

MAP2K6 is associated with radiation resistance and adverse prognosis for locally advanced nasopharyngeal carcinoma patients

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Background: Although radiotherapy is the primary therapeutic option for nasopharyngeal carcinoma (NPC), local recurrence and distant metastasis caused by radioresistance are still the major barriers for some NPC patients who cannot benefit from radiotherapy. In this study, we analyzed the association between MAP2K6 expression and radioresistance in patients with locally advanced NPC.

Methods: We collected 120 NPC patients who received radiotherapy in the Xiangya Hospital of Central South University from August 2008 to July 2012. The clinical data and tissue samples of patients were collected. Detection of MAP2K6 was performed using immunohistochemical staining.

Results: The rates of two groups were 19.4% and 4.2%, and significant difference was observed between MAP2K6 high expression group and low expression group ($\chi^2=5.817$, $P=0.016$). The Kaplan–Meier analysis suggested a significant difference in the survival rate between two groups ($P<0.05$). The results from multivariate Cox regression indicated that the MAP2K6 was independently related to adverse prognosis in NPC patients (HR=3.40, 95% CI=1.13–10.26, $P=0.030$).

Conclusion: The present study indicated that MAP2K6 was correlated with radioresistance, and the elevated expression of MAP2K6 predicted poor prognosis in NPC patients. MAP2K6 may be a new therapy target for radioresistance of NPC.

Keywords: Mitogen-activated protein kinase 6, progression free-survival, radioresistance

Introduction

Nasopharyngeal carcinoma (NPC) is one of the most prevalent cancers in Southern China and Southeast Asia, which seriously affects the local people's health.¹ Radiotherapy is the prior selection for NPC therapy. In the past 10 years, radiation therapy technology has been developing by leaps and bounds with the intervention of electronic computers. The application of intensity-modulated radiation therapy (IMRT) has made the curative effect of NPC radiotherapy continue to improve.^{2–4} However, there are still some patients who cannot benefit from this therapy. The local recurrence and distant metastasis caused by radioresistance become the main obstacles for the patients with NPC to obtain satisfactory efficacy. Blindly increasing the dose of radiotherapy can only increase the local control rate and reduce the distant metastasis rate, but it will also increase the radiation damage of the surrounding normal tissues and organs such as the brain stem and spinal cord and even result in serious short-term and long-term adverse reactions.⁵ It was reported that 5-year disease-free survival rate of NPC patients with stages I–IVB was 83.3%.⁶ However, ~20% of NPC patients have local recurrence and

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distant metastasis.⁷ Previous studies had preliminarily identified the risk factors and possible mechanisms of NPC, but the specific mechanism of local recurrence and metastasis caused by radiotherapy resistance is not yet clear.⁸ Therefore, it will provide new potential targets for radiotherapy sensitization treatment of NPC via exploring molecular mechanisms and key regulating factors of radiotherapy resistance.

Protein kinases belong to a class of enzymes that catalyze phosphorylation of proteins, and they can transfer gamma-phosphate from adenosine triphosphate (usually ATP) to serine, threonine, or tyrosine residues by phosphorylation. Approximately 2% of eukaryotic genes are protein kinases, and there are >500 kinases in the mammalian genome. Protein kinases, such as receptor tyrosine kinases, cyclin-dependent kinases, and MAP kinases (MAPKs) accounts for the majority of all kinases, and they play crucial roles in cellular metabolism, cell signaling, protein regulation, cell transport, cell division, and many other forms of cancer cell pathways.⁹ Mitogen-activated protein kinase kinase 6 (MAP2K6) is an upstream kinase of the p38/MAPK signal pathway.¹⁰ MAP2K6 is involved in various physiological and pathological processes such as cell growth, development, division, and inflammatory reactions.¹¹ In recent years, some studies have found that MAP2K6 may be associated with the occurrence and progression of tumors. In fact, 20% of human cancers are related to the occurrence of MAPK. Parry et al¹² use immune fluorescence technology to systematically evaluate the MAP2K6 protein expression of different types of human tumor samples. The results suggested that MAP2K6 expression was significantly increased in esophageal cancer, gastric cancer, and colon cancer compared with the control, and the high expression of MAP2K6 was also found in the kidney, intestine, and lung cancer primary tissues. These results showed that MAP2K6 was probably involved in the occurrence and progression of human tumors and could be potentially treated as a new diagnostic or prognostic biomarker for cancers. In this study, we explored the association between MAP2K6 and radioresistance in patients with locally advanced NPC.

Methods

Patients and tissue samples

We retrospectively selected 120 NPC patients from the Department of Radiation, Xiangya Hospital of Central South University, between August 2008 and July 2012. The clinical data and paraffin section samples were collected according to the following criteria: 1) all patients

were confirmed by pathology examination; 2) no distant metastasis and other tumors when patients were confirmed; 3) paraffin-embedded tissue samples were available for immunohistochemistry analysis; and 4) patients did not receive other anticancer treatment before chemoradiotherapy or surgery treatment after radiotherapy. TNM staging referred to American Joint Committee on Cancer (AJCC) staging system for NPC. The follow-up ended on May 31, 2016. The primary outcome was recurrent or metastatic. Progression-free survival (PFS) time was defined from therapy to recurrence and metastasis. All outcomes must be confirmed by clinical examination. NPC patients were divided into radioresistance group and radiosensitive group.¹³ This study was approved by the ethics committee of the Xiangya Hospital Central South University, and all patients provided written informed consent, and this study was conducted in accordance with the Declaration of Helsinki.

All patients received external beam IMRT. IMRT was applied as follows: 68–74 Gy/30–33 fraction to planned targeted volume (PTV) nx, 68–70 Gy/30–33 fraction to PTVnd, 60–64 Gy/30–33 fraction to PTV1, and 50–56 Gy/30–33 fraction to PTV2, and received two periods of induction chemotherapy with docetaxel (65 mg/m²/d₁) plus nedaplatin (NDP; 80 mg/m²/d₁) every 3 weeks, followed by six cycles of concurrent chemotherapy with NDP (40 mg/m²/d₁) every week. Besides, some patients received sodium glycididazole through intravenous 750 mg/m²/fraction, 3–5 fraction/week, within 1 or 2 hours before radiotherapy. Some patients received nimotuzumab (100–200 mg/week) and cetuximab (400 mg/m²/week).

Immunohistochemical staining

Immunostaining detection was conducted under the guidelines of ChemMate kit (Dako Denmark A/S, Glostrup, Denmark) and 3,3'-diaminobenzidine as the chromogen, which has been described in the previous studies.¹⁴ First, paraffin section was dewaxed, sealed by 3% H₂O₂ for 10 minutes, and followed by distilled water wash for 5 minutes in the shake bed. Second, slice was placed in heat-resistant glass containers with sodium citrate buffer, 92°C–95°C, 5 minutes; remove the residue and replenish preheated sodium decanoate buffer; 92°C–95°C, 5 minutes. Third, specimens were incubated overnight at 4°C with the anti-MAP2K6 antibody at a dilution of 1:400 (Cell Signaling Technology, Inc., Beverly, MA, USA). Detection of immunostaining was performed using the ChemMate kit (Dako Denmark A/S)

with 3,3'-diaminobenzidine as the chromogen. The non-immune isotype antibodies replaced the primary antibody as the negative control.

The evaluation of staining was conducted by two experienced pathologists. The pathologists scanned the tissue specimens under the low-power ($\times 40$) and high-power ($\times 400$) magnification. The positive and negative results were according to the staining intensity and scope. The criteria were as follows: sample with no positive cells were negative (–), $<25\%$ of cells: weakly positive (\pm), $25\%–50\%$: positive (+), $>50\%$ of cells: strongly positive (++).¹⁵ We defined the negative or weak positive as the MAP2K6 low expression group and the positive or strong positive as MAP2K6 high expression group.

Statistical analyses

First, we divided NPC patients into radioresistant and radiosensitive groups. The clinical parameters between low and high expression groups were compared by using chi-squared test for the qualitative data (age: ≥ 45 vs <45 years; gender: male vs female; radiotherapy: ≥ 73.27 vs <73.27 Gy; AJCC: stage IVa vs III; chemotherapy: no vs yes; targeted therapy: no vs yes; sensitization therapy: no vs yes) and Student's *t*-test for the quantitative data. Second, NPC patients were divided into low expression group and high expression of MAP2K6. The chi-squared and Student's *t*-test were also used for comparison of clinical feature between two groups. HR with 95% CIs was calculated for assessing the relationship between MAP2K6 and prognosis through univariate and multivariate Cox regression analysis. The survival curves of MAP2K6 low and high expression was plotted by Kaplan–Meier analysis. We performed the whole analyses by using SPSS 20.0 (IBM Corporation, Armonk, NY, USA) and GraphPad Prism 6.0 (GraphPad Software, Inc., La Jolla, CA, USA). $P < 0.05$ was considered to be statistically significant.

Results

General characteristics of NPC patients

One hundred and twenty NPC patients were included in the follow-up, including 16 radioresistance patients and 104 radiation-sensitive patients, including 52 females and 68 males. The median age was 47 years, ranging from 28 to 70 years. The follow-up duration was from 12 to 71.8 months, with the median of 42 months. There were 52 patients younger than 45 years and 68 patients aged 45 years or older. Fifty-two patients were classified as stage III and 68 patients as stage IVa. Eleven patients did not receive chemotherapy. Sixty-six patients receive sensitization therapy before radiotherapy. Fourteen patients took nimotuzumab. Clinical

and histopathological data of NPC patients are presented in Table 1. The MAP2K6 expression level in radioresistant group was significantly higher than that in the radiosensitive group ($P=0.016$). There were no significant differences in age ($P=0.971$), gender ratio ($P=0.263$), radiotherapy dose ($P=0.543$), AJCC stage ($P=0.263$), chemotherapy ($P=0.620$), target therapy ($P=0.428$), and sensitization ($P=0.517$) between two groups.

We also divided NPC patients into MAP2K6 high expression group and MAP2K6 low expression group according to the MAP2K6 expression level. Clinical characteristics were also compared between two groups. The results are listed in Table 2. No significant differences were observed in clinical parameters between two groups ($P > 0.05$).

MAP2K6 and radiotherapy resistance

According to the results of immunohistochemical, MAP2K6 was mainly located in the nucleus and cytosol of cell, and the MAP2K6 in the radioresistant group was significantly stronger than in the radiosensitive group (Figure 1). Fourteen patients were confirmed as radioresistance in the MAP2K6 high expression group, and there were two patients in the MAP2K6 low expression group. The rates of two groups were 19.4% and 4.2%, and significant difference was observed between the high expression group and low expression group ($\chi^2=5.817$, $P=0.016$).

MAP2K6 and survival

The Kaplan–Meier analysis showed a significant difference in the survival curve between the high expression group and low expression group ($P < 0.05$, Figure 2). We conducted the univariate and multivariate Cox regression analyses to identify the relationship between MAP2K6 expression and PFS in the NPC patients, including the following parameters: age, gender, therapy dose, chemotherapy, AJCC stage, targeted therapy, sensitization therapy, and MAP2K6 expression. The univariate results showed that MAP2K6 was associated with adverse prognosis in NPC patients (HR = 3.46, 95% CI = 1.17–10.22, $P=0.025$). The other variables are presented in Table 3. The multivariate results indicated that the MAP2K6 high expression was independently related to adverse prognosis in NPC patients (HR = 3.40, 95% CI = 1.13–10.26, $P=0.030$). Other variables were not associated with PFS in NPC patients (Table 4).

Discussion

The current study qualitatively evaluated the expression of MAP2K6 in NPC tissues via immunohistochemistry, and

Table 1 General characteristics of study patients

Parameters	Radiation sensitivity, n (%)		χ^2	P-value
	Radiation-resistant group	Radiation-sensitive group		
Age (years)			0.001	0.971
<45	7 (13.5)	45 (86.5)		
≥45	9 (13.2)	59 (86.8)		
Gender			1.254	0.263
Female	9 (10.3)	43 (82.7)		
Male	7 (17.3)	61 (89.7)		
Radiotherapy dose			0.371	0.543
<73.27	9 (15.3)	50 (84.7)		
≥73.27	7 (11.5)	54 (88.5)		
AJCC stage			1.254	0.263
III	9 (17.3)	43 (82.7)		
IVa	7 (10.3)	61 (89.7)		
Chemotherapy			0.246	0.620
Yes	14 (12.8)	95 (87.2)		
No	2 (18.2)	9 (81.8)		
Targeted therapy			0.526	0.428
Yes	1 (7.1)	13 (92.9)		
No	15 (14.2)	91 (85.8)		
Sensitization therapy			0.420	0.517
Yes	10 (15.2)	56 (84.8)		
No	6 (11.1)	48 (88.9)		
MAP2K6			5.817	0.016
Low	2 (4.2)	46 (95.8)		
High	14 (19.4)	58 (80.6)		

Abbreviation: AJCC, American Joint Committee on Cancer.

Table 2 Correlation between MAP2K6 expression and clinical pathological features in patients with NPC

Parameters	MAP2K6 expression, n (%)		χ^2	P-value
	Low expression	High expression		
Age (years)			0.684	0.408
<45	23 (44.2)	29 (55.8)		
≥45	25 (36.8)	43 (63.2)		
Gender			2.494	0.114
Female	25 (66.2)	27 (51.9)		
Male	23 (33.8)	45 (48.1)		
Radiotherapy dose			1.801	0.179
<73.27	20 (33.9)	39 (66.1)		
≥73.27	28 (45.9)	33 (54.1)		
AJCC stage			2.042	0.153
III	17 (32.7)	35 (67.3)		
IVa	31 (45.6)	37 (54.4)		
Chemotherapy			1.505	0.220
Yes	46 (42.2)	63 (57.8)		
No	2 (18.2)	9 (81.8)		
Targeted therapy			0.054	0.816
Yes	1 (7.1)	13 (92.9)		
No	15 (14.2)	91 (85.8)		
Sensitization therapy			0.275	0.600
Yes	25 (37.9)	41 (62.1)		
No	23 (42.6)	31 (57.4)		
Therapy			5.817	0.016
Sensitivity	46 (95.8)	58 (80.6)		
Resistance	2 (4.2)	14 (19.4)		

Abbreviations: AJCC, American Joint Committee on Cancer; NPC, nasopharyngeal carcinoma.

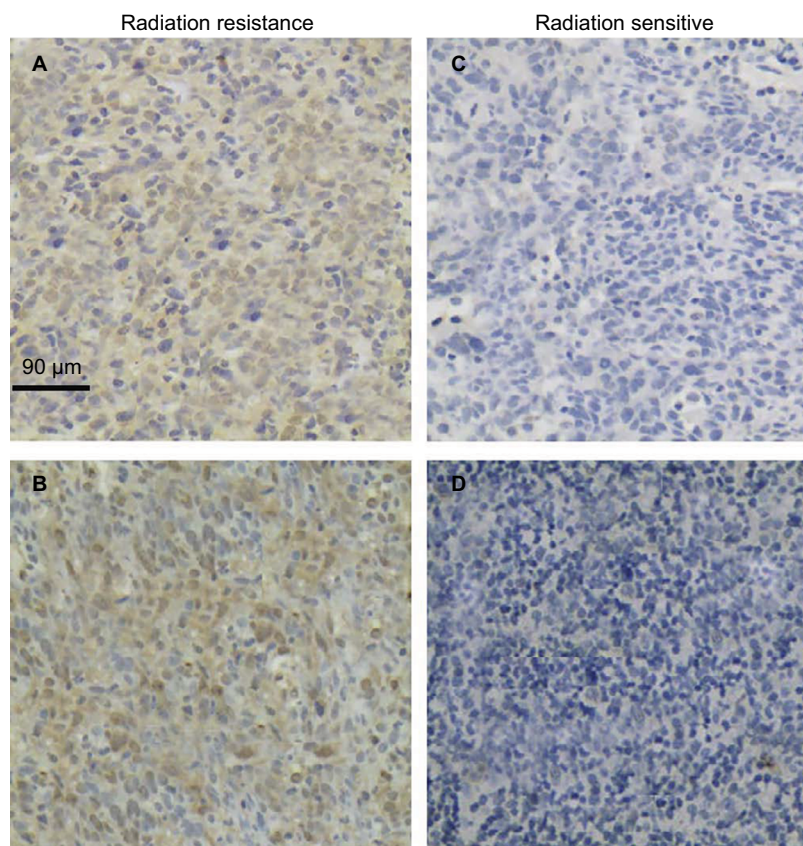


Figure 1 Expression of MAP2K6 in radiation-resistant (upper) and radiation sensitive (lower) NPC tissues – radiation resistant: (A) 60% and (B) 0%; radiation sensitive: (C) 5% and (D) 10%.

Abbreviation: NPC, nasopharyngeal carcinoma.

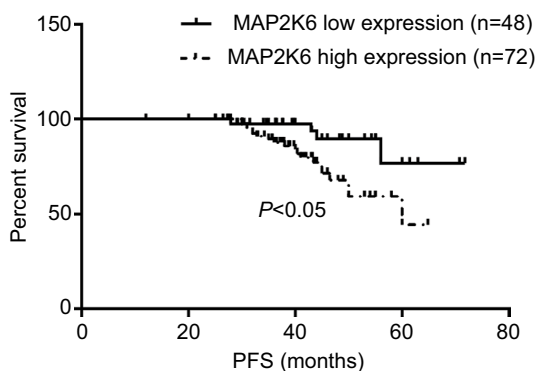


Figure 2 The significant difference in the PFS rate was observed in the patients with low MAP2K6 expression and high MAP2K6 expression.

Abbreviation: PFS, progression-free survival.

our results indicated that the expression of MAP2K6 was significantly elevated in the radioresistance NPC tissues. The further analyses indicated that the MAP2K6 expression was associated with the prognosis of NPC patients, and the high expression of MAP2K6 predicted a worse prognosis of patients with NPC.

Although radiotherapy is the primary therapeutic option for NPC, local recurrence and distant metastasis caused by radioresistance are still the major barriers for some NPC patients who cannot benefit from radiotherapy. Therefore, it is significantly important to deeply explore the molecular mechanisms and to identify the key effect factor and related signaling pathways of NPC radioresistance for intervention. Previous studies had reported some radioresistance biomarkers.^{16–18}

MAP2K6 belongs to one of the families of dual specificity protein kinase and can activate the MAPK. The MAPK can regulate the extracellular signal pathway and is the overlap points of several signals.¹⁹ MAP2K6 can phosphorylate and activate the p38/MAPK signal pathway under the status of inflammatory and stress. Previous studies have suggested that MAP2K6 plays important roles in cell cycle regulation, transcription, and cell apoptosis.^{20,21} Parry et al²² found that the expression of MAP2K6 was significantly higher in esophageal, stomach, and colon cancers than controls using Western blotting and immunofluorescence assay. Overexpression of MAP2K6 is indicative of its role in human cancers and

Table 3 Univariate Cox regression analyses of clinical prognosis in patients with NPC

Factors	β	SE	Wald χ^2	HR	95% CI	P-value
Age (≥ 45 vs 45 years)	-0.250	0.439	0.325	0.78	0.33–1.84	0.569
Gender (male vs female)	0.295	0.438	0.453	1.34	0.57–3.17	0.501
Radiotherapy dose (≥ 73.27 vs <73.27)	-0.525	0.451	1.356	0.59	0.24–1.43	0.244
AJCC stage (IVa vs III)	-0.529	0.438	1.459	0.59	0.25–1.39	0.227
Chemotherapy (no vs yes)	0.919	1.026	0.801	2.51	0.33–18.72	0.371
Targeted therapy (no vs yes)	-0.203	0.745	0.074	0.82	0.19–3.52	0.785
Sensitization therapy (no vs yes)	-0.194	0.446	0.189	0.82	0.34–1.98	0.664
MAP2K6 (high vs low)	1.240	0.553	5.020	3.46	1.17–10.22	0.025

Abbreviations: AJCC, American Joint Committee on Cancer; NPC, nasopharyngeal carcinoma.

Table 4 Multivariate Cox regression analyses of clinical prognosis in patients with NPC

Factors	β	SE	Wald χ^2	HR	95% CI	P-value
Age (≥ 45 vs 45 years)	-0.226	0.444	0.358	0.77	0.32–1.83	0.550
Gender (male vs female)	0.222	0.426	0.272	1.25	0.54–2.88	0.602
Radiotherapy dose (≥ 73.27 vs <73.27)	-0.655	0.458	2.043	0.52	0.21–1.28	0.519
AJCC stage (IVa vs III)	-0.217	0.446	0.237	0.81	0.34–1.93	0.626
Chemotherapy (no vs yes)	1.189	1.032	1.327	3.28	0.43–24.84	0.249
Targeted therapy (no vs yes)	0.118	0.643	0.034	1.13	0.32–3.97	0.854
Sensitization therapy (no vs yes)	0.006	0.455	0.000	1.01	0.41–2.45	0.990
MAP2K6 (high vs low)	1.224	0.564	4.714	3.40	1.13–10.26	0.030

Abbreviations: AJCC, American Joint Committee on Cancer; NPC, nasopharyngeal carcinoma.

could possibly be a novel diagnostic or prognostic biomarker in these cancers. It was reported that the fusion protein TAT-OSBP-MAP2K6 (E) can induce the autophagic cell death by activating the P38MAPK signaling pathway, thereby inhibiting the proliferation, invasion, and metastasis of ovarian cancer cisplatin-resistant SKOV3/DDP cells cisplatin sensitivity. To a certain extent, it can reverse cisplatin resistance. It is suggested that the fusion protein TAT-OSBP-MAP2K6 (E) is expected to be a new chemotherapeutic agent in the treatment of cisplatin in ovarian cancer and provide a new therapeutic mode for the treatment of ovarian cancer.^{23,24} Galan-Moya et al²⁵ reported that the balance between MKK6 and MKK3 mediated p38/MAPK-associated resistance to cisplatin in non-small-cell lung cancer cell lines, and these results were also confirmed in head and neck carcinoma-derived cell line. Studies about the role of MAP2K6 are few in patients with NPC. In fact, we simply compared the expression level of MAP2K6 between normal tissue and NPC tissue using some database from Gene Expression Omnibus (GEO), and no significant difference was observed. However, it was suggested that phosphorylation in the Ser207 and Thre211 can activate the MAP2K under the stress stimulations such as radiation, inflammatory reaction, osmotic shock, and hypoxia.²⁶ It is possible that MAP2K6 can be activated by radioactive X-rays.

Protein kinases such as MAP2K6 are important components of MAPK signal pathway. The dual specificity protein

kinases consist of the following kinases. MAP3K3 mainly catalyzes the phosphorylation of threonine and tyrosine of MAPK such as p38, MAPK11, and so on, which regulated the cellular responses to inflammation factors and types of stress, while MAP2K6 is the activator of MAPK during the process of response.²⁷ MAPK consisted of three types, c-Jun amino-terminal kinase, p38, and extracellular signal-regulated protein kinase. They can be activated by phosphorylation in Tyr and Ser, which increased the expression of some proliferation gene. Schiemann et al²⁸ first reported that leukemia inhibitory factor (LIF) receptor was linked to MAPK. An obvious dose–response relationship was observed when 3T30L1 (cell extracts) processed by LIF in vitro was bound to the intracellular domain of LIF receptor. The activity of MAPK increased with LIF concentration. This result indicated that MAPK was correlated with LIF phosphorylation. MAPK was activated and combined to the Ser1044 area of intracellular domain of LIF receptor. Further study found that LIFR could be a substrate of MAPK. The phosphorylation of serine location accelerated the procession of itself intracellular phagocytosis. In other words, the activity of MAPK quantitatively regulated the LIF receptor in the surface membranes of cells. Recent study reported that serum LIF was associated with radiosensitivity in NPC patients.²⁹ In this study, the author first found that LIF significantly promoted the NPC tumor growth and enhanced radioresistance in vitro and vivo. LIF

treatment can activate mTORC1/p70S6K signaling pathway and enhanced tumor growth and radioresistance. Treatment with LIF receptor or the mTOR inhibitor rapamycin reversed the LIF-mediated effect. LIF-mediated effect, which is the growth of NPC tumor was in arrest and the sensitivity to irradiation was increased. We assumed that this is the mechanism of MAP2K6-mediated radioresistance in NPC tumor. This speculation gives us an important clue. Of course, further research was required.

Some study limitations should be addressed. The present study has a relatively small sample size, and larger sample size is needed to validate the results. Moreover, the present study did not explore the molecular mechanism, and experiments in vivo and vitro should be performed. Besides the adjusted factors are limited, some potential factors may affect the results. Finally, few patients received operation treatment. The adjacent tissues were unavailable for now. We did not compare the expression levels of MAP2K6 in the tissue. According to the GEO data set (GSE53819 and GSE12452), expression of MAP2K6 cannot be confirmed. But it is reported that MAP2K6 was involved in the signaling pathway of NPC, including stress-induced cell cycle arrest, transcription activation, and apoptosis. This can explain the potential role of MAP2K6 in the NPC radiotherapy.

Taken together, the present results indicated that MAP2K6 was correlated with radioresistance in NPC patients, and the elevated expression levels of MAP2K6 predicted poor prognosis. MAP2K6 may be a valuable biomarker for predicting tumor prognosis and a new therapy target. Future study should focus on the molecular mechanism, especially in the LIF/mTOR/p70S6K signal pathway.

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Disclosure

The authors report no conflicts of interest in this work.

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