Differential expression of the Na\(^+\)/I\(^-\) symporter protein in thyroid cancer and adjacent normal and nodular goiter tissues

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Abstract. The ability of differentiated thyroid cancer and adjacent thyroid cells to concentrate iodine is dependent on their expression of a functional Na\(^+\)/I\(^-\) symporter (NIS). Thyroid cancer is insensitive to \(^{131}\)I treatment if the thyroid cells lack the ability to concentrate iodide. Thus, in this study, we aimed to determine whether the NIS protein was differentially expressed in thyroid cancer and various surrounding tissues. We recruited 114 cases of papillary thyroid carcinoma (PTC) and divided them into two groups: 60 patients of 9 males and 51 females with a mean age of 49.55 years who had PTC with surrounding nodular goiter tissue (simplified as GNG), and 54 patients of 8 males and 46 females with a mean age of 45.78 years who had PTC with surrounding normal tissue (Gnormal) after total or near total thyroidectomy. Formalin-fixed and paraffin-embedded tissue sections were prepared for immunohistochemical staining of the NIS protein and semi-quantitative analysis. The NIS protein was expressed in the basolateral membrane of the normal epithelium, while PTC and nodular goiter cells expressed NIS in the cytoplasm and basolateral membrane. The expression levels of the NIS protein were higher in the adjacent normal tissues compared with those of the surrounding nodular goiter tissues (P=0.002) and expression levels of the NIS protein were higher in PTC tissues compared with the surrounding nodular goiter tissues (P=0.008). The data from this study indicate that cancer-surrounding tissues may play a significant role in mediating the sensitivity of PTC patients to radioactive iodine treatment.

Introduction

Thyroid cancer is the most common endocrine neoplasm. Histopathologically, thyroid cancers can be classified into papillary, follicular, medullary and anaplastic carcinomas. Papillary and follicular carcinomas are derived from thyroid follicular epithelial cells and account for the vast majority of thyroid cancer. These two types of thyroid cancers are differentiated tumors with low grades. The prognosis of thyroid cancer is associated with the histological type of the cancer and the stage at diagnosis. For papillary thyroid cancer (PTC), the overall prognosis is excellent following proper treatment, including thyroidectomy, lobectomy or radioactive iodine (RAI) therapy (1).

RAI ablation, which removes all remnant or residual normal thyroid tissues, is an important element of therapy following initial surgery in patients with papillary and follicular thyroid carcinomas (1). The underlying mechanism of RAI treatment of thyroid cancer is based on the ability of thyroid follicular cells to concentrate iodine, which is dependent on the functional Na\(^+\)/I\(^-\) symporter (NIS) (2). NIS is a transmembrane glycoprotein with a molecular weight of 87 kDa, which transports two Na\(^+\) for each I\(^-\) into thyroid follicular cells. The loss of NIS expression in thyroid follicular cells may cause goiters or hypothyroidism. During thyroid cancer development, NIS expression has been reported to be reduced or lost. Thus, the detection of NIS expression was able to predict the outcome of RAI treatment in patients with thyroid cancer. Previous studies have demonstrated that the NIS protein was differentially expressed in differentiated thyroid carcinomas compared with normal thyroid tissues (2-4). By contrast, certain previous studies have found that PTC patients with normal NIS expression respond to RAI therapy (3,5,6). However, to date, few studies reporting NIS expression in PTC and their surrounding tissues have been published. Therefore, in the present study, we evaluated NIS expression in adjacent normal thyroid tissues in comparison to nodular goiter in PTC patients. The detection of NIS expression in surrounding normal thyroid tissues was able to predict the iodine uptake activity during RAI therapy of differentiated thyroid carcinomas.
Materials and methods

Study population. In this study, we first identified and reviewed the medical records from 600 patients who were diagnosed with PTC and underwent total or near-total thyroidectomy at the Affiliated Hospital of Qingdao Medical College between January 1, 2008 and January 1, 2011. Histology sections from these 600 patients were carefully re-examined by two pathologists for confirmation of the original diagnosis of PTC. Specimens of PTC and the surrounding normal tissues or the surrounding nodular goiter tissues were used in the current study, which generated two groups of cases: a group of 60 patients (52.63%; 9 males and 51 females with a mean age of 49.55±11.29 years) whose PTC had surrounding nodular goiter tissues (abbreviated as G_ng) and a second group of 54 patients (47.37%; 8 males and 46 females with a mean age of 45.78±11.12 years) whose PTC had surrounding normal tissues (abbreviated as G_normal) (Table I). 1. The study was approved by the Ethics Committee of The Affiliated Hospital of Qingdao Medical College, Qingdao, China. Written informed patient consent was obtained from the patient.

Immunohistochemistry. Formalin-fixed and paraffin-embedded tissue blocks from PTCs with G_ng and G_normal were obtained from the Department of Pathology at the Affiliated Hospital of Qingdao Medical College, and prepared in 5-µm thick sections for immunohistochemistry. Briefly, all tissue sections were deparaffinized in xylene, rehydrated in graded alcohol (100-50%) and endogenous peroxidase activity was blocked in 3% H₂O₂ solution in methanol for 5 min and the sections were washed with PBS three times for 2 min each. Next, the sections were incubated with 20% normal serum for 30 min and then with a rabbit polyclonal anti-NIS antibody (#696557; American Basic Gene Associate Bioscience, Inc., Chicago, IL, USA) at a dilution of 1:400 in PBS at 4°C overnight. The next day, the sections were washed three times with PBS for 2 min each and incubated with a secondary antibody from a PV-6000 kit (Zhongshan Golden Bridge Biotechnology, Beijing, China) for 15 min at room temperature. Next, the sections were incubated with 3,3’-diaminobenzidine tetrahydrochloride solution (DAB; Zhongshan Golden Bridge Biotechnology) after washing three times with PBS. The color reaction was stopped after a suitable color had developed or after a maximum of 10 min. The sections were briefly counterstained with hematoxylin. Finally, all sections were washed with distilled water, dehydrated through ascending alcohol and xylene washes and mounted with cover slips with a drop of mounting medium. Both positive and negative controls were used for each sample of tumor tissues and surrounding tissues.

Review and score of the immunostained tissue sections. All the immunostained tissue sections were reviewed and scored under a microscope for expression and localization of NIS protein by two pathologists independently and blindly. The scores of each section were compared and if there was a discrepancy, the two pathologists reviewed them again and reached a consensus. Briefly, five high-powered fields under the microscope were randomly chosen and one hundred cells in each field were counted. The staining scores (IHS) were calculated by combining an estimate of the percentage of immunoreactive cells (quantity score) with an estimate of the staining intensity (staining intensity score) (7). For the percentage of staining, 0 indicated no staining; 1, 1-10% of cells stained; 2, 11-50%; 3, 51-80%; 4, 81-100%. Staining intensity scores were as follows: 0, negative; 1, weak; 2, moderate; and 3, strong. The raw data were converted to the IHS by multiplying the quantity and staining intensity scores. The scores of IHS ranged from 0 to 12, with 0 indicating negative, 1 to 4 indicating weak, 5 to 8 indicating moderate and 9 to 12 indicating strong immunoreactivity. For multifocal immunoreactivity or significant differences in staining intensities between foci, the average of the least and most intense staining was recorded.

Statistical analyses. All statistical analyses were performed with SPSS version 17.0 (SPSS, Inc., Chicago, IL, USA). Statistical independent sample t-tests or t’ tests were used to determine the correlation of NIS expression between tumor and normal tissues. P<0.05 was considered to indicate a statistically significant result.

Results

Clinicopathological characteristics. In this study, we collected a total of 114 cases of PTC and divided them into two groups: 60 patients with PTC whose resected tissues contained adjacent normal thyroid epithelium (G_normal) and 54 patients with PTC whose surgical specimens contained nodular goiter tissues (G_ng). We found that tumor size (P=0.0004) and lymph node metastases (P=0.0000) were significantly different between these two groups. Additionally, G_normal patients had larger tumor sizes and more lymph node metastases than G_ng patients (Tables I and II).

Expression of NIS. We then assessed NIS expression in these tissues and found that NIS protein was expressed in the basolateral membrane of the normal epithelium, while nodular goiter cells expressed NIS in the cytoplasm and occasionally in the basolateral membrane (Fig. 1). By contrast, NIS protein was mostly expressed in the cytoplasm and rarely in the basolateral membrane of PTC cells (Fig. 1). Based on the percentage of staining and the staining intensity of the NIS antibody, we summarized NIS expression levels as ‘points’ in each section of PTC, G_normal and G_ng tissue. The 114 cases of thyroid cancer exhibited NIS staining with a score ranging between 0 and 12 points. The total NIS protein scores of the PTC and G_ng tissues were 348 and 276 points, respectively (P=0.008). In addition, we associated NIS expression with clinicopathological data from the PTC patients with surrounding G_ng tissue. We found that age <45 years (P=0.000), female gender (P=0.003), tumor size ≥1 cm (P=0.000), TNM stage I (P=0.017) and lymph node and distant metastases (P=0.026 and P=0.008, respectively) were associated with NIS expression.

IHS scores. Furthermore, as shown in Table II, the total scores of NIS expression in PTC and G_normal tissues were 366 and 351 points, respectively, which was not statistically significant (P=0.675). By contrast, the expression levels of
NIS protein in G\textsubscript{normal} tissue of tumors <1 cm in size were lower (21/351 points) compared with cancerous tissue (36/366; \(P=0.037\)). Moreover, NIS expression in cancerous tissue of G\textsubscript{normal} and G\textsubscript{NG} did not reach statistical significance.
In conclusion, NIS protein was differentially expressed in different surrounding tissues from PTC patients. This is the first study to report that expression levels of NIS protein were lower in surrounding nodular goiter tissues than in surrounding normal thyroid tissues. A previous study revealed that immunodetection of NIS protein predicted radioiodine uptake in thyroid cancer tissues (3), recurrent lesions (5), metastatic and recurrent disease (6). Thus, we suggest that the impaired NIS protein localization or various expression levels in the surrounding tissues may also affect the sensitivity of PTC to RAI therapy. Moreover, the current data further indicated that plasma membrane localization of NIS protein or induction of NIS cell membrane expression may improve the sensitivity of PTC to RAI therapy. Thus, detection of NIS protein expression and the localization of NIS in surrounding thyroid tissues may be useful to predict RAI therapy outcomes in PTC patients. In addition, based on NIS protein expression and localization, a physician administering nuclear medicine may be able to modify the $^{131}$I dose when treating PTC.

In conclusion, NIS protein was differentially expressed in surrounding normal thyroid tissues and nodular goiter tissues from PTC patients and may regulate the sensitivity of PTC patients to RAI therapy. Future studies should evaluate the association between NIS protein expression, sensitivity of RAI treatment and serum levels of iodine in PTC patients.
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References


