Overexpression of long non-coding RNA HOTAIR predicts a poor prognosis in patients with acute myeloid leukemia

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Abstract. The long non-coding RNA, HOX transcript antisense intergenic RNA (HOTAIR), has been indicated to have involvement in a number of cancers, however, its role in acute myeloid leukemia (AML) is unknown. The present study aimed to investigate the pattern of HOTAIR expression in AML and to evaluate its clinical significance in tumor progression. Quantitative polymerase chain reaction was performed to examine the HOTAIR expression in mononuclear cells from the bone marrow (BM) or peripheral blood specimens of 85 patients with newly diagnosed AML. The association of HOTAIR expression with the clinicopathological factors and prognosis of AML patients was statistically analyzed. The expression of HOTAIR was significantly upregulated in the AML patients compared with the healthy controls (mean expression value, 3.87±0.29 vs. 1.28±0.09; P<0.001), and markedly decreased in the patients post-treatment compared with pre-treatment (4.76±0.47 vs. 2.81±0.27; P<0.001). Moreover, high levels of HOTAIR were associated with higher white blood cell and BM blast counts (P<0.001 and P=0.001, respectively), and lower hemoglobin and platelet counts (P=0.007 and 0.001, respectively). Patients with a high level of HOTAIR expression had relatively poor overall survival (OS; 20.5 vs. 32.1 months, P=0.001) and relapse-free survival (21.5 vs. 33.6 months, P=0.001) times compared with those with a low level of HOTAIR expression. These data demonstrated that HOTAIR expression was upregulated in newly diagnosed AML patients and was associated with leukemic burden, and DFS and OS times. HOTAIR may represent a biomarker of a poor prognosis and is a potential therapeutic target for AML treatment.

Introduction

Acute myeloid leukemia (AML) is a heterogeneous hematopoietic malignancy characterized by the rapid accumulation and malignant proliferation of immature myeloid progenitors in the bone marrow (BM) and peripheral blood (PB) (1). Without treatment, AML quickly becomes fatal, and historically, it has always been associated with a poor prognosis. However, AML treatment has markedly improved over the last few decades, with improvements in risk assessment, post-remission chemotherapy and hematopoietic stem-cell transplantation (2,3). However, even though complete remission (CR) is achieved after chemotherapy by the majority of AML patients, only ~20% of obtain relatively long-term relapse-free survival (RFS) (4). Thus, in order to improve the diagnosis, prevention and treatment of this disease, detailed knowledge of the mechanisms that form the basis of AML development and progression must be acquired. Recently, it has been shown that long non-coding RNAs (lncRNAs) play a crucial role in hematopoietic differentiation and hematological malignancies, including AML (5).

Transcriptome analysis by tiling arrays and RNA sequencing has revealed that only 2% of the human genome is dedicated to the transcription of protein coding sequences and that >90% of the genome is transcribed as non-coding RNAs (6,7). IncRNAs are transcripts of >200 nucleotides and conventionally cannot be translated into proteins to participate in a large number of biological processes (8,9). However, recent studies hypothesized that a number of IncRNAs are key developmental regulators that are involved in cell homeostasis and proliferation (9,10). Notably, increasing numbers of studies are indicating that the abnormal expression of certain IncRNAs is associated with tumor growth, carcinogenesis or metastasis in a range of malignancies (11-15).

Hox transcript antisense intergenic RNA (HOTAIR) is a 2,158-bp IncRNA that is transcribed from the antisense strand of the homeobox C gene locus of chromosome 12. HOTAIR coordinates with chromatin-modifying enzymes and regulates gene silencing (16). Several recent studies have identified the aberrant expression of HOTAIR in a number of cancer types, including breast, colon, bladder, renal, pancreatic, cervical and lung cancer, and a high level of HOTAIR expression has been correlated with enhanced breast, colon and gastric cancer
metastasis. In addition, HOTAIR-knockdown is able to inhibit the invasion and proliferation of cells, as well as altering cell cycle progression and inducing cells apoptosis, thus indicating that HOTAIR may function in the modulation of cancer progression (17-19).

To the best of our knowledge, no previous studies exist concerning the expression status, prognostic value and role of HOTAIR in AML. Thus, the aim of the present study was to investigate the correlation of HOTAIR expression with clinicopathological features and the prognosis of the patients with AML. The findings may improve our understanding of the roles and the clinic implications of HOTAIR in the development and progression of AML.

Patients and methods

Patients and specimens. This study was approved by the Research Ethics Committee of Wenzhou Central Hospital (Wenzhou, Zhejiang, China). Written informed consent was obtained from all patients according to the committee’s regulations. Between February 2011 and August 2014, 85 patients in the Department of Hematology (Wenzhou Central Hospital) were diagnosed with AML according to the World Health Organization classification system (21). The clinical characteristics of all the AML patients are summarized in Table I. A total of 66 patients received standard cytarabine (100 mg/m² daily, days 1-7) plus daunorubicin (45 mg/m² daily, days 1-3) 7+3 induction chemotherapy. Specimens were obtained from the BM or PB (peripheral blood cell; HGB, hemoglobin; PLT, platelet). Of the 85 enrolled patients, 5 patients presented with AML of type M1, 30 of type M2, 12 of type M4, 32 of type M5 and 6 of type M6, according to the French-American-British (FAB) criteria (20). The cohort consisted of 45 males and 40 females, with a medium age of 45.2 years (range, 19.3-72.4 years). The median leukocyte count at diagnosis was 52,897/µl (range, 793-327,100/µl; normal range, 4,000-11,000/µl). Of the 85 enrolled patients, 5 patients presented with AML of type M1, 30 of type M2, 12 of type M4, 32 of type M5 and 6 of type M6, according to the World Health Organization classification system (21). The clinical characteristics of all the AML patients are summarized in Table I. A total of 66 patients received standard cytarabine (100 mg/m² daily, days 1-7) plus daunorubicin (45 mg/m² daily, days 1-3) 7+3 induction chemotherapy. Specimens were obtained from the BM or PB (peripheral white blood cells; >50x10⁹/µl) of the patients at the time of diagnosis, and from 33 patients who were in CR following two cycles of chemotherapy. Additionally, 40 PB specimens were obtained from healthy donors as negative controls. Patients who achieved CR were then administered high- or medium-dose cytarabine-based chemotherapy for consolidation according to their physical condition. The 33 patients with CR were then followed up for a median time of 22 months (range, 9-40 months), and the data were censored when the patients relapsed or succumbed.

RNA extraction and reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Mononuclear cells from PB or BM specimens were isolated by Ficoll density gradient centrifugation (400 x g, 30 min; Hao Yang Biological Manufacture, Tianjin, China), and then washed and pelletted. Additionally, PB cluster of differentiation (CD)34⁺ cells from healthy donors were obtained using magnetic bead separation (EasySep Human CD34 Positive Selection kit; Stem Cell Technologies, Vancouver, BC, Canada). Total mRNA was extracted using TRIzol reagent (Invitrogen Life Technologies, Grand Island, NY, USA) according to the manufacturer’s instructions. The quality and concentration of RNA were determined using a Nanodrop 2000 (Thermo Fisher Scientific Inc., Wilmington, DE, USA). Next, 1 µg total RNA was reverse transcribed from each sample to synthesize cDNA using the RT reagent kit (Fermentas, Glen Burnie, MD, USA) according to the manufacturer’s instructions. qPCR was performed using the ABI 7300 Sequence Detection System with primer pairs and SYBR Green PCR Master Mix (Applied Biosystems, Foster City, CA, USA). The primer sequences used were as follows: HOTAIR forward, 5'-CAGTGGGGGAACCTGTGACGTG-3' and reverse, 5'-GTGGCTCTGGTGCTCCTTACC-3'; β-actin forward, 5'-CACCATGGCAATGAGCGGTTCC -3' and reverse, 5'-TGAAGTGCTGAGGAGCGTCC-3' and reverse, 5'-CAGTGGGGGAACCTGTGACGTG-3' and reverse, 5'-GTGGCTCTGGTGCTCCTTACC-3'. The amplification profile was 95°C for 5 min, followed by 42 cycles of denaturation at 95°C for 15 sec, then annealing and extension at 60°C for 60 sec. The comparative Ct method (ΔΔCt) was used for the quantification of gene expression. The relative expression of HOTAIR to β-actin was calculated using the equation 2⁻ΔΔCt, where ΔΔCt = ΔCt HOTAIR - ΔCt β-actin. Each sample was analyzed in triplicate and the mean expression level was calculated.
Statistical analysis. Statistical analysis was performed with SPSS 16.0 for Windows (SPSS Inc., Chicago, IL, USA). Data are expressed as the mean ± standard deviation. The Kruskal-Wallis non-parametric test was used to evaluate the difference in HOTAIR expression between the AML patients and the healthy controls. The paired t-test was used to evaluate the difference in HOTAIR expression prior to and following chemotherapy. Pearson’s χ² test was used to evaluate the association between HOTAIR expression and clinicopathological characteristics. Survival curves were plotted using the Kaplan-Meier product-limit method, and differences between survival curves were tested using the log-rank test. RFS was defined as the time between the achievement of CR and the time of the hematological relapse or the last follow-up. Overall survival (OS) was defined as the time between the moment of diagnosis and mortality or the last follow-up. Cox regression analysis in a forward stepwise method was used to evaluate the effect of multiple independent prognostic factors on survival outcome. Differences were considered to be statistically significant when P<0.05.

Results

HOTAIR is overexpressed in AML patients. The HOTAIR expression levels were detected in BM/PB samples from the patients with AML and the healthy controls by RT-qPCR. As shown in Fig. 1A, the expression of HOTAIR was significantly upregulated in the AML patients compared with the healthy controls (mean expression value, 3.87±0.29 vs. 1.28±0.09; P<0.001). Additionally, the 33 AML patients who achieved CR following one or two cycles of chemotherapy were monitored for HOTAIR expression during the course of treatment. The mean expression value of these AML patients markedly decreased when CR was achieved after chemotherapy (mean expression value, 4.76±0.47 vs. 2.81±0.27; P<0.001).

Correlations between the expression of HOTAIR and the clinicopathological factors in AML patients. To identify the clinical relevance of HOTAIR expression in AML patients, the correlation between HOTAIR expression and clinicopathological parameters was assessed. Those AML patients with HOTAIR expression levels at less than the median value (3.87) were assigned to the low expression group (mean expression value, 1.98; n=52), and those with expression above the median value were assigned to the high expression group (mean expression value, 6.84; n=33). As shown in Table I, high levels of HOTAIR were associated with higher white blood cell and BM blast counts (P=0.001 and P=0.001; respectively), and a lower hemoglobin level and platelet count (P=0.007 and 0.001; respectively). However, other clinical characteristics, including age, gender and FAB subtype were not directly associated with the high expression of HOTAIR.

Association between HOTAIR expression and clinical outcomes of AML patients. In total, 66 patients received standard induction chemotherapy, The CR rate following two cycles of chemotherapy was 27.3% (9/33) in the high expression group compared with 46.2% (24/52) in the low expression group (P=0.082). Despite the high CR rate in the low expression group, there was no statistically significant difference between the values of the two groups (P=0.082). The 33 patients who achieved a CR were followed up for a median time of 22 months (range, 9-40 months). OS and RFS survival curves in high expression and low expression groups are shown in Fig 2. Patients with high HOTAIR expression exhibited significantly poorer OS (hazard ratio, 3.37; 95% confidence interval, 0.99‑8.31; P=0.008). Statistical values for HOTAIR expression and other clinical parameters derived from the Cox stepwise proportional hazards model are presented in Table II.

Discussion

It is becoming evident that mammalian genomes encode thousands of IncRNAs, and multiple lines of evidence increasingly support the idea that certain IncRNAs could be used as biomarkers that predict the prognosis of tumor targets of human cancer (22,23). HOTAIR was first identified as one of 231 IncRNAs that are associated with the human HOX loci, which binds to polycomb repressive complex 2 (PRC2), a transcriptional co-repressor, and recruits it to silence the target genes (18). HOTAIR is also able to interact with a second histone
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Table II. Cox multivariate analysis of factors for overall survival in AML patients.

<table>
<thead>
<tr>
<th>Factors</th>
<th>Hazard ratio</th>
<th>95% confidence interval</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>WBC</td>
<td>0.87</td>
<td>0.44-1.39</td>
<td>0.047</td>
</tr>
<tr>
<td>HGB</td>
<td>0.93</td>
<td>0.74-1.79</td>
<td>0.088</td>
</tr>
<tr>
<td>PLT</td>
<td>2.07</td>
<td>1.13-3.62</td>
<td>0.034</td>
</tr>
<tr>
<td>Blasts in BM</td>
<td>1.15</td>
<td>0.41-1.77</td>
<td>0.078</td>
</tr>
<tr>
<td>Complete remission</td>
<td>1.78</td>
<td>1.14-2.55</td>
<td>0.031</td>
</tr>
<tr>
<td>HOTAIR expression</td>
<td>3.37</td>
<td>0.99-8.31</td>
<td>0.008</td>
</tr>
</tbody>
</table>

WBC, white blood cell; HGB, hemoglobin; PLT, platelet; BM, bone marrow; HOTAIR, HOX transcript antisense intergenic RNA.

Figure 2. Kaplan-Meier survival curves for overall survival (OS) and relapse-free survival (RFS) according to HOX transcript antisense intergenic RNA (HOTAIR) expression from 32 AML patients in CR. (A) The OS rate of AML patients in complete remission (CR) with high or low HOTAIR expression. (B) The RFS rate of AML patients in CR with high or low HOTAIR expression. The P-value was calculated using the log-rank test.

modulation complex, the LSD1/CoREST/REST complex, which functions by coordinating the targeting of PRC2 and LSD1 to chromatin for methylation of coupled histone H3K27 and demethylation of K4 (24). HOTAIR is believed to be an oncogene due to its elevated expression levels in a number of cancer types, and due to its ability to mediate the invasion and metastasis of cancer cells. For example, Huang et al revealed that HOTAIR expression in cervical cancer tissues was significantly upregulated compared with the matched non-tumorous tissues, and increased HOTAIR expression was significantly correlated with the International Federation of Gynecology and Obstetrics stage, lymph node metastasis, depth of cervical invasion and tumor size (25). HOTAIR was associated with the carcinogenesis and invasion of gastric adenocarcinoma, HOTAIR-targeted RNA interference can reduce the proliferation, invasion and migration abilities of gastric cancer cell lines (26). Similarly, dysregulation of HOXA5 expression has also been reported in association with tumorigenesis and progression in lung cancer (27-29). These observations suggest that HOTAIR has a direct role in the modulation of cancer progression and may be useful in patients with cancer as a novel prognostic or progression marker. However, in AML, the HOTAIR expression status and its prognostic roles are unclear.

In the present study, through the use of qPCR, it was confirmed for the first time that the expression of IncRNA HOTAIR was markedly unregulated in patients with newly diagnosed AML compared with healthy controls; these results were consistent with other studies regarding solid tumors. Moreover, the level of HOTAIR expression was significantly decreased following chemotherapy when patients achieved CR, indicating that HOTAIR expression is consistent with tumor burden, and that HOTAIR expression can be used as a prognostic marker of relapse. In addition, the present results indicated that the upregulation of HOTAIR in AML patients was significantly correlated with higher white blood cell and BM blast counts, and a lower hemoglobin level and platelet count, which represented more aggressive clinicopathological features. Finally, AML patients with high HOTAIR expression tended to have poorer OS and RFS times compared with those with low HOTAIR expression, indicating that the expression of HOTAIR is significant in the classification of AML prognosis. Taken together, these data suggest that HOTAIR may function as an oncogene in the development of AML, and may represent a candidate prognostic biomarker for AML patients.

The aforementioned findings that HOTAIR overexpression was associated with aggressive tumor progression indicated that its possible prognostic value in AML patients should be investigated in the present study. According to the univariate and multivariate analyses, HOTAIR overexpression was identified as an independent predictor for the OS of AML patients, which was in agreement with recent findings in NSCLC and cervical cancer (25,26), suggesting that the detection of increased HOTAIR expression may aid in the identification of AML patients with a poor prognosis, and could therefore be a novel prognostic marker for AML patients.

In summary, the present study provides evidence for the first time that HOTAIR may act as an oncogenic gene in AML, and that it may represent a potential biomarker of poor prognosis and a potential therapeutic target for AML intervention. However, the precise molecular mechanisms behind the involvement of HOTAIR in AML require further investigation.

References