Association of CXCR4 mRNA Expression with Clinicopathological Aspects of Invasive Breast Carcinoma

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Abstract

Introduction: Breast cancer is the most common malignancy in women of which majority histological type is Invasive (Ductal) Carcinoma of No Special Type (NST). The prognosis in breast carcinoma is influenced by many factors such as age, tumor size, degree of histology, and lymph node metastasis. Another factor in the development and metastasis of breast cancer is the chemokine receptor CXCR4 and its ligand, CXCL12. Studies state that the expression of CXCR4 in Breast Invasive Carcinoma associated with clinicopathologic aspects remain unclear. This study aims to determine differences in the level of CXCR4 mRNA expression between clinicopathologic aspects in breast carcinoma.
Methods: A total of 50 samples of formalin-fixed paraffin-embedded (FFPE) tissues diagnosed as invasive breast carcinoma (NST) are used in this study. Samples are divided into groups, namely with and without lymph node metastasis, age <45 years and >45 years, small and large size, low grade and high grade. CXCR4 mRNA expression is quantitatively examined by qRT-PCR. CXCR4 mRNA expression differences between various clinicopathologic aspects were analyzed by T-test or Kruskal-Wallis.

Results: Of the 50 samples, 26 samples (52%) revealed increased expression of CXCR4 mRNA compared to normal tissue. There were no significant differences in mRNA expression of CXCR4 between various prognostic factors (p> 0.05) such as the status of lymph node metastasis, histologic grading, size, and age. However, the expression of CXCR4 mRNA is increased in breast carcinoma when compared to normal breast tissue. Nonetheless the level of CXCR4 expression alone is not associated to clinicopathologic aspects in invasive breast carcinoma.

Conclusion: CXCR4 mRNA expression did not differ significantly between the various clinicopathological aspects of invasive breast carcinoma.

Keyword: invasive breast carcinoma; mRNA of CXCR4; clinicopathologic aspects.

INTRODUCTION

Breast carcinoma is the most common malignancy in women. The incidence is about 23% of all malignancies in women in developed countries and increasing in developing countries1,2. Currently more than half the incidence of breast cancer occurs in developing countries2,3. The most common histologic type of breast cancer is invasive (ductal) carcinoma of no special type (NST)1.

The prognosis in breast carcinoma is strongly influenced by age, menopausal status, tumor size, lymph node metastasis, histological grade and tumor stage at diagnosis1. Breast carcinoma has a favorable prognosis when diagnosed early, whereas in metastatic cases the mortality rate is very high. The five-year survival rate in patients with localized breast carcinoma is around 90% and decreased dramatically to 20% if it has metastasized4.

Metastasis is the multistep and multifactorial process that includes many factors5,6. The process begins with local invasion in primary organ followed by intravasation of tumor cells into blood or lymph vessels, following blood or lymph flow, then by extravasation, the tumor cells undergo proliferation and colonization in new site. One of the factors that play a role in the tumor growth and metastasis is chemokine CXCR44,7.

CXCR4 is a G protein-coupled receptors that bind chemokines CXCL12, a chemoattractant, that functions in the activation, differentiation and cell circulation. The interaction of CXCR4 and CXCL12 plays a role in the process of angiogenesis, proliferation, apoptosis and metastasis of tumor cells4,7. CXCR4 gene expression can be examined semi quantitatively using immunohistochemical techniques (IHC) or quantitatively using qRT PCR technique (Quantitative Real Time PCR)8,9. From existing, CXCR4 is associated with poor prognosis in breast carcinoma9–11. However, studies of CXCR4 in association with clinicopathological aspects, especially metastatic status in invasive breast carcinoma remain unclear. This Study is aimed to investigate the association between CXCR4 mRNA expression and metastatic status in invasive breast carcinoma. The study of CXCR4 expression in invasive breast carcinoma can further guide in determining prognosis and "molecularly targeted therapy" especially in the metastatic ones. This study
aims to see variations within the level of CXCR4 mRNA expression between clinicopathologic aspects in breast malignant neoplastic disease.

**MATERIAL AND METHOD**

**Study Design and Subject**

A cross-sectional study was performed between September 2015 until December 2016. A total of 50 formalin-fixed paraffin-embedded (FFPE) tissue samples of breast cancer which met the inclusion criteria obtained from the government hospital and private laboratories in Yogyakarta. The inclusion criteria include women, radical mastectomy procedure with lymph nodes, invasive breast carcinoma of no special type, no residual or recurrent case. The samples used consisted of 25 samples with lymph node metastasis and 25 samples without lymph node metastasis. Normal breast tissue is used as normal control. The data of lymph node status, tumor size, histologic grade, and age of the patient were secondarily collected from medical record. CXCR4 mRNA expression was obtained from qRT PCR examination.

**RNA Extraction**

RNA extraction and qRT-PCR process was performed in Anatomic Pathology laboratory, Faculty of Medicine, Universitas Gadjah Mada. Total RNA Mini Kit (Tissue) Protocol was used for RNA extraction. The protocol included sample preparation, cell lysis, RNA binding, washing, and RNA Elution. Samples prepared by cutting paraffin blocks with microtome in 7µ of thickness of as much as 5-8 pieces. Deparaffinization is then performed by adding a solution of xylol followed by absolute ethanol. Cell lysis is performed by adding 400µl of RB Buffer and 4 µl of β-mercaptopethanol into tissue pellet. 400 µl of 70% ethanol prepared in ddH2O (RNase and DNase-free) is used for RNA Binding. Washing step is performed by adding Wash Buffer (make sure ethanol was added). This process is repeated three times. Next is the addition of 50 µl of RNase-free water followed by centrifugation at 14,000 rpm for 1 minute to elute the purified RNA. Purified RNA are kept at -70 °C.

**PCR Analysis**

RNA amplification and PCR analysis was performed using One-Step qRT-PCR protocol. The composition of each component used are as follows: 6.4 µl of PCR-grade water, 10 µl KAPA SYBR FAST qPCR Master Mix, 0.4 µl deoxyuridine triphosphate, 0.4 µl Forward Primer (GAPDHF atau CXCR4 F), 0.4 µl Reverse Primer (GAPDHR atau CXCR4 R), 0.4µl KAPA RT Mix and 2µl Template RNA. Primers used to make standard GAPDH and CXCR4 are presented in Table 1.

**Table 1. GAPDH and CXCR4 primer sequences**

<table>
<thead>
<tr>
<th>Primer</th>
<th>Forward Primer</th>
<th>Reverse Primer</th>
</tr>
</thead>
<tbody>
<tr>
<td>GAPDH</td>
<td>5’-GCA TCC TGG GCT ACA CTG AG-3’</td>
<td>5’-TCC ACC CTG TTG CTG TA-3’</td>
</tr>
<tr>
<td>CXCR4</td>
<td>5’-AGC ATG ACG GAC AAG ATG GAC TAC C-3’</td>
<td>5’-GAT GAT ACC CTT ACA.C3</td>
</tr>
</tbody>
</table>

**Table 2. Cycling Protocol of RT-PCR**

<table>
<thead>
<tr>
<th>Step</th>
<th>Temp</th>
<th>Duration</th>
<th>Cycles</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reverse Transcription</td>
<td>42 °C</td>
<td>5 min</td>
<td>-</td>
</tr>
<tr>
<td>Enzyme inactivation</td>
<td>95 °C</td>
<td>3 min</td>
<td>-</td>
</tr>
<tr>
<td>Denaturation</td>
<td>95 °C</td>
<td>3 sec</td>
<td>40</td>
</tr>
<tr>
<td>Annealing/extension</td>
<td>60 °C</td>
<td>≥20 sec</td>
<td>40</td>
</tr>
</tbody>
</table>
RT PCR results in the form of a graph and number which is then converted into expression value using a formula as shown below:

\[ \text{CXCR expression} = 2^{-\delta \delta CT} \]

\[ \delta CT : \text{PCR score of CXCR4} - \text{PCR Score GAPDH} \]

\[ \delta \delta CT: \delta CT \text{ sample} - \delta CT \text{ normal tissue} \]

**Statistical Analysis**

The data obtained were statistically processed to find differences of CXCR4 mRNA expression level between variables. The normally distributed data were analyzed using T-test and non-normally distributed data were analyzed using Mann-Whitney U Test or Kruskal-Wallis Test. All data analysis were performed in SPSS version 21 (IBM). A p-value < 0.05 was considered statistically significant.

The ethical problem that may arise in this study is the confidentiality of the patient's identity. In this study, the data to be submitted was in the form of a code without mentioning the name and laboratory number of the research subject. The identity will be converted into a code according to each group.

The ethical clearance letter issued by Medical and Health Research Ethics Committe (MHREC) Faculty of Medicine Gadjah Mada University with Ref number: KE / FK / 210 / EC / 2015.

**RESULT**

Fifty samples of breast cancer FFPE tissues are used in this study. The samples were divided into two groups, as many as 25 samples with axillary lymph node metastases and 25 samples without metastatic lymph nodes. Of the 25 samples with metastatic lymphnodes, 13 samples were from private laboratory while the rest were from the government hospital. Of the 25 tissue samples without metastasis, 12 samples were from private lab and the rest were from the government hospital. Normal breast tissue from government hospital was used as normal control.

Of the 50 samples obtained, there were 26 samples (52%) which revealed increased expression of CXCR4 mRNA compared to normal tissue. The samples which produce higher level of expression both for group with (10 of 12 samples) and without lymph node metastasis (11 of 13 samples) were derived from government hospital tissues. These results are in contrast with samples coming from private laboratories where only 2 of 13 samples had high expression in the group with lymph node metastasis and 4 of 12 samples in the group without lymph node metastasis. This study also obtained three samples with very high CXCR4 expression (more than 1000) in non-metastatic group compared to the normal control (36.4). (data not shown)

There were no significant differences between CXCR4 mRNA expression level with respect to lymph node status, tumor size, histologic grade, as well as age of patient observed in this study (table3).

<table>
<thead>
<tr>
<th>Variable</th>
<th>CXCR4</th>
<th>p (sig)</th>
</tr>
</thead>
<tbody>
<tr>
<td>lymph node status</td>
<td></td>
<td></td>
</tr>
<tr>
<td>negative</td>
<td>36.75 (5.27-3326.98)</td>
<td>0.907 (NS)</td>
</tr>
<tr>
<td>positive</td>
<td>29.85 (2.14-388.02)</td>
<td></td>
</tr>
<tr>
<td>Tumor size</td>
<td></td>
<td></td>
</tr>
<tr>
<td>T1 (n=2)</td>
<td>71.83 (6.49-137.18)</td>
<td></td>
</tr>
<tr>
<td>T2 (n=19)</td>
<td>39.39 (2.14-1351.17)</td>
<td></td>
</tr>
<tr>
<td>T3 (n=29)</td>
<td>29.85 (4.28-3326.98)</td>
<td>0.756(NS)</td>
</tr>
<tr>
<td>Grade</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 (n=0)</td>
<td>1.65 (0.99)</td>
<td>0.99(NS)</td>
</tr>
<tr>
<td>2 (n=9)</td>
<td>1.65 (0.65)</td>
<td></td>
</tr>
<tr>
<td>3 (n=41)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Age & ≤ 45 y & 1.51 (0.51) & 0.41 (NS) \\> 45 y & 1.70 (0.78) \\

**DISCUSSION**

Various biological markers have been used as marker for prognosis and treatment of breast cancer. Currently the most commonly used marker in breast cancer is hormonal receptors such as ER, PR, and Her2\(^1\). Other markers that are currently being researched and developed is chemokine receptor. Chemokines are chemotactic molecules which play role in inflammatory process, immune system, and also plays a role in tumor progression. Several studies have proven the role of chemokines in determining the tendency and sites of tumor metastasis\(^4\).

This study discusses mRNA expression of CXCR4 associated with clinicopathological aspects especially metastatic status. qRT-PCR method is used to assess the level of CXCR4 mRNA expression quantitatively. qRT-PCR examination using formalin-fixed paraffin-embedded (FFPE) is routinely used and the results are quite satisfactory. Although the expression of mRNA from FFPE is lower than fresh tissue (without fixation), but recent studies showed accurate value in qRT-PCR using FFPE(12–14). Since the frozen sample is limited and requires a large area thus the use of FFPE for qRT PCR examination become a solution\(^15\). Factors that should be considered in qRT-PCR examination using FFPE include:

- Formalin used is a buffer with normal pH because the acidic conditions can lead to degradation of mRNA\(^15\).
- Time fixation from tissue obtained is immediately after removal from the bodies to prevent autolysis process\(^13,16\).
- Optimal fixation time is 12-48 hours, because in some studies revealed decreasing mRNA expression after fixed more than 24 hours\(^13,15,17\).
- Temperature and storage time of FFPE. Storage temperature should be 4 °C with a storage time of no more than one year, although this is still being debated\(^15,15\).

In this study, the level of expression of CXCR4 mRNA were higher in tissues derived from hospital compared to tissues derived from a private lab. It is possible because private laboratories receiving tissue from different areas with different types and duration of fixation which may affect mRNA expression. The use of non-buffered formaline and delayed tissue fixation also decreased mRNA expression as experienced by private laboratories who does not have facilities as complete as government hospitals.

In this study, the level of CXCR4 mRNA expression between breast cancer with lymph node metastasis and without metastasis were not significantly different. This is consistent with several studies that have been done before where the expression of CXCR4 was not associated with incidence of lymph node metastasis however is more associated with a poor prognosis\(^10,18\). Moreover in some studies CXCR4 expression in the breast is associated with bone metastasis\(^18,19,20\). This may be due to CXCR4 needs its ligands, namely CXCL12, wherein the difference in expression levels of CXCL12 between primary organ and sites of metastasis has an important role for metastasis\(^6,8,11,21,22\). Existing studies revealed that CXCL12 expression have more roles in the process of metastasis than CXCR4 expression\(^21,22\). As the study conducted by Wei Wu et al stated the significant difference between the levels of CXCL12 in primary organs and lymph nodes, while tumors with low CXCL12 levels at the primary organ and high levels at the lymph nodes have a tendency to metastasize\(^22\). Even the study about ovarian tumor has proven that CXCL12 can be used as a predictive factor of metastasis while CXCR4 cannot\(^23\). According
to Hassan et al, mortality of breast cancer increases in patients with both high expression of CXCR4 in the tumor and low CXCL12 in the plasma compared to the increase CXCR4 in tumor alone or lower CXCL12 level in the plasma alone. Nevertheless, several other studies have the opposite results where the expression of CXCR4 is associated with lymph node metastasis in breast cancer. Although there is association between the increased expression of CXCR4 with incidence of lymph node metastasis, they convey varying results. A study by Kato et al found significant differences in the expression of CXCR4 protein via IHC staining on the expression pattern and the number of lymph nodes, although there was no significant difference between the groups with and without metastasis of lymph node metastasis. Variations of the expression pattern and vague cut-off point made it difficult to certainly conclude the role of CXCR4 in breast carcinoma using immunohistochemistry.

The process of metastasis is a complex process influenced by many factors. Pretranscription and posttranscription factors, epigenetic changes, and microenvironment greatly affect the expression of CXCR4. Normal tissue cells such as fibroblasts and endothelial cells express CXCR4 albeit in smaller amounts when compared with tumor tissue. In addition, the hypoxic conditions in primary tumor can lead to increased expression of CXCR4. This may explain why there are three samples of breast cancer without lymph node metastasis have CXCR4 mRNA expression levels were very high. After exploring the patient history, two of the three samples obtained had undergone neoadjuvant chemotherapy. Chemotherapy causes cells undergo hypoxia and necrosis. On the other hand hypoxic conditions triggered tumor cells to increase the expression of CXCR4. Subsequently hypoxic conditions will trigger cancer cells to express CXCR4 binding to CXCL12 endothelial cells involved in angiogenesis and migration of cancer cells into the bloodstream. Nevertheless, the increased expression of CXCR4 alone without the support of other microenvironment factors are not enough to cause metastasis.

Prognostic factors such as size, grading or age appear not directly related to the level of CXCR4 expression. This is in line with previous studies that revealed CXCR4 expression was not directly related to the clinicopathological features such as age, age and tumor size.

Hypoxic conditions is greatly influenced by the size and histological grade of the tumor. The larger the size and the degree of histological tumor would cause cancer cells susceptible to hypoxic conditions, which in turn triggers the process of angiogenesis and metastasis. Younger age in breast cancer is also associated with poorer prognosis because it deals with higher progressivity. But in study there has been no significant difference in mRNA expression levels of CXCR4 in all three clinicopathological aspects above which is in line with other existing research. This is possible because the complexity of environmental factors and processes that confound the influence so that CXCR4 expression alone can not be used as the sole predicting factor.

Although this study found no significant association between CXCR4 mRNA expression with clinicopathological features in breast carcinoma, the role of CXCR4 on the breast should not be ignored. The administration of drug that inhibits the expression of CXCR4 proved to inhibit metastasis when given with chemotherapy. This is possible because the administration of an antagonist of CXCR4 can inhibit CXCR4-CXCL12 binding that plays a role in the
progression, invasion and metastasis of tumors. 

This study has several limitations. One of them is limited clinical information from a sample in which the status of distant metastases (lung, bone, liver and other tissues) is unknown. So it is still possible samples without lymph node metastasis with higher expression of CXCR4 mRNA have metastasized to other organs. Sample variations and pre-analytic conditions such as type and duration of fixation can not be controlled because samples came from a wide area and diverse handling system so that it can affect the level of expression of CXCR4 mRNA. In this study, the samples used are not separated between the tumor tissue and surrounding tissue which can affect the outcome, even though it has been minimized by using the controls of normal breast tissue as a correction factor.

CONCLUSION

From this study we can conclude that CXCR4 mRNA expression was not associated with clinicopathological features of invasive ductal carcinoma NST. There were increased expression of CXCR4 mRNA in tumor tissue compared to normal breast tissue.

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