p53 Immunohistochemical Expression of Egyptian Cervical Carcinoma

Howayda ABD EL ALL 1*, Annie RYE,2 Pierre DUVILLARD3

1 Department of Pathology, Faculty of Medicine, Suez Canal University, Egypt; 2 Department of Statistics and 3 Department of Pathology, Institute Gustave Roussy, Villejuif, France

Introduction

Cervical carcinoma is the second leading cause of death in women worldwide. In Egypt, cervical carcinoma represents 59.58% of female genital tract malignancies according to the statistics of the National Cancer Institute of Egypt. Many factors are implicated in the process of cervical carcinogenesis. In the western population, there is evidence that human papilloma virus infection (HPV) especially 16 and 18 subtypes are involved in the development of cervical carcinoma. The suppressor gene p53 plays a pivotal role in protection against the development of cancer. Its wild type monitors the integrity of the genome. The mechanism of p53 activation in response to DNA damage depends on DNA-protein kinase. Studies on the value of p53 in cervical carcinoma are controversial. Some studies showed that p53 protein accumulation may be an early event in carcinogenesis, while for others, p53 overexpression or mutation did not seem to play a significant role in cervical carcinogenesis. The mechanism of inactivation or loss of the wild type in the development of cervical carcinoma, has been attributed to either mutation within the genome or the presence of virally encoded p53 binding protein. Mutant p53 could form an oligomeric complex with wild type preventing the latter from functioning, or alternatively, mutant p53 could gain a new oncogenic function that overcomes the negative regulation by small quantities of infection in Egypt are lacking and information regarding the possible role of schistosomiasis in the process of cervical carcinogenesis are debatable.

Data concerning the expression of p53 in cervical carcinoma, one of the leading cause of death in developing countries, are still confusing. This study was designed to identify p53 in Egyptian cervical carcinoma in an attempt to evaluate its prognostic significance. Eleven chronic cervicitis and 38 invasive carcinoma (31 squamous cell carcinoma (sqcc) and 7 adenocarcinoma, ranging from stage IB to IVB), were stained with the monoclonal antibody anti p53, DO7, using the microwave for antigen retrieval. No immunoreactivity was detected in chronic cervicitis, while nuclear p53 reactivity was detected in all carcinoma and in squamous intraepithelial lesions (SIL) overlying 8 sqcc. P53 immunohistochemical (IHC) expression was more pronounced in early clinical stages (p=0.007) and in adenocarcinoma compared to sqcc (p=0.015). A positive correlation was present between p53 and heat shock protein 70 (hsp70) expressions (p=0.005). No correlation could be found between p53 expression and tumor infiltrating lymphocytes, the presence or absence of either schistosomiasis or HPV infections. It can be concluded, that in the Egyptian population, p53 immunoreactivity appears to be an early event in cervical neoplasm, and seems to play an important role together with other cell regulatory proteins in the process of carcinogenesis, which could be different between sqcc and adenocarcinoma. (Pathology Oncology Research Vol 5, No 4, 280–284, 1999)

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Correspondence: Howayda ABD EL ALL, M.D., Department of Pathology, Faculty of Medicine, Suez Canal University, Ismailia, Egypt; Tel: + (202) 274-32-31; Fax: + (202) 274-36-70; e-mail: howayda@intouch.com
Abbreviations: sqcc: squamous cell carcinoma; IHC: immunohistochemistry; SIL: squamous intraepithelial lesion; HPV: human papilloma virus infection
wild type. Based upon recent advances, several therapeutic strategies had been envisioned concerning the structure and function of the p53 protein, its interaction with cellular and viral proteins and its roles in repairing DNA, regulating cell division and promoting apoptosis and other approaches had focused on restoring p53 function in human tumors.

We performed this work in order to study the significance of IHC expression of p53 in invasive cervical carcinoma in the Egyptian population and to correlate this expression with the FIGO clinical staging, the histopathologic type and grade, the presence or absence of HPV and schistosomiasis.

**Materials and Methods**

**Patients**

Samples were available from 38 untreated patients with invasive cervical carcinoma, 23 by punch biopsy and 15 by radical hysterectomy. All patients were included in two previous studies. As the treatment modality for stages IB and IIA is different from stages IIB, III and IV, and for statistical purposes, we designated the former as early clinical stages and the later as late stages. The control tissues were cervixes diagnosed as non neoplastic cervical tissues. All carcinomas were stained with the p53 DO7, supplied manual with the Kreatech In Situ Hybridization Detection Kit. Negative controls were non-specific DNA probes supplied with the kits. Tissue specimens, condyloma and SIL known for their positivity with the probes were used as positive controls.

**Immunohistochemistry**

For immunostaining, 5 µm thick sections were dewaxed in xylene over night; dehydrated in decreasing concentrations of alcohol ending in phosphate buffer saline (PBS). The step of deparaffinization was critical, since slides immersion in xylene for 15 to 30 minutes were negative for staining and few cells were stained by immersion for 60 minutes. The microwave oven heating method was used for antigen retrieval. Sections were irradiated at 750w for 15 minutes in citrate buffer (pH 6.0) (3 x 5 minutes) cycles with 1 minute interval between cycles to check out the fluid level. Slides were left to cool for 15 minutes, rinsed in distilled water twice and finally in PBS. The staining steps was performed according to the manual supplied with the labeled streptavidin biotin kit (DAKO) and the mouse monoclonal antibody DO-7 (DAKO) that react with the wild and mutant type p53 at a dilution of 1:50. The duration of incubation performed at room temperature was 90 minutes. Positive staining was visualized with 3,3 diaminobenzidine tetrachloride (DAB). For counterstaining methyl green was used. Only nuclear staining was considered positive. Both the number or percentage of positive cells and the intensity of staining were assessed. The staining intensity was graded as follows: absent (-), mild (+), moderate (++) , severe (+++). The percentage of positive cells was assessed in a semi-quantitative manner in 10 high power fields as follows: 1. 1-5% positive cells, 2. 6-25%, 3. 26-50%, 4. 51-75% and 5. >75%. The p53 score was the sum of the intensity and percentage of positive. Negative controls were parallel sections treated as above with omission of the primary antibodies.

**In situ hybridization**

This was previously done and detailed. Briefly, the OmniProbe and the DAKO in situ detection system for biotinylated probes were used for initial screening of all the tumors (DAKO). OmniProbe, is a wide spectrum HPV biotinylated probe for which the targets are the genomic DNAs of HPV types 6, 11, 16, 18, 30, 33, 35, 45, 51 and 52. Positive staining was present in 5 cases for which further subtyping with the probes directed against HPV 6/11, 16 and 18 (Kreatech) were performed according to the supplied manual with the Kreatech In Situ Hybridization Detection Kit. Negative controls were non-specific DNA probes supplied with the kits. Tissue specimens, condyloma and SIL known for their positivity with the probes were used as positive controls.

**Statistical analysis**

The correlation between the p53 and the different parameters was performed using the Chi square test. The 95% confidence interval for the rho values were calculated.

**Results**

There was no detectable p53 protein in non neoplastic tissues. All carcinomas were stained with the p53 DO7, with 7/38 (18%) having the maximum score 8 (Figure 1A). The staining was only nuclear either as fine or as coarse granular dots. The expression was greater in the peripheral cells of tumor islands in sqcc especially in the large cell keratinized subtype. There was marked heterogeneity between and within tumors. There being within the same section, islands of cells that were virtually negative, others having 1 to 5% of reactive cells and others with more than 75% positive cells. This heterogeneity was marked in adenocarcinoma (Figure 1B). In low grade SIL, the nuclear staining was confined to the basal layer until reaching the superficial one in high grade SIL. Tumor cells beneath these SIL had the same staining intensity but their number were increased. P53 immunoreactivity was detected in HPV positive and negative cases without significant differences between both groups. Similarly, p53 was detected in tumors with and without bilharziasis (Table 1). Metastatic cancerous cells infiltrating lymph nodes
showed the same staining intensity with the anti p53 as their primary tumors. P53 was more expressed in early clinical stages \( (p=0.0007) \) and in adenocarcinoma compared to sqcc \( (p=0.015) \) (Table 2 and Figure 2). A positive correlation was noted between p53 and hsp70 previously studied \( (p=0.005) \). No correlation was present with the other variable.

**Discussion**

The results of IHC of p53 in cervical carcinoma have been somewhat confusing. p53 immunoreactivity was detected in normal and inflammatory lesions of the cervix, explained as a result of overexpression or stabilization of the wild type as noted by Bosari et al., or as an effect of the microwave oven heating in the absence of mutations. In the present study search for p53 using the microwave oven heating, showed lack of expression of the protein in chronic cervicitis, an observation in agreement with other studies. However in the latter study, p53 was not expressed in condyloma, dysplastic tissues and carcinoma in situ while in the present work p53 immunoreactivity started in SIL. The expression began to be seen in few cells in the basal layer of low grade SIL and increased to parallel the extension of neoplastic cells in high grade SIL and carcinoma in situ and was more extensive in invasative lesions. The same observation was documented in previous studies. This may suggest that abnormal p53 expression participate alone or with other factors in the transformation of non neoplastic epithelium to SIL and latter on in the development of invasive cervical lesions. A similar finding in colorectal tumorigenesis was detected, where p53 alterations take place at the transition from adenoma to carcinoma.

Looking for the association of p53 with staging, studies are controversial. A significant pronounced expression of p53 in early clinical stages, has been reported in the work of Lakshmi et al. Most of the studies showed a lack of asso-

**Table 1. Expression of p53 in bilharziasis and HPV associated cervical carcinoma.**

<table>
<thead>
<tr>
<th>%p53</th>
<th>Bilharziasis</th>
<th>HPV</th>
</tr>
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<tbody>
<tr>
<td>Ova*</td>
<td>TTT**</td>
<td>16</td>
</tr>
<tr>
<td>1-5</td>
<td>3</td>
<td>5</td>
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<td>6-25</td>
<td>1</td>
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<td>26-50</td>
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<tr>
<td>51-75</td>
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<td>1</td>
</tr>
<tr>
<td>&gt;75</td>
<td>2</td>
<td>1</td>
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* ova in the neoplastic tissue
** treatment for urinary bilharziasis
Oka et al. noted that p53 CM1 expression in 49/119 stage III squamous cell carcinoma cases was not a predictive factor for the prognosis in cervical carcinoma. Helland et al. in 50/92 stage I-IV cases failed to find significant association between NCL-CM1 p53 protein expression and the clinical stage. Kainz et al. using the monoclonal antibody BP53-12 on 109 squamous cell carcinoma stage IB-IIIA failed to show any correlation. Using the same antibody in this study p53 DO7, Hunt et al. did not find relation between p53 and staging.

What is the significance of the early expression found in our study on our patients? We have noted in 2 cases of adenocarcinoma stage IIA, that p53 overexpression was associated with nodal metastasis and recurrence. Unfortunately, more than half of our patients were lost for follow up and we cannot find the significance of such expression on the clinical outcome of the patients. A long term study is needed to clarify this point.

We have previously shown an association between schistosomal infection and cervical carcinoma (p=0.005) and advanced clinical staging (p=0.03). The mechanism underlying the carcinogenic effect of bilharziasis has been attributed to several mechanisms, one of them was a possible involvement of loci on chromosome 11 in controlling the level of chromosomal breakage caused by oxidative damage of the inflammatory cells. We expected to find a relationship between schistosomiasis and p53 as we have found association between p53 and early clinical staging, but this was not the case.

Similarly, we failed to find association between p53 and HPV. Overexpression of p53 in tissues has generally been assumed to reflect p53 mutations in HPV negative cervical carcinoma, since in HPV positive cases, the wild type p53 complex with E6 of HPV 16 or 18, is degraded and cannot be detected by IHC. Did the expression of p53 in 33/38 HPV negative cases reflected mutation? We believe so, as we found an increased expression of c-myc and hsp70 in the same cases, and we found a direct correlation between IHC nuclear expression of hsp70 and p53 (p=0.005). In addition, examination of serial sections stained for hsp70 and p53 revealed co-localization of both in the nucleus of the same cell. Early studies showed interactions between mutant p53 protein and hsp70. Recently, mutations in the p53 protein has been shown to alter the tertiary structure of the central DNA binding domain, thus exposing high affinity hsp70 binding sites that are cryptic in the wild type molecule, therefore allowing the interaction between both mutant and hsp70. This finding in cervical carcinoma contradicted studies on breast carcinoma where only part of the p53 pool was found to be bound to hsp70, and that the nuclear accumulation of p53 was not associated with cytoplasmic or nuclear hsp70.

Regarding the over-expression of p53 in the HPV positive cervical carcinoma cases, we think that this might be due to non mutational stabilization of the p53 protein. Wynford-Thomas 38,39 suggested that the IHC detection of p53 may requires not only mutation of the gene but some additional secondary unknown mechanism of non mutational stability conferred by the neoplastic cells and that not all p53 detected by IHC should be considered mutated. Does this stabilization was conferred by the co-association of p53 with other molecules such as hsp70 and/or c-myc? A point needing further evaluation on molecular basis.

The finding of a pronounced p53 expression in adenocarcinoma compared to squamous cell carcinoma in the literature was less fortunate than its squamous counterpart. McCluggage and colleagues found the expression in 23/33 cases of adenocarcinoma, 2/10 adenocarcinoma in situ, and concerning the non neoplastic tissue, in 2/10 tubo-endometrial metaplasia, 1/10 microglandular hyperplasia, and 1/17 normal endocervical tissues. In another study, on 25 cervical adenocarcinoma and 7 adenosquamous carcinomas, Uchiyama et al. showed that HPV-negative or p53-positive adenocarcinoma may be a biologically distinct subset with a poorer prognosis.
Other studies failed to demonstrate staining in 13 adenocarcinoma examined.24

In conclusion, we have documented p53 immunoreactivity in all cervical carcinoma either HPV positive or negative and in cases with or without schistosomiasis. This expression was significantly associated with early clinical stages and adenocarcinoma and appears to be related to p53 mutation.

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