

Abnormality of p16/p38MAPK/p53/Wipl pathway in papillary thyroid cancer

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Objective: To investigate the expression of the p16/p38MAPK/p53/Wipl pathway in patients with papillary thyroid cancer (PTC) and its clinical significance.

Methods: The protein expressions of Wipl, p53, p38MAPK, and p16 in 70 cases of PTC tissues and 20 cases of normal thyroid tissues were detected by immunohistochemical staining. The correlations of Wipl protein high-expression with p53, p38MAPK, and p16 protein expressions were analyzed.

Results: The high-expression rate of Wipl protein in the PTC tissue was 64.3% (45/70), which was significantly different than that in normal thyroid tissue (0/20) ($P < 0.01$). There were no significant differences between the paired groups in terms of age, gender, tumor size, and lymph node metastasis (all $P > 0.05$). The Wipl protein high-expression was negatively correlated with the expressions of p38MAPK, p53 and p16 (r value was -0.620 , 0.356 and 0.550 , respectively, and all $P < 0.01$).

Conclusions: The p16/p38MAPK/p53/Wipl pathway is abnormal in PTC, and this abnormality may possibly be associated with the aberrantly up-regulated Wipl, which can induce inhibition of p38MAPK, p53 and p16.

Keywords: Thyroid neoplasms; wild-type p53-induced phosphatase 1; carcinoma, papillary; immunohistochemistry



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Introduction

Wild-type p53 induced phosphatase 1 (Wip1) is a member of the PP2C family of evolutionarily conserved protein phosphatases and a novel proto-oncogene (1,2). Originally described as a p53-regulated gene. In recent years, analysis of Wip1 has focused primarily on its role in tumorigenesis because of its overexpression in human tumors. Overexpression or activation of Wip1, as commonly seen in primary human tumors, could have a more general effect on multiple signaling pathways contributing to deregulation of several molecular networks.

p53 and p16 are key anti-oncogenes, whereas p38 mitogen-activated protein kinase (MAPK) is a key cell cycle regulator (3). Wip1 is a major inhibitor of p53 functions. The mutation or missing of p53 gene is the cause of many tumors (4) which is a main carcinogenic mechanism of

Wip1. Wip1 has been implicated as a negative regulator of p53 via its ability to attenuate p38 MAPK activity (5,6). p16, also known as mitotic inhibitor or multiple tumor suppressor gene, is one of the cell cycle regulator. p16 is a key anti-oncogene. Moreover, higher Wip1 expression is associated with lower p16 levels in primary human mammary carcinomas (7). And Wip1 can potentially down-regulate p16 expression by suppressing p38MAPK (8). However the downregulation of p53 and the DNA damage and DNA repair responses by WIP1 has cancer implications, particularly since the DNA damage response has recently been shown to play a crucial role as an early anti-cancer barrier (9-11). It is known that p38 mitogen-activated protein (MAP) kinase controls cellular pathways for proliferation, differentiation, development of the inflammatory response, and apoptosis. Wip1 also negatively regulates the stress responsive p38 MAPK pathway by

directly inactivating p38 through dephosphorylation (12). However phosphorylated p38 MAP kinase phosphorylates and activates p53 to cause cell cycle arrest or apoptosis. And Wip1 can control a feedback loop in the p38 MAP kinase-p53 signaling pathway (12). Thus, p16/p38MAPK/p53/Wip1 signal transduction pathways may play a crucial role in human tumors.

Not surprisingly, Wip1 has been implicated as an oncogene and Wip1 gene amplification and/or protein overexpression have been reported in a variety of human tumors including breast cancer, neuroblastoma, pancreatic cancer, ovarian cancer, and gastric cancer (7,13). However, its expression in papillary thyroid carcinoma (PTC) remains unclear. Up to now few studies have explored the role of p16/p38MAPK/p53/Wip1 pathway in the occurrence and development of PTC. In this study, by detecting the protein expressions of Wip1, p53, p38MAPK, and p16 in 70 cases of PTC tissues and 20 cases of normal thyroid tissues using immunohistochemical staining and analyzing their mutual correlation, we investigated the role of p16/p38MAPK/p53/Wip1 pathway in PTC. To our knowledge, this is the first report establishing the existence of the signaling pathway in PTC and may be a potential target for therapeutic intervention for treatment of PTC.

Materials and methods

Specimens and general information

PTC group

The specimens and clinical data of 70 patients with pathologically confirmed PTC who were treated in our hospital from March 2010 to December 2010 were retrospectively collected. There were 16 males and 54 females aged 15-64 years (mean: 41 years). Lymph node metastasis was noted in 92 patients, while 37 patients had no lymph node metastasis. TNM staging showed 39 cases in stage I, 12 in stage II, 14 in stage III, and 5 in stage IV. All patients were naive to chemotherapy and radiotherapy before the surgery. PTC specimens were obtained from these tissues immediately after surgical resection.

Control group

Pathologically confirmed normal thyroid tissues were obtained from 20 cases. There were 4 males and 16 females aged 20-55 years (mean: 42.3 years).

Main reagents

The main reagents included rabbit anti-human Wip1 polyclonal antibody (Santa Cruze); rabbit anti-human p-p38MAPK monoclonal antibody (Cell Signaling); and mouse anti-human p53/p16 monoclonal antibody and streptavidin-proxidase (SP) reagent kit (Fuzhou Maxim Biotech Inc., China).

Experimental methods

Immunohistochemical staining

Specimens were fixed in 10% neutral buffered formalin, embedded in paraffin, cut into 4 μ m sections and stained with hematoxylin and eosin (HE) or SP methods. Known positive sections were used as a positive control and phosphate buffer saline (PBS) was substituted for the primary antibody for a negative control.

Determination of results

The positive protein expression of Wip1 was mostly in nucleus and only a few in cytoplasm. Positive protein expressions of p-p38MAPK, p53, and p16 were found in the nucleus as brown/yellow granules (diamino benzidine (DAB) staining). The numbers of positive cells in five random high-definition fields under the light microscope (\times 400) were then counted, and their proportions relative to the total cells were calculated. Wip1: a stained cell count =0 or <10% as negative (-), 10-30% as (+), and >30% as (++); negative (-) to (+) was regarded as low expression and (++) as high expression. p53, p38, and p16: a stained cell count =0 or <10% as negative (-) and \geq 10% as positive (+).

Statistical analysis

Data were analyzed using SPSS 13.0 software. Rates were compared using Chi square test and correlation was analyzed with Spearman's rank correlation coefficient. $P < 0.05$ was considered significantly different.

Results

Expression of Wip1 in normal thyroid tissues and PTC tissues

The immunohistochemical staining of Wip1 was negative or weak in the follicular epithelial cells of normal thyroid tissues (*Figure 1A*), whereas the positive expression of Wip1 in PTC tissues was mostly in nucleus (*Figure 1B*). The

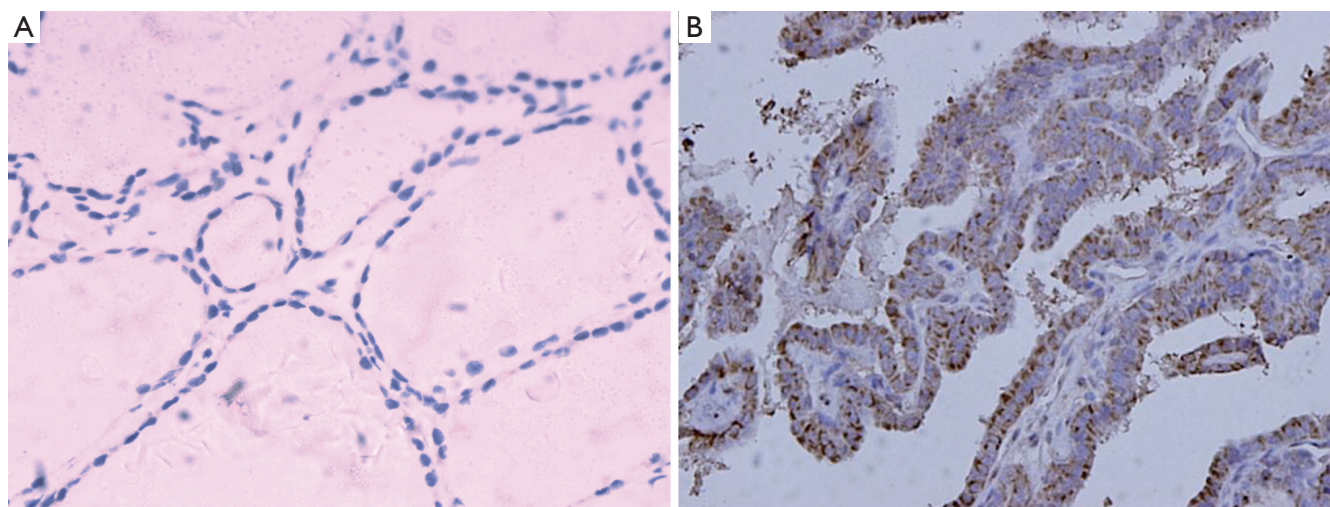


Figure 1 Expression of Wip1 (DAB staining, $\times 400$). A: Normal thyroid tissue, without Wip1 positive expression; B: PTC tissue, with Wip1 positive expression (mostly in nucleus).

high-expression rate of Wip1 protein in the PTC tissue was 64.3% (45/70), which was significantly higher than that in normal thyroid tissue (0/20) ($\chi^2=25.714$, $P=0.00039$).

Wip1 high expression and its relationship with the clinicopathological factors of thyroid carcinoma

Wip1 high expression showed no correlation with age, gender, tumor size, lymph node metastasis, and pathological stage (all $P>0.05$). These factors also showed no significant difference among the groups ($\chi^2=0.311$, 0.029, 0.002, 0.796, and 0.194, respectively; $P=0.577$, 0.865, 0.961, 0.372, and 0.659, respectively) (Table 1).

Protein expressions of p38MAPK, p53, and p16 in PTC tissues their correlations with the Wip1 high expression

The positive protein expressions of p38, p53, and p16 in PTC tissues were all in nucleus (Figure 2). The Wip1 protein high-expression was negatively correlated with p53, p38MAPK, and p16 protein expressions (Table 2).

Discussion

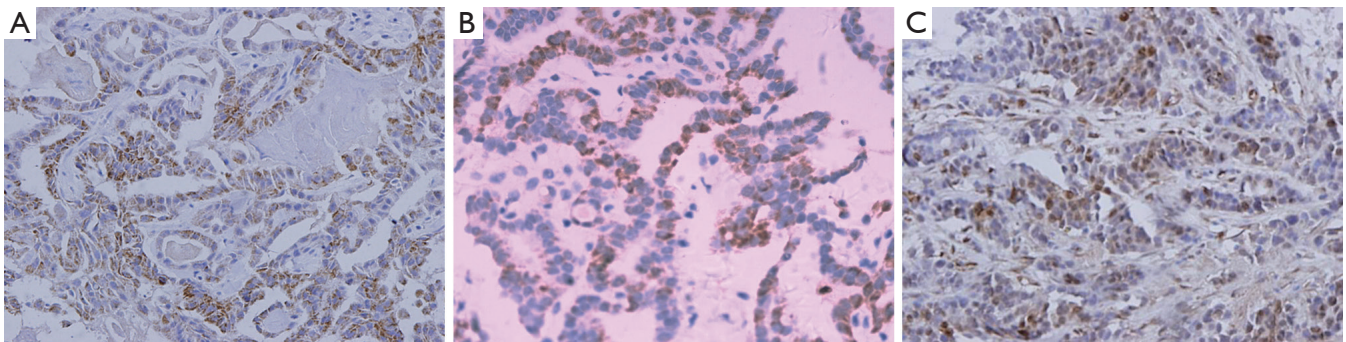
Previous study has shown that the occurrence and development of human tumors are closely associated with the changes in intracellular genome, among which the activation of proto-oncogenes, the inactivation or missing

of anti-oncogenes, and the changes in the functions of many cell cycle regulatory genes play key roles (14). As shown in our current study, the high-expression rate of Wip1 protein in the PTC tissue was 64.3% (45/70), which was significantly higher than that in normal thyroid tissue ($P<0.01$), indicating that Wip1 is involved in the progression of PTC. Correlation analysis showed that Wip1 high expression was not correlated with the clinicopathological factors including age, gender, tumor size, lymph node metastasis, and pathologic stage, which are consistent with our previous findings (15). Therefore, Wip1 can not be used to predict the prognosis.

Also in our study, the high protein expression of Wip1 in PTC tissues was negatively correlated with the protein expressions of p38MAPK, p53, and p16; in other words, PTC tissues with Wip1 high protein expression were often associated with the depletion of the protein expression of p38MAPK, p53, or p16, and vice versa. As a stress regulator, Wip1 can inactivate p38MAPK (6,16). p38MAPK, a key cell cycle regulator, plays an important role in controlling the G1/S and G2/M checkpoints; the inactivation of p38MAPK increases the risk of tumorigenesis (3). p53, as one of the most important anti-oncogenes, plays an important role in the apoptosis induced by DNA damage (17,18). p53 has two subtypes: wild type and mutant type. The subtype detected by immunohistochemical methods is mainly mutant type. As shown in our study, by inhibiting the functions of p53,

Table 1 Wip1 high expression and its relationship with the clinicopathological factors of thyroid carcinoma

Clinopathological factors	n	Wip1 high expression		χ^2	P
		n	(%)		
Age					
<45	45	30	66.7	0.311	0.577
≥45	25	15	60.0		
Gender					
Male	16	10	62.5	0.029	0.865
Female	54	35	64.8		
Tumor size					
>40 mm	11	7	63.6	0.002	0.961
≤40 mm	59	38	64.4		
Lymph node metastasis					
Yes	33	23	69.7	0.796	0.372
No	37	22	59.5		
TNM stage					
Stage I - II	51	32	62.7	0.194	0.659
Stage III - IV	19	13	68.4		

**Figure 2** Immunohistochemical staining of p38MAPK, p53, and p16 in PTC tissues (DAB staining, ×400). A: positive expression of p38MAPK; B: positive expression of p53; and C: positive expression of p16.**Table 2** Correlation of Wipl protein high-expression with p53, p38MAPK, and p16 protein expressions in PTC tissues

Wip1	p38		p53		p16	
	(+)	(-)	(+)	(-)	(+)	(-)
(++)	9	36	14	31	12	33
(-)(+)	21	4	17	8	21	4
r	-0.620		-0.356		-0.550	
P	<0.01		<0.01		<0.01	

Wip1 reduced the selective mutation of p53 gene.

As a key anti-oncogene, p16 can specifically inhibit CDK4/6 and thus prevent the transition of a cell from G1 to S phase. It adjusts the mitosis via negative feedback and prevents the abnormal proliferation of cells. And its inactivation or missing can dramatically accelerate the growth of cells, therefore play a crucial role in the progression of PTC.

In addition, we found that the p16/p38MAPK/p53/Wip1 pathway is abnormal in PTC, and this abnormality may possibly be associated with the aberrantly up-regulated Wip1, which can induce inhibition of p38MAPK, p53 and p16. It is therefore speculated that Wip1 may be a therapeutic target for PTC; particularly, the inhibition of Wip1 function may enhance the activities of anti-oncogenes and thus exert therapeutic effect on PTC. Research also has shown that arsenic trioxide can inhibit Wip1, during which Wip1 is served as the direct molecular target. Currently, arsenic trioxide is mainly used for the treatment of acute promyelocytic leukemia (19). The prevalence of thyroid cancer has shown an increasing trend in China in recent years (20); unfortunately, this disease is not sensitive to conventional chemotherapy drugs.

In conclusions, the p16/p38MAPK/p53/Wip1 pathway is abnormal in PTC, and this abnormality may possibly be associated with the aberrantly up-regulated Wip1, which can induce inhibition of p38MAPK, p53 and p16. The arsenic trioxide may exert its anti-tumor efficacy in the targeted therapy for thyroid cancer.

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