Visual inspection with Acetic Acid (VIA) as an Alternative to Colposcopy for detection of Cervical Carcinoma in low recourses setting

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Abstract

Objective: To evaluate visual inspection with acetic acid (VIA) as an alternative to colposcopy in detection of cervical carcinoma in low resource setting. Introduction: Cervical cancer is a major public health problem in developing countries, with India itself accounting for one-fifth of the global burden of the disease. In low resource settings, screening of CIN and carcinoma cervix by colposcopy can be replaced by cheaper and easily available visual methods like VIA, which has the high sensitivity to detect any grade of dysplasia. Method: VIA was performed in two hundred symptomatic women as a pre colposcopy attending the gynaecology department at National Institute of Medical Science (NIMS) Hospital Jaipur. Sensitivity of VIA was calculated. Result: Out of 200 patients VIA was positive in 48 (24%) cases. In VIA positive cases, Colposcopy was abnormal in 30 (62.5%). In VIA negative cases colposcopy was abnormal in only in 2 (1.3%) cases. 32 cases with abnormal colposcopic finding were subjected to biopsy. Conclusion: The high Sensitivity of 86.6% of VIA shows that most of the preinvasive and invasive lesion were picked up by VIA. Thus VIA could be valuable in detection of precancerous lesion of the cervix in low resource setting.

Key Words: Visual Inspection with Acetic Acid, Colposcopy, Cervical Cancer, Low Resource Setting

Introduction

Cervical cancer is an avoidable cause of death among women with 528000 new cases every year. Cervical cancer is the fourth most common cancer affecting women worldwide, after breast, colorectal and lung cancers; it is most notable in the lower-resource countries of sub-Saharan Africa. It is also the fourth most common cause of cancer death (266000 deaths in 2012) in women worldwide. Almost 70% of the global burden falls in areas with lower levels of development and more than one fifth of all new cases are diagnosed in India [1].

Cervical cancer is preceded by a long phase of cytological changes, known as cervical intraepithelial neoplasia (CIN) and takes a long period of 15-20 years before the invasive cancer develops [2].

Thus cervical cancer can be prevented if cellular changes are detected and treated in early stage. In western countries, cervical cancer incidence and death have been brought under control using organised cytology based screening program (pap smear)[3]. Though cytology (pap smear) is reliable, the laboratory infrastructure and

Manuscript received: 24th June 2014 Reviewed: 07th July 2014 Author Corrected: 14th July 2014 Accepted for Publication: 26th July 2014 logistic including technical expertise may not be available in low resource setting. The cost factor in testing for human papillomavirus using DNA testing, coupled with requirement of trained manpower makes this option non-viable for developing countries [4]. India with its highest share of global burden of cervical cancer has to implement a population based cervical cancer control programme to reduce the number of death [5]. Visual inspection using acetic acid (VIA) has emerged as a promising, cost effective, non-cytology based, "see and treat" alternative for economically underprivileged geographic regions. The present study was undertaken to find the value of VIA in picking up pre-invasive or invasive cancer in an abnormal cervix. All VIA positive cases were further evaluated with a colposcopy. A directed biopsy was taken in cases with abnormal colposcopy

Material and Methods

This prospective study was conducted in department of Obstetrics and Gynaecology National Institute of Medical Science (NIMS) Jaipur (Rajasthan). 200 symptomatic women over a period of one year attending Gynaecology clinic were studied. Approval of the institutional ethical committee and informed consent was

obtained from each women. Relevant obstetric and gynaecological history was obtained and recorded.

Inclusion criteria:

Age - >25 yrs

Marrital life >3yrs

Parous

Non Pregnant

Symptomatic cases

Exclusion criteria:

Virgin

Pregnant

In menstrual phase

H/o hysterectomy

Confirmed cases of carcinoma cervix

After General physical examination, local examination, cervix was exposed by Cusco's vaginal speculum. Any evidence of infection, ectopy, any visible growth etc were examined. Then 3-5 % acetic acid solution was applied to the cervix for 1 minute (with the help of swab stick)

Observation

Results are tabulated and discussed as follow.

Table 1: Demographic distribution of cases

Age Group (years)	No. of Women	Parity			
		1-2	3-4	5-6	7 or above
18-24	12	07	01	04	00
29-34	62	01	37	23	01
35-44	94	02	64	28	00
45-54	25	00	12	11	02
>54	07	00	03	01	03

Out of 200 symptomatic women, most women belong to age group of 35-44 yrs and parity between 3-4. The mean age of cases was 38.175 yrs with mean parity of 4.19. The commonest presenting complaint was vaginal discharge (62% of the patient). Other presenting complaints were intermenstrual bleeding (25 % of the patient), postcoital bleeding (8% of the patient) and postmenopausal bleeding (5% of the patient).

On visual inspection of cervix with 3-5% acetic acid, 48 cases (24 %) were VIA positive, Of these 30 cases (62.5%) had abnormal colposcopic finding and 18 cases (37.5 %) had normal colposcopic finding. 152 cases (76%) were VIA negative, Of these 2 cases (1.3%) had abnormal colposcopic finding and 150 cases (98.6 5) had normal colposcopic finding.

Table 2: Correlations of histopathology with VIA result

HPR	VIA	Total	
	Negative	Positive	Total
CIN 1	2	10	12
CIN 2		4	4
CIN 3		2	2
Invasive Cancer		1	1
Normal/Negative		13	13
Total	2	30	32

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and then the cervix was carefully inspected for any acetowhite lesions, particularly in the transformation zone. The normal squamous epithelium of the cervix is pink. On application of acetic acid, cervical intraepithelial neoplasia (CIN) lesion takes on white colour due to increased nuclear proteins and cytokeratins in the cervical epithelium [7].

Criteria for classifying via positive

- Location of lesion
- Colour intensity of acetowhite lesion
- Borders and demarcation of acetowhite lesion
- Size of lesion
- Extension of acetowhite lesion

After VIA patient were then examined by Colposcope with 12X-15X magnification. Biopsy was taken only from those who showed abnormal colposcopic appearances. Colposcopy and biopsy was taken as a reference standard for statistical calculation.

30 cases of VIA positive correlated with positive histopathological finding. Remaining 18 VIA positive cases have normal histopathology.

Table 3: Correlations of colposcopy and histopathology

	Colposcopy		
HPR	Negative	Positive	Total
CIN 1		12	12
CIN 2		4	4
CIN 3		2	2
Invasive Cancer		1	1
Chronic cervicitis		13	13
Total	0	32	32

As per table all the cases with abnormal colposcopy findings have positive histopathology results.

Table 4: Correlations of colposcopy and VIA with biopsy result

HPR	Sensitivity		PPV	
	VIA	Colposcopy	VIA	Colposcopy
CIN 1	10/12	12/12		
CIN 2	4/4	4/4		
CIN 3	2/2	2/2		
Invasive Cancer	1/1	1/1		
Overall	17/19	19/19	17/30	19/32

It is clear from table that for CIN 1 stage sensitivity of VIA is less. For advanced lesion sensitivity of VIA & colposcopy was 100 %.

Table 5: Sensitivity and positive predictive value of VIA and colposcopy

	Sensitivity		PPV	
HPR	VIA	Colposcopy	VIA	Colposcopy
CIN 1	83.33%	100.00%		
CIN 2	100.00%	100.00%		
CIN 3	100.00%	100.00%		
Invasive Cancer	100.00%	100.00%		
Overall	89.47%	100.00%	56.67%	59.375%

The most sensitive test for all grades of CIN and invasive cancer is colposcopy followed by VIA which is less sensitive in CIN1.

Pearson Chi-Square= 101.609 DF=1 'p' value <0.001

Kappa value=0.691 'p' value <0.001

As above table shows that there was 15% of positive cases and 75% of negative cases were showing agreement. On application of Chi-square test and kappa statistics there were significant association (Chi-square – 101.609 and p value < 0.001) and statistically significant (k - 0.691, p < 0.001).

Thus from above table it is clear that there is statistically significant association between colposcopy and VIA. Thus VIA is an alternative of Colposcopy.

Discussion

Cervical cancer is a potentially preventable cancer. It is preceded by premalignant lesions which may take 5-15 years to progress to invasive cancer. If detected and treated timely, pre-invasive disease has nearly 100 per cent cure rate with simple surgical procedure, while advanced cancers have less than 35 per cent survival rates [8].

However, in developing countries like India, universal screening has not been achieved. The main screening method (Pap smear) is available to a small percentage of population. Cytology based screening programmes are difficult to organize owing to limited infrastructure, trained personnel and funds [9].

It has been estimated that in India, even with a major effort to expand cytology services, it will not be possible to screen even one-fourth of the population once in a lifetime [10].

Moreover, screening programmes in India are mostly institution based and are restricted to urban centres [11]. Thus, in developing countries, there is a need for alternative strategies for early detection of premalignant cervical lesions.

Thus VIA is a cost effective method which can differentiate a normal cervix from a precancerous cervix with reasonable accuracy. Hence, if practiced by gynaecologists routinely, it would help to avoid many panic hysterectomies. Besides, in the absence of organized cytology screening program and limited availability of colposcopy facilities, VIA can be propagated on a wider scale.

In the present study, women from all the age group were included. The mean age was 38.17 years and majority were parous women. This observation agrees with the studies of singh kavita N et al [12].

N Yusuf et al studies correspond well with this study that CIN is more prone in sexually active women. Therefore, Screening for cervical carcinoma should start ideally at the onset of sexually activity and all the women should be screened at least once by the age of 30-35 years.

In our study sensitivity of VIA was 89.47% which shows that most of the pre-invasive and invasive lesions were picked up by VIA. Only two patients were missed (false

negative), one with post -menopausal bleeding where transformation zone receded inside and other case was of adenocarcinoma.

Shankarnarayanan and Mahe [13] have published results from a randomized intervention trial in India comparing VIA to cytology and to HPV DNA testing, and found that all three had similar detection rates of CIN 2 and 3 lesions, and the range of sensitivity for VIA was 67-79% and specificity 49-86%.

The higher sensitivity of our study can be explained on the basis that the study was done on a group of women in susceptible age group (mean 38.48 ± 8.9 years). Secondly, the VIA was performed using colposcope halogen bulb (24V 250 Watt) by a gynaecologist and not by a nurse or a paramedical person.

In JHPIEGO cervical cancer project at Zimbabwe, VIA was done by trained nurses and paramedical person on 10,934 women and was found to have sensitivity of 76.4% and specificity of 64% in picking up pre-invasive lesions [14].

The study also inferred that higher test qualities of VIA are likely to be observed under better service delivery conditions like good lighting, examination tables and specula, and more standardized VIA training. They observed that in more than 75% of cases when a lesion was found on colposcopy or biopsy, it was also visible on VIA [15].

Ottaviano and Torre [16], in the first study of its kind, have published results of simultaneous VIA and colposcopy on 2,400 unselected patients with normal or abnormal cervical cytology.

In our study, VIA and biopsy correlation is less for CIN1 which is 83.3%., but the sensitivity increased up to 100% in picking up CIN 2, CIN 3 which is indeed a true cancer precursor and early invasive cancer . Colposcopic magnification is a complimentary method to VIA and not essential to identify a cervix with higher grade lesions.

VIA can also guide a practicing gynaecologist regarding the site to be biopsied. Although colposcope accurately identifies the most abnormal area for biopsy,VIA will demarcate the site precisely and give better results than a blind biopsy of ectocervix, in absence of facilities for colposcopy [17].

In our study it is clear that there is statistically significant association between colposcopy and VIA. Thus VIA is an alternative of Colposcopy.

Conclusion

VIA is attractive test in low resource settings like India. It is simple, inexpensive low technology test that requires minimal infrastructure.

The majority of women belonging to low socioeconomic status, remain devoid of any screening test, thus time has come to integrate VIA based screening program at the primary care level of health services and to down stage cancer cervix in our country.

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