Cancer cervix is considered to be an ideal gynaecological malignancy for screening as it meets both test and disease 61 criteria for screening. It has a long latent phase during which it can be detected as identifiable and treatable 62 premalignant lesions which precede the invasive disease and the benefit of conducting screening for carcinoma

cervix exceeds the cost involved. [4] Despite the success of cervical cancer screening programs, questions remain

<sup>64</sup> about the appropriate time to begin and end screening. This review explores epidemiologic and contextual data

on cervical cancer screening to inform decisions about when screening should begin and end. The incidence and mortality rates from, cervical cancer that have had a Pap smear within 3 years have decreased since 2000.

mortality rates from, cervical cancer that have had a Pap smear within 3 years have decreased since 2000.
In this study, more than half (54.50%) were aged between 31 to 45 years followed by 20.50% between 46 to 60
years. The mean age of patients with cancer in the present study was 51.94 years. This is close to that found by
Biswas et al [5] and Missaoui et al. [6] Although, invasive cancer cervix is reported at all ages; it has two peaks,
one at about 35 years and another above 50 years. The highest age of cervical cancer in the present study was

71 73 years and the lowest was 26 years. The mean age for non-cancer cases was 39.53 years. In this study, the

most common symptoms was discharge per vaginum (58%) followed by irregular bleeding in 47% of the patients.
Patients with cancer also presented with post-coital bleeding and in cases of older age group post menopausal

 $^{74}$   $\,$  bleeding was seen. Symptomatic presentation was similar to some extent as seen by Ikram et al [7] .

In this study, 59% patients had the cytological diagnosis of benign/ inflammatory and carcinoma was present in 10% of the cases. This is comparable to Saha and Thapa [8] in which benign cases were 51.16% and carcinoma was diagnosed in 6.97% of the cases. Most common cancer in the present study was squamous cell carcinoma (85.18%). This study showed results similar to those seen by Ikram et al [7] (83.33%).

As regards the various histopathological varieties of SCC, the present study found an incidence of 67.39% for moderately differentiated SCC, 23.91% for well differentiated, 8.70% for poorly differentiated. Thus, the findings of the present study are consistent with that

#### 82 8 Conclusions

It is concluded that most commonly seen problem, infection, can be controlled with good hygiene. Cervical carcinoma is seen in large number of patients. Pap is a relatively less invasive and a simple procedure to diagnose cervical lesions in developing countries. But sometimes, there can be obscuring of the cellular details by blood, especially in malignant cases. In such cases, biopsy is helpful and confirmatory. of Missaoui et al [6] in that moderately differentiated large

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Figure 1: P



Figure 2:



Figure 3: Figure 1 :



Figure 4: Figure 2 : Figure 3 : Figure 4 :



Figure 5:

#### 1

Age group (Years) 18-30 31-45 46-60 > 60	Distribution (n=200) No.	%age 29	9 14.50 109 54.50 41 20.8
Total	200		100
Table 2 : showing duration of symptoms	5		
Duration (Years)	Distribution $(n=200)$ No.		%age
Up to 1	158		79.00
1-3	25		12.50
4-6	11		05.50
>6	06		03.00
Total	200		100
Table 3 : showing cytological diagnosis			
Diagnosis	Distribution (n= $200$ ) No. %age		
Unsatisfactory smear	08	4.00	
Inflammatory	118	59.00	
ASCUS/H	19	9.50	
LSIL	18	9.00	
HSIL	17	8.50	
Frank malignancy	20	10.00	
Total	200	100	
	Age group (Years) 18-30 31-45 46-60 > 60 Total Table 2 : showing duration of symptoms Duration (Years) Up to 1 1-3 4-6 >6 Total Table 3 : showing cytological diagnosis Diagnosis Unsatisfactory smear Inflammatory ASCUS/H LSIL HSIL Frank malignancy Total	Age group (Years) 18-30 31-45 46-60 >Distribution (n=200) No.60200Total200Table 2 : showing duration of symptomsDistribution (n=200) No.Duration (Years)Distribution (n=200) No.Up to 11581-3254-611>606Total200Table 3 : showing cytological diagnosisDistribution (n=200) No.DiagnosisDistribution (n=200) No.Unsatisfactory smear08Inflammatory118ASCUS/H19LSIL18HSIL17Frank malignancy20Total200	Age group (Years) 18-30 31-45 46-60 >Distribution $(n=200)$ No. %age 2960Total200Table 2 : showing duration of symptomsDuration (Years)Distribution $(n=200)$ No.Up to 11581-3254-611>606Total200Table 3 : showing cytological diagnosisDiagnosisDistribution $(n=200)$ No. %ageUnsatisfactory smear084.611>606Total200Table 3 : showing cytological diagnosisDiagnosisDistribution $(n=200)$ No. %ageUnsatisfactory smear084.00Inflammatory118ASCUS/H199.00HSIL178.50Frank malignancy20100

### Figure 6: Table 1 :

#### $\mathbf{5}$

Histopathological No.	Cytological Diagnosis Unsatis	sfactory Infla	ammatory	ASCUS/H LSIL	HSIL
Diagnosis					
Infections 115	-	108	07	-	-
Carcinoma 54	08	-	-	12	14
Dysplasia 25	-	04	12	06	03
Benign tumors 06	-	06	-	-	-
Total 200	08	118	19	18	17
	Table 6 : showing means age				
Variable	Cervical Ca $(n=54)$ Mean SD	)	No Ca	(n=146) Mean	$\mathbf{SI}$
Mean	51.94	12.30	39.53		09.66
Т		7.469			
Df		198			
Р		< 0.001			
Significance V.		Highly Significant			

Figure 7: Table 5 :

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