Mast cells density in fibrotic capsule of enchondroma and well-differentiated chondrosarcoma: a method for histopathologic differentiation

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Abstract

Background: An enchondroma is a benign and a well-differentiated chondrosarcoma is an invasive chondroid tumor with high recurrence potential. In spite of biologic differences, these two tumors have very similar histopathologic appearance. It has been shown that the biologic nature of the connective tissue around benign and malignant tumors varies in the number of mast cells. The aim of this study was to study the histopathologic distinction of enchondroma and well-differentiated chondrosarcoma using the density of the mast cells in fibrotic capsule.

Methods: Twelve enchondroma and 15 well-differentiated chondrosarcoma were collected from Pathology department of Cancer Institute and Central Pathology department of Imam Khomeini Hospital in Tehran. 3 micron paraffin embedded tissue sections were stained by toluidine blue for mast cells counting. Mast cells were counted in fibrous capsule of all cases. Mast cells counts were accomplished in 10 high power fields .The average number of mast cells in 10HPF was determined as an index for each lesion. Mann-Whitney U test was used for statistical analysis.

Results: Mean index in enchondroma and well-differentiated chondrosarcoma groups were 0.1±0.12 and 0.31±0.33 respectively, showing a significant difference between number of mast cells in the fibrotic capsule in these two lesions (p=0.028). Comparison of the corresponding points in ROC curve, showed a cut-off point = 0.15, with positive predictive value of 61%, negative predictive value 71%, specificity of 33.3% and sensitivity of 66.7%, (p=0.025).

Conclusion: Average density of the mast cells in the surrounding fibrotic capsules of enchondroma and well-differentiated chondrosarcoma along with other criterions, could be a beneficial factor for histologically differentiation between these two lesions.

Keywords: Enchondroma, Well-differentiated Chondrosarcoma, Mast cell, Toluidine blue staining.

Introduction

An enchondroma is a benign chondroid tumor formed by mature hyaline cartilage and often involves medullary bone. Most of tumors are small with less than 3 cm in greatest dimension and with limited growth potential. Asymptomatic and small lesions do not need any treatment except periodic follow up (1).

A chondrosarcoma is a slow growing malignant tumor. It is locally invasive and has

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high recurrence potential. This tumor can extend to adjacent tissues and may result in limiting the range of joints motion. Surgery is the primary treatment for chondro-sarcoma with radical surgical excision be-ing the most effective mode of therapy (1).

Despite biologic differences, these two tumors have the same histologic appearances. The exact distinction between enchondroma and well-differentiated chondrosarcoma is a major problem in bone pathology.

In histologic evaluation, both lesions con-tain a mass of mature hyaline cartilage with a lobular pattern and small chondrocytes in lacunar spaces. Small to large foci of calci-fication may be seen (1-3). Because of this similarity, distinction between these two tumors on the basis of pathologic features is not feasible, and differentiation is achieved by clinical, pathologic, radiographic fea-tures, and complementary immune histo-chemical staining (3-7).

These diagnostic methods sometimes have false results and cause problems in diagnosis and treatment procedures. Be-cause of the difference between biologic nature of these two lesions, a distinguishing method for differentiation is necessary for proper identification.

Mast cells are members of the hemato-logic system that secrete various mediators. These cells play an important role in patho-logic, immunologic and physiologic events (8).

The effect of these mediators on tumors includes angiogenesis, suppression of the immune system, destruction of the extracel-lular matrix, proliferation and apoptosis (9).

It has been reported that mast cells are involved in gastric carcinoma and skin tu-mors, lung adenocarcinoma, and breast cancer (10-13). These cells often accumu-late in the margin of the solid tumors within stroma of the fibrous tissue that surrounding benign and malignant tumors (14).

According to investigations, the biologic nature of the connective tissue of benign and malignant tumors differs in the pattern of presence of mast cells (13-15-16).

Since the biologic nature of benign and malignant tumors bears an association with the density of mast cells, it seems that we can use this index to distinguish these two tumor .The aim of this study was to explore the difference between mast cells densities within fibrotic capsule surrounding enchon-droma and well-differentiated chondro-sarcoma.

Methods

The study was a descriptive-analytic, ret-rospective. Sample cases were collected from archives of Pathology Deptartment of Cancer Institute and Central Pathology Deptartment of Imam Khomeini Hospital between 1999 and 2008. In all cases diag-nosed as chondroma, enchondroma, poorly differentiated chondro-sarcoma, well differ-entiated chondrosarcoma and grade I chondrosarcoma were selected.

Proper histopathologic slides with enough capsular tissue were chosen. 12 enchondroma and 15 well- differentiated chondrosarcoma were selected for this study. Toluidine blue was used for mast cells staining. The slides were dried in ov-en, dipped in alcohol for a few seconds, and then put in Xylol (Xylene) solution for 2 seconds, then the sections were once more dipped in alcohol for a few seconds, and stained by 1% aqueous toluidine blue for 30 seconds. To complete the staining process, the specimens were washed in distilled wa-ter, dried, passed through alcohol, and placed in Xylol and mounted one slides.

Table1.The index of mast cells counting in enchondroma and well-differentiated chondrosarcoma groups

Lesion	cases														
Enchondroma	0.3	0.3	0	0	0.1	0	0.2	0.1	0	0	0	0.2			
Well-differentiated chondrosarcoma	0.2	0.2	0.5	0.2	0.1	0	0.1	0.3	0.3	0.7	1.3	0.1	0.3	0.4	0

Mast cells were counted by light microscopy (Zeiss,Germany) at ×1000 magnification. Mast cells of capsules were counted in 10 high power fields. The average number of mast cells in 10 high power fields was evaluated and determined as an index for each corresponding lesion. Each lesion counts twice and counting was performed blindly.

Statistical Analysis

We calculated the mean of the number of mast cells in 10HPF as an index for each sample and also mean and standard deviation of mast cell index for each lesion. Mann-Whitney test was used to examine the results. All the results were expressed in mean±SD. Sensitivity, specificity, positive and negative predictive values were determined. We also conducted receiver operating characteristic (ROC) analysis to measure the effectiveness of mast cells density parameter in differentiating enchondroma and well-differentiated chondrosarcoma. The ROC curve and area under curve were also determined using SPSS (version 15.0, Chicago, US) software. Area under the ROC curve was used as a summary index of diagnostic effectiveness. P<0.05 was considered statistically significant.

Results

With toluidine blue staining, mast cells were blue and contained purple granules (fig 1 and 2).

The minimum average number of mast cells in both enchondroma and well-differentiated chondrosarcoma was 0. The maximum average number of mast cells in enchondroma and well-differentiated chondrosarcoma were 0.3 and 1.3, respectively. The index of mast cells counting in two groups is shown in Table1.

The average index for enchondroma was 0.1±0.12 and for well-differentiated chondrosarcoma was 0.13±0.33, which showed a significant difference in the number of mast cells in the fibrous capsule of two lesions (p=0.028).
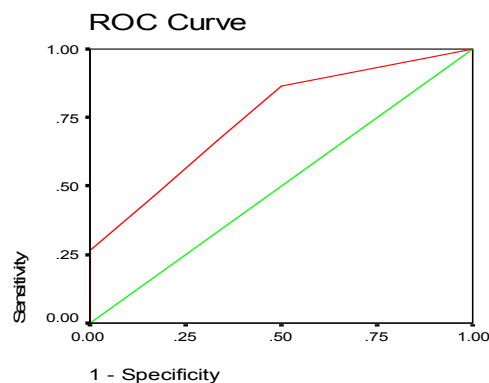
The area under ROC curve was 0.75

Chart. 1. The area under the ROC curve was 0.75. Comparison of the corresponding points in ROC curve, depict a cut-off point = 0.15, with positive predictive value 61%, negative predictive value 71%, specificity of 33.3% and sensitivity of 66.7%, achieved.
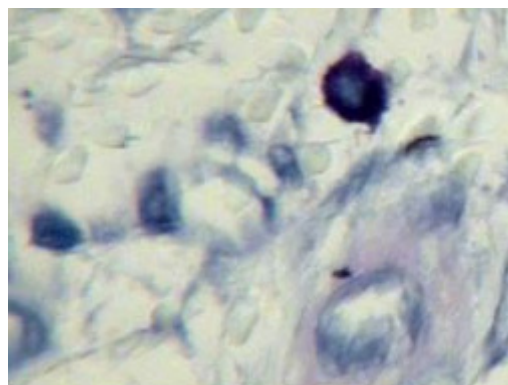

Fig1. Mast cell within fibrous capsule (Toluidine blue staining, ×1000).

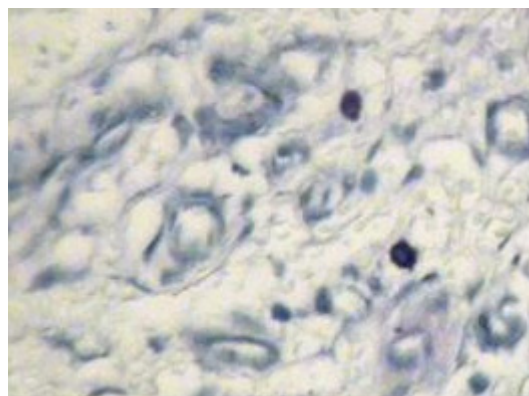Fig 2.Two mast cells between blood vessels (Toluidine blue staining, ×400).

(Chart1). Comparison of the corresponding points in ROC curve, revealed a cut-off point = 0.15, with positive predictive value 61%, negative predictive value 71%, specificity of 33.3% and sensitivity of 66.7%, achieved (p=0.025). As a result, if the aver-

age density of mast cells in the fibrous capsule surrounding the lesions was 0.15 or higher, the lesion could be a well-differentiated chondrosarcoma with a probability of 61% and if it was lower than it could be 0.15 could be a enchondroma with probability of 71% (positive predictive value 61% and negative predictive value 71%,).

Discussion

In this study, density of mast cells in fibrous capsule of enchondroma and well-differentiated chondrosarcoma were different. Mast cells density in chondrosarcoma was more than in enchondroma and this difference was statistically significant (p=0.028). We found that if the average number of mast cells was 0.15 or higher the lesion should be a well-differentiated chondrosarcoma with the probability of 61%, and if it was less than 0.15 an enchondroma with a probability of 71%.

Previous studies have shown that the mast cells numbers will be increased as a lesion undergoes malignant transformation (17-20).

By increasing the degree of malignancy, mast cells migration will be more and their presence will be more remarkable (13-15-16-21). Previous evidence on increased number of mast cells in malignant lesions versus benign ones is in harmony with findings of the current study which showed an increase in number of mast cells in chondrosarcoma (12,13).

A greater number of mast cells within the capsule of chondrosarcoma could also be an indicative of presence of absorptive cytokines emanating from tumoral tissue. With the biologic activity of tumor, cytokines attract mast cells and by increasing the tumor extent, mast cells attraction will be increased further.

Kessler et al. showed that implantation of tumoral cells within the human fetus causes a 40 folds increase of mast cells density around the implantation site in comparison to normal tissue (17).

In another study, Poole et al, showed that mast cells migrate towards factors that released from various types of tumors (22).

Release of some chemical materials such as IL-1, IL-3, FGF-2, and SCF from tumor cells and formation of a paracrine loop cause migration of mast cells. The level of these chemical materials in the stroma surrounding the tumors is related to degree of progression, invasion and tumor grade (23-25).

Presence of mast cells in connective tissue around tumors suggests two theories, either, the presence of these cells is induced by tumor and play an active role in carcinogenesis and its intensity, or helps the immune system with its inhibitory activity to arrest the extension of the tumor.

It has been shown that secretory heparin produced by mast cells in the presence of fibroblasts is a powerful inhibitor of growth for tumoral cells .This finding indicates the inhibitory role of mast cells in carcinogenesis (20).

It has been shown that injection of mast cells suspension to animal models causes acceleration of tumor growth, but decreasing the number of mast cells could inhibit tumor growth (26).

Now, this raises a question: 'where do these cells have an inhibitory role and where have a stimulating effect on tumor cells?' Samoszuk etal, found the answer of this question in 2005. They explained the mast cells double role in carcinogenesis: first, mast cells granules contain various active biologic compounds such as heparin, histamine, tryptase and chymase. Some of these mediators have stimulatory effects and some like heparin have inhibitory and stimulatory effects at the same time. There are also other mediators such as fibroblasts and endothelial cells of blood vessels that produced by stromal cells adjacent to mast cells. This mediators affect on tumor growth. Thus, the net effect on tumor growth is a result of numerous and complex interplay of various components of mast cells granules and composition of other cells like fibroblast and vessels endothelium. Second, secretion manner of mast cells

changes on the basis of the environment in which they are located. Different compositions have various roles in different environment (20).

On the basis of mast cells role, it is obvious that these cells have an essential role in pathogenesis of tumors. Hence, it seems that we can control tumor growth using the medications affecting mast cells secretions. It has been shown that inhibition of degranulation of mast cells in animals by disodium cromoglycate (cromolyn) obviously suppresses tumor growth (27).

This study showed that mast cells density within fibrotic capsule of well differentiated chondrosarcoma is more than enchondroma and this difference is statistically significant. As a result, mast cells counting along other criteria can be a useful method in histopathologic differentiating of enchondroma from well-differentiated chondrosarcoma. Arriving at more confident results would be possible by utilizing more cases and study of specific mediators.

Conclusion

Average density of the mast cells in the surrounding fibrotic capsules of enchondroma and well-differentiated chondrosarcoma along with other criterions, could be a beneficial factor for histologically differentiation between these two lesions.

Acknowledgment

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5

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