



## Colposcopy in Inflammatory PAP SMEARS

**Dr Aakanksha  
agarwal**

M.S Resident , Department of Obs & Gyane, NIMS Medical College & Hospital,  
NIMS University, Jaipur

**Dr Beena  
Bhatnagar**

HOD and professor, Department of Obs & Gyane, NIMS Medical College &  
Hospital, NIMS University, Jaipur

**Dr Priyanka Suhag**

DNB Resident ,Department of Obs & Gyane, NIMS Medical College & Hospital,  
NIMS University, Jaipur

**ABSTRACT****INTRODUCTION**

*Cervical is the second most common cancer in the world and the leading cause of death. Frequency and mortality are significantly reduced by screening through cytological Papanicolaou test. However PAP test has low sensitivity and a high negative rate of 9-40%. Epithelial abnormalities on PAP test can be managed according to ACCP guidelines. However for management of inflammatory smears there are no such algorithms available. The incidence of inflammatory PAP smears is very high 14-19%. and subjecting each patient to colposcopy and HPV DNA testing might become costly. This study was done to evaluate inflammatory PAP smears with colposcopy and directed biopsy to find out any premalignant changes in the cervix and whether colposcopy should be done to help triage these women.*

**MATERIAL AND METHOD**

*100 women were screened at NIMS Hospital with PAP smear. However only those women who had inflammatory smears were subjected to colposcopy and directed biopsies.*

**RESULT**

*Out of 91 patients , 28 women (30%) had normal colposcopy, and 63 women (70%) had abnormal colposcopy of which 6.5% had polyp, 15.38% had CIN1, 3.29% had CIN 2, 3.29% had CIN 3 and 1.09% had invasive cancer.*

**CONCLUSION**

*Nearly 70% women had abnormal colposcopy with PAP smear showing inflammatory smear. Hence a large no of women with abnormal colposcopy and CIN (around 10%) would be missed if inflammatory PAP smears are not evaluated further.*

**KEYWORDS : Colposcopy, Inflammatory Pap smear, CIN**

**INTRODUCTION**

Cervical is the second most common cancer in the world and the leading cause of death. Frequency and mortality are significantly reduced by screening through cytological Papanicolaou test. However PAP test has low sensitivity and a high negative rate of 9-40%.<sup>1,2,3</sup> Epithelial abnormalities on PAP test can be managed according to ACCP guidelines. However for management of inflammatory smears there are no such algorithms available. The incidence of inflammatory PAP smears is very high 14-19%.<sup>4,5,6,7</sup> and subjecting each patient to colposcopy and HPV DNA testing might become costly. This study was done to evaluate inflammatory PAP smears with colposcopy and directed biopsy to find out any premalignant changes in the cervix and whether colposcopy should be done to help triage these women.

**MATERIAL AND METHOD**

The study was conducted at NIMS Hospital and Medical College, a tertiary level teaching hospital after obtaining clearance from the Ethical Committee of the Institution. 100 women presenting with symptoms like discharge per vaginum, pain in lower abdomen, lower backache, pruritis vulvae or vagina were enrolled. Women who were pregnant or postmenopausal, with proven CIN or obvious cervical growth, IUCD users, with diabetes mellitus, and with multiple sexual partners were excluded from the study. Informed consent was taken from all women. After a detailed history and general physical examination, a detailed gynecological examination was carried out in all. This included examination of the breast, local examination of vulva, per speculum examination with Pap smear, and per vaginum examination. Women with inflammatory Pap smear were then included in the study. They were then subjected to colposcopy. The presence of any acetowhite or iodine negative areas was noted and classified according to colposcopic terminology by IFCPC 8 [Table 1]. The patients with suspicious lesions on colposcopy were then subjected to cervical biopsy from these abnormal areas along with endocervical brush cytology and

endocervical curettage. The biopsy specimens were fixed in 10% formalin and sent for histopathological evaluation. The results were then interpreted. The statistical analysis was done using Student's *t*-test.

**Colposcopic evaluation by IFCPC table 1**

|     |  |
|-----|--|
| I   | Normal colposcopic findings                        |
|     | Original squamous epithelium                       |
|     | Columnar epithelium                                |
|     | Transformation zone                                |
| II  | Abnormal colposcopic findings                      |
|     | Flat acetowhite epithelium                         |
|     | Dense acetowhite epithelium                        |
|     | Fine mosaic  |
|     | Coarse mosaic                                      |
|     | Fine punctation                                    |
|     | Coarse punctation                                  |
|     | Iodine partial positivity                          |
|     | Iodine negativity                                  |
|     | Atypical vessels                                   |
| III | Colposcopic features suggestive of invasive cancer |
| IV  | Unsatisfactory colposcopy                          |
|     | Squamocolumnar junction not visible                |
|     | Severe inflammation, severe atrophy, trauma        |
|     | Cervix not visible                                 |

### V Miscellaneous findings

Condylimata

Keratosis

Erosion

Inflammation

Atrophy

Deciduosis

Polyps

### RESULT

one hundred women were recruited from the OPD in our hospital. Out of 100 women, 91 women (91%) had inflammatory smear. The mean age of the women with inflammatory Pap smear was  $37.99 \pm 7.92$  years. The mean age at onset of sexual activity was  $16.98 \pm 1.48$  years. Out of 91 patients, 28 women (30%) had normal colposcopy, and 63 women (70%) had abnormal colposcopy of which 6.5% had polyp, 15.38% had CIN1, 3.29% had CIN 2, 3.29% had CIN 3 and 1.09% had invasive cancer. No tissue was identified in the endocervical curettings in any patient and the endo-cervical brush cytology showed only inflammation. On correlating the histopathology reports with the colposcopic findings, it was found that the women with chronic cervicitis had grade I-II acetowhite changes and fine mosaic and punctation pattern, while the patients with CIN had grade II-III acetowhite change with coarse punctation.

### DISCUSSION

Inflammation on Pap smear is a very common finding. Its prevalence in various Indian studies is reported to vary between 70% and 80.5%.<sup>6,9</sup> In our study the prevalence is 91%.

The mean age of the women with inflammatory Pap smear was  $37.99 \pm 7.92$  years. The mean age of women with CIN in our study was  $41 \pm 7.86$  years ( $P=0.1188$ ).

The mean age at onset of sexual activity in inflammatory pap smear was  $16.98 \pm 1.48$  years where as the mean age of onset of sexual activity in women with CIN in our study was  $16.57 \pm 1.47$  years ( $p=0.2544$ ).

The mean parity was  $3.167 \pm 0.974$  and  $3.142 \pm 1.014$ , respectively ( $P=0.9164$ ).

In our study out of 91 with inflammatory pap smear, 21.96% had CIN and 1.09% had invasive cancer. Various studies have found the possibility of CIN with a report of inflammatory smear to range from 18% to 35%.<sup>4,5</sup>

According to various studies, ASCUS on Pap smear has a 10%-20% chance of harboring CIN.<sup>10,11</sup> This is the reason why we triage women with ASCUS on Pap smear with either repeat cytology, HPV DNA testing, or colposcopy.<sup>12</sup> Our study has shown that 7.2% of women with inflammation on Pap smear could be harboring CIN. Hence, all women with inflammation on Pap smear should be subjected to further evaluation. HPV DNA testing can be an alternative to colposcopy; however, the use of HPV DNA testing to triage women with inflammatory pap smears has not been recommended. Further studies are required to establish the cost effectiveness of HPV DNA testing in triage of women with inflammatory pap smear.

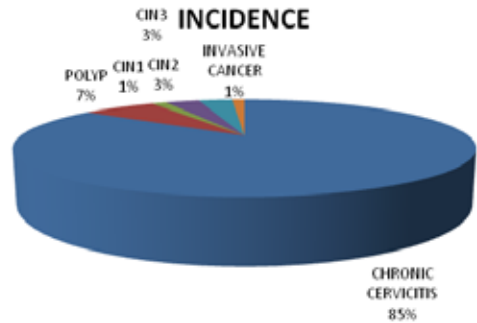
In our study, we found a 21.96% incidence of CIN in women with inflammatory pap smear. The incidence of CIN and invasive carcinoma in women with persistent inflammatory pap smears over just 2 weeks was found to be 20.6% and 0.7%, respectively, in a study by Dasari *et al.*<sup>13</sup> Hence, by waiting for a longer period of time before repeating the pap smear may lead to a delay in diagnosis of CIN in a high percentage of cases.

To conclude, one must not see a report of inflammation on Pap smear in isolation and ignore it as being absolutely insignificant. If it persists in any patient then we must consider evaluating the patient further by colposcopy.

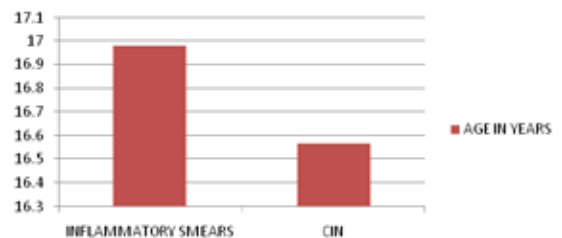
### CONCLUSION

Nearly 70% women had abnormal colposcopy with PAP smear showing inflammatory smear. Hence a large no of women with abnormal

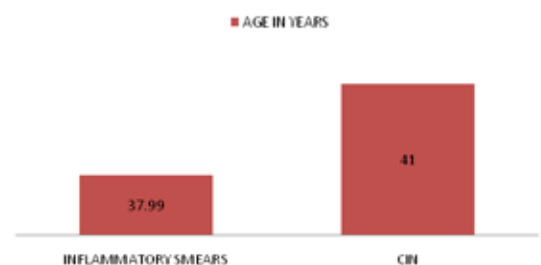
colposcopy and CIN (around 10%) would be missed if inflammatory PAP smears are not evaluated further.



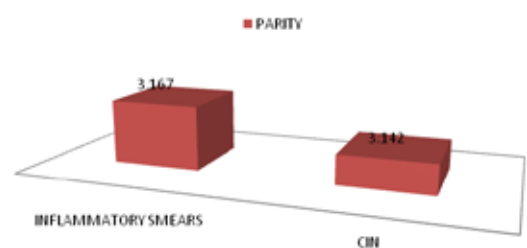
### AGE OF ONSET OF SEXUAL ACTIVITY IN YEARS



### AGE IN YEARS



### PARITY



## REFERENCES

- DiBonito L, Falconieri G, Tomasic G, Colautti I, Bonifacio D, Dudine S. Cervical cytopathology: An evaluation of its accuracy based on cytohistologic comparison. *Cancer* 1993;72:3002-6. | Wilkinson EJ. Papanicolaou smear and screening for cervical neoplasia. *Obstet Gynecol* 1990;35:817-25. | Gay JD, Donaldson LD, Goellner JR. False negative result in cervical cytological studies. *Acta Cytol* 1985;29:1043-6. | Seçkin NC, Turhan NO. Ozmen S, Ersan F, Ayar F, Ustün H. Routine colposcopic evaluation of patients with persistent inflammatory cellular changes on Pap smear. *Int J Gynaecol Obstet* 1997;59:25-9. | McLachlan N, Patwardhan JR, Ayer B, Pacey NF. Management of suboptimal cytological smears. *Acta Cytol* 1994;38:531-6. | Parashari A, Singh V, Gupta MM, Satyanarayana L, Chattopadhyaya D, Sodhani P, et al. Significance of inflammatory cervical smears. *APMIS* 1995;103:273-8. | Eckert LO, Koutsky LA, Kiviat NB, Krone MR, Stevens CE, Eschenbach DA. The inflammatory Papanicolaou smear: What does it mean? *Obstet Gynecol* 1995;86:360-6. | 8. Walker P, Dexeus S, De Palo G, Barrasso R, Campion M, Girardi F, et al. International terminology of colposcopy: An updated report from the International Federation for Cervical Pathology and Colposcopy. *Obstet Gynecol* 2003;101:175-7. | Mali BN, Joshi JU, Bhavé GG, Wagle UD. Cervical cytology in prostitutes of Bombay (India). *Genitourin Med* 1992;68:62-3. | 10. Wright TC, Sun XW, Luolos J. Comparison of management algorithms for the evaluation of women with low grade cytologic abnormalities *Obstet Gynecol* 1995;85:202-10 | Lonky NM, Navarre GL, Saunders S, Sadeghi M, Wolde-Tsadiq G. Low-grade Papanicolaou smears and the Bethesda system: A prospective cytohistopathologic analysis. *Obstet Gynecol* 1995;85:716-20. | Wright TC, Massad LM, Dunton CJ, Spitzer M, Wilkinson EJ, Solomon D, et al. 2006 consensus guidelines for the management of women with abnormal cervical cancer screening tests. *Am J Obstet Gynecol* 2007;197:346-55. | Dasari P, Rajathi S, Kumar SV. Colposcopic evaluation of persistent inflammatory Pap smear: A prospective analytical study. *Cytojournal* 2010;7:16 |