Alpha-fetoprotein as a differential marker between bladder cancer and Schistosomiasis among Egyptian patients


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Abstract

Introduction: Schistosomiasis is considered as a widespread problem that affects Egyptians at different ages (WHO, 1993). It is well known that the high incidence of bladder carcinoma in schistosomal patients also represents a great risk to Egyptian society.

Objective: This study represents a survey on the changes which took place in Alpha-Fetoprotein (AFP) levels as a differential marker among Egyptian patients suffering from urinary bilharzial infection and bladder cancer.

Methodology: A partial study was carried out on four post-operative patients and followed up for sixty three days after chemotherapeutic treatment using immunoenzymatic mediated assay.

Results: It was found that bladder cancer leads to obvious significant increase in AFP levels, while schistosomal infection causes a non-significant decrease as compared to healthy control. Although the combination between schistosomal infection and cancer leads to a significant decrease in AFP level compared to non bilharzial cancer cases, there was no significant correlation with healthy cases.

Conclusion: It should be pointed out that the somewhat constant level of AFP in the sera of all patients was due to chemotherapeutic treatment. Also, it can be noticed that there is a significant positive correlation with progress of cancer grades.

Key words: Alpha-Fetoprotein, schistosomal infection, urinary bilharzial infection and bladder cancer.

Abbrevations: AFP: Alpha-Fetoprotein

Introduction

Where schistosomiasis remains endemic, a great deal has been published from Egypt about the role played by schistosoma parasite in development of malignant lesions in bladder (Ammal et al., 1992).

Carcinoma of the urinary bladder is the most common malignancy in the Middle-East and parts of Africa where schistosomiasis is a widespread problem. Much evidence supports the association between schistosomiasis and bladder cancer: this includes the geographical correlation between the two conditions, the distinctive pattern of gender and age at diagnosis, the clinical and pathological identity of schistosoma- associated bladder cancer, and extensive evidence in experimentally infected animals (Mostafa et al., 1999). So, the size of bilharzial problem and its complication needs more careful studies to demonstrate new marker to clarify the effect of bilharziasis in the development of carcinogenesis (Gu et al., 1987). So, alpha-fetoprotein (AFP) is the marker of choice to study in this research. AFP is a serum glycoprotein, initially identified in human sera in 1956 as one of the two major proteins in fetal blood (Bergstrand and Czar, 1956). It is marked for hepato-cellular and germ cell carcinoma. Its synthesis by embryonal liver cells is virtually inhibited during adult life. Thus, it is not demonstrable in normal tissues but has been found in traces in adult human sera, this ratio began to increase with some body disorders and can be determined by some more sensitive
radioimmunoassay (Albert, 1979). In testicular cancer, lactate dehydrogenase (LDH), AFP, and human chorionic gonadotropin (hCG) are essential markers not only to determine the tumor stage but also for the prognosis of the patients (Shimazui et al., 2004). Asymptomatic individuals were screened on a voluntary basis using a panel of tumor markers, including AFP, CA 125, CA 15-3, CA 19-9, carcinoembryonic antigen (CEA), prostate specific antigen (PSA), chromogranin A (CgA), and squamous cell specific antigen (SCC) for cancer at Chang Gung Memorial Hospital in Taiwan (Tsao et al., 2006).

Complete ablation rates after a single session of radiofrequency ablation (RFA) of hepatocellular carcinoma (HCC) vary from 48% to 97%. Limited data are available regarding risk factors and prognostic significance of incomplete ablation. By univariate analysis, it was found that absence of previous transarterial chemoembolization (TACE), preoperative serum alfa-fetoprotein was less or equal 100 µg/mL, and complete response after further treatment of incomplete ablation were associated with better overall survival in patients with incomplete ablation (Lam et al., 2008). Antigen A2/3 is alternative to alfa-fetoprotein (AFP) it was induced by heavy metal salts (Pb2+ and Cd2+) in the liver of adult rats and AFP+/A2/3(-) clones of hepatomas; the attenuation of AFP synthesis occurred simultaneously (Poltoranina et al., 2007).

Several evidences support the association between the two diseases. So, our works represents a survey on the changes which took place in Alpha-Fetoprotein (AFP) levels as differential marker among Egyptian patients suffering from urinary bilharzial infection and bladder cancer.

Subjects and Methods

Immunoenzymatic mediated assay (IEMA):

The immunoenzymatic mediated assay is based on the principle of a “sandwich” formed of the analyte to be detected between two specific monoclonal antibodies directed against two different epitopes on the analyte molecule. Measurement of the activity is performed by adding a colorless chromogen solution. The action on chromogen produces a color which can be measured by ELISA reader (SORIN-BIO-MEDICA).

Subjects:

This work was conducted on 40 subjects including 36 patients submitted to the Surgical and Internal Medicine Departments in National Cancer Institute, Cairo University. The investigated patients were divided into five main groups according to the histological diagnostic reports supplied by the pathology department in the institute. The bilharzial infection was based on the stool and urine examinations documented in the disease history reports associated with patients submitted to Theodor Bilharz Institute.

The patients groups were:
1-Patients having bladder cancerous at three grades (12 patients at grades I, II and III)
2-Patients having bilharzial bladder cancer (12 patients at grades I, II and I
3-Patients were suffering from urinary schistosomiasis (4 patients).
4-Patients having bladder cancer and submitted to chemotherapy treatment (4 patients).
5-Patients suffering from bladder cancer post-operative (4 patients).
Four healthy persons were included in this study as a control group.

Specimen collection:

Venous blood samples were collected from all the previous groups and sera were separated and kept at -70°C until the assay.

Statistical analysis:

Statistical analysis was done using ANOVA test and regression line analysis.

Results

One of the most clinically useful tumor associated antigens of this class is AFP which has been widely studied in patients with gastrointestinal cancer, therefore we try to demonstrate the changes which might took place in AFP levels as a tumor marker among Egyptian patients
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suffering from both bilharzial infestation and cancer bladder.

Table (2) show the effect of schistosomal infection which leads to slight decrease in AFP levels, with percent change equal 12.5% referred to control. The difference was statistically insignificant (P < 0.05) by ANOVA.

Data summarized in tables (2-4) demonstrated the AFP levels in bilharzial and non-bilharzial bladder cancer compared to their levels in healthy cases. It is clear that cancerous infiltration resulted with highly significant increase which was higher in grade III than grade II and I in the two types of cancer. Although the combination between schistosomal infection and cancer lead to significant decrease in AFP level compared to non-bilharzial cancer cases, there was no significant difference with healthy cases. On the other hand, the non-bilharzial cancer leads to significant elevation of AFP as compared to control cases. It is worthy mentioned that this decrease was significantly marked with the conversion to malignancy (Figs. 1, 2).

The specific treatment protocol for refractory carcinoma in situ of the bladder remains elusive. It is ultimately the combined decision of the clinician and patient to determine which course of management is most beneficial (Kim and Steinberg, 2001). In the present study, we used AFP to differentiate between post-operative and after therapy; whereas the addition of chemotherapy to surgery improved both disease-free and overall survival rates (El-Mawla et al., 2001). The statistical analysis showed that there has been no significant difference after-operative or chemotherapy whereas P<0.05 by using ANOVA test (Table 5). On the other hand, there was a significant difference post-operative and post-chemotherapy as compared to control.

**Table (1)**
α-Fetoprotein concentrations in serum of normal subjects and patients with cancer bladder, bilharzial cancer bladder and Bilharzzziiasis (ng/ml).

<table>
<thead>
<tr>
<th>Groups</th>
<th>α-fetoprotein ng/ml</th>
<th>Mean ± S.D</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bladder cancer Grade I</td>
<td>5.0-4.8-4.9-4.9</td>
<td>4.9 ± 0.07</td>
</tr>
<tr>
<td>Bladder cancer Grade II</td>
<td>5.4-4.9-5.1-5.0</td>
<td>5.1 ± 0.18</td>
</tr>
<tr>
<td>Bladder cancer Grade III</td>
<td>5.6-6.3-6.0-5.1</td>
<td>5.75 ± 0.4</td>
</tr>
<tr>
<td>Bilharzial bladder cancer</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Grade I</td>
<td>4.2-3.8-3.8-4.5</td>
<td>4 ± 0.2</td>
</tr>
<tr>
<td>Grade II</td>
<td>4.7-4.0-3.9-4.5</td>
<td>4.28 ± 0.3</td>
</tr>
<tr>
<td>Grade III</td>
<td>4.9-4.6-3.9-4.5</td>
<td>4.48 ± 0.36</td>
</tr>
<tr>
<td>Bilharzial cases</td>
<td>3.5-3.6-3.2-3.8</td>
<td>3.5 ± 0.2</td>
</tr>
<tr>
<td>Post – Opriteive</td>
<td>1.6-1.4-1.8-1.6</td>
<td>1.6 ± 0.14</td>
</tr>
<tr>
<td>Post – Chemotherapy</td>
<td>1.5-1.5-2.0-1.8</td>
<td>1.7 ± 0.2</td>
</tr>
<tr>
<td>Normal cases</td>
<td>3.8-4.3-4.1-3.6</td>
<td>3.95 ± 0.26</td>
</tr>
</tbody>
</table>
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Table 2: α-Fetoprotein concentrations (ng/mL) in serum of urinary Schistosomiasis, bladder cancer and bilharzial cancer patient as compared to control cases.

| Groups                  | Mean ± 0.26 | % change | P <  
|-------------------------|-------------|----------|------
| Control                 | 4.0 ± 0.26  | —        | —    |
| Urinary Schistosomiasis | 3.5 ± 0.2   | 12.5     | 0.05 |
| Bladder cancer I        | 4.9 ± 0.01  | 22.5     | 0.001|
| II                      | 5.1 ± 0.18  | 27.8     | 0.001|
| III                     | 5.8 ± 0.8   | 45.0     | 0.001|
| Bilharzial              | 4.1 ± 0.2   | 2.5      | 0.001|
| Bladder II              | 4.3 ± 0.3   | 7.5      | 0.001|
| Cancer III              | 4.5 ± 0.36  | 12.5     | 0.001|

Table 3 α-Fetoprotein concentrations (ng/mL) in serum of bilharzial and non bilharzial bladder cancer patients as compared with Schistosomiasis cases.

| Groups                  | Mean ± 0.26 | % change | P <  
|-------------------------|-------------|----------|------
| Urinary Schistosomiasis | 3.5 ± 0.2   | —        | —    |
| Bladder cancer I        | 4.9 ± 0.07  | 40       | 0.001|
| II                      | 5.1 ± 0.18  | 45.7     | 0.001|
| III                     | 5.8 ± 0.4   | 65.7     | 0.001|
| Bilharzial              | 4.1 ± 0.2   | 17.14    | 0.01 |
| Bladder II              | 4.3 ± 0.3   | 22.9     | 0.01 |
| Cancer III              | 4.5 ± 0.36  | 28.6     | 0.001|

Table 4 The effect of bilharzial infection on serum (AFP) concentrations (ng/mL) on Bladder cancer patients.

| Groups                  | Mean ± 0.26 | % change | P <  
|-------------------------|-------------|----------|------
| Bladder cancer          | 4.25 ± 0.19 | —        | —    |
| Bilh-bladder cancer     | 5.23 ± 0.03 | 23.1     | 0.001|
| Bilharzial I            | 4.1 ± 0.2   | 17.14    | 0.01 |
| Bladder II              | 4.3 ± 0.3   | 22.9     | 0.01 |
| Cancer III              | 4.5 ± 0.36  | 28.6     | 0.001|

Table 5 – The effect of Chemotherapeutical treatment on serum (AFP) concentrations (ng/mL) in Patient post operative.

| Groups                  | Mean ± 0.26 | % change | P <  
|-------------------------|-------------|----------|------
| Post-operative          | 1.6 ± 0.14  | 60       | 0.001|
| Post-Chemotherapy       | 1.7 ± 0.2   | 57.5     | 0.001|
| Control                 | 4 ± 0.26    | —        | —    |
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Figure 1: Levels of alpha fetoprotein according to health status:

![Bar chart showing levels of alpha fetoprotein according to health status](image1)

Figure 2: The difference of the alpha fetoprotein post-operative and after chemotherapy.

![3D bar chart showing the difference of alpha fetoprotein](image2)

Figure 2: The difference of the alpha fetoprotein post-operative and after chemotherapy.
Discussion

The search for specific tumors markers is of considerable importance as new and more effective therapies for many neoplasms are becoming available. The ideal markers should be specific for the organ involved by the tumor and elevated in the majority of patients with neoplastic disease not in benign one (Berlin, 1981). A variety of substances circulating in serum are potentially useful as tumor markers such as certain glycoproteins. Some biochemical markers give a more precise and earlier diagnosis than the usual ones (Goffinet et al., 2001).

Our results were as similar to those obtained by Alsbti (1978) where as two hundred and eighty-four patients of various ages from an endemic area, diagnosed as having bilharziasis by the presence of living Schistosoma haematobium's ova in the mid-stream urine sample, have been investigated for the presence of AFP in the serum using radioimmunoassay. One hundred and forty-one (49.6%), were positive. Wespic and Kirkatrick (1979) suggested that AFP elevated in 5-10% of patients with large liver metastases virtually exclusively from gastric and pancreatic patients. Also when we evaluated the effect of progressive grading of carcinoma on AFP levels, our data showed that AFP level increase progressively as cancer grades progresses reached the maximum values or grade II. So, there is a gradual significant positive correlation with progress of cancer grades. These results were in line with that detected by Joan et al. (1988) who found gradually increase in AFP in normal pregnant women and in complication with cancer in all grades. The same results were obtained in comparing the grades among the non bilharzial cancer. This can be attributed to the fact that in case of early grade the cells have more benign course rather than late grades, and this is also supported by Trojan et al. (1983) who studied the localization of AFP in different tissues including normal tissues in rat, baboon and mouse teratocarcinoma and they suggested that the incorporation of AFP was associated with the degree of differentiation. AFP and prostate-specific antigen (PSA) in serum are widely used as tumor markers in the evaluation of prognosis and management of patients with heptato-cellular carcinoma and prostate cancer, respectively. Reverse transcriptase polymerase chain reaction (RT-PCR) for AFP and PSA were used to identify circulating cancer cells in the blood of cancer patients. Broad expression of AFP was in several tissues and a large amount of AFP mRNA was found in fetal liver. PSA was expressed in prostate, salivary gland, pancreas and uterus. By RT-PCR, AFP and PSA mRNA were detected in several tumors, including salivary pleomorphic adenoma, hilar bile duct carcinoma, pancreatic carcinoma, transitional cell carcinoma of urinary bladder and thyroid papillary carcinoma. Furthermore, AFP and PSA mRNAs were frequently detected by RT-PCR, even in peripheral blood cells from healthy volunteers (Ishikawa et al., 1998). The early diagnosis of bladder cancer allows for effective local treatment and optimizes the success of surgical therapy. Basic fetoprotein (BFP) has no correlation with tumor grade, while cytology had a strong association. Linear regression analysis showed the significant correlation between BFP level and tumor size. The detection rate of bladder cancer was higher by the combination of BFP and cytology than by using one alone (Ichikawa et al., 2000).

Conclusion

In conclusion, we can use AFP level as a differential marker among Egyptian patients suffering from urinary bilharzial infection and bladder cancer infestation using immunoenzymatic mediated assay.

References

تقييم مستويات البروتين الجنيني-ألفا كعامل مميز للتفرقة بين سرطان المثانة والمثانة البلهارسي في المرضى المصريين

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"قسم الكيمياء العلاجية - مركز التميز العلمي للتعليم المقدم - المركز القومي للبحوث.

قسم تكنولوجيا المختبرات الطبية - كلية العلوم الطبية التطبيقية - جامعة مصر للعلوم والتكنولوجيا.

قسم طب الأورام - المعهد القومي للأورام.

يعد البروتين الجنيني-ألفا من دلالات الأورام ذات الطبيعة البروتينية والتي تتكون أساساً في الخلايا الجنينية وأثناء نمو الخلية السرطانية لذلك فإن قياس معدلات تباين مستوياته في المرضى المصابين بسرطان المثانة والمثانة البلهارسي ذو أهمية للتفرقة بينهم. لذلك يهدف البحث إلى دراسة مضاعفات الإصابة ببطفل البلهارسية وتشمل تلك الأورام التي يمكن أن تؤدي إلى التهابات مزمنة للثدي مما ينتج عنه إفراز لبعض البروتينات والتي تظهر في الحالات السرطانية (سرطان الثدي، سرطان المثانة موضع الدراسة) مثل البروتين الجنيني-ألفا، إضافة إلى مقارنة نسبته في مرضى سرطان الثدي فقط لمعرفة تأثير عدوى البلهارسية وتطورها إلى مضاعفات سرطان الثدي على إنتاج هذا البروتين الجنيني. استنتجت النتائج أن الأورام السرطانية في الثدي تظهر تباين درجة قياس أنه يمكن أن يؤدي إلى ارتفاع ملحوظ في مستويات البروتين الجنيني-ألفا في حين كانت للإصابة ببكتريا البلهارسية تثبيط غير معنوي لمستواه مقارنة بالمجموعة الضابطة الأصحاء. كما أوضحت الدراسة أن تطور عدوى البلهارسية تثني على أن مستويات البروتين الجنيني-ألفا مقارنة بحالات سرطان الثدي غير البلهارسي. كما أثبتت الدراسة أيضاً أن العلاج الكيميائي المستخدم يؤدي إلى ثبات مستويات البروتين الجنيني-ألفا إلى حد ما. كما يلاحظ أن هناك علاقة إيجابية تدريجية بين مستويات البروتين الجنيني-ألفا ودرجات تقدم المرض.