

Up-regulation of peroxiredoxin 3 by high-risk human papillomavirus in cervical cancer cells

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Abstract

Introduction: Peroxiredoxin 3 (PRX3) is a member of the PRX family with antioxidant functions by scavenging hydrogen peroxide. Since the development of cervical cancer is causally linked to high-risk human papillomavirus (HPV) that induces oxidative stress, we conducted the present study to investigate the response of PRX3 to high-risk HPV infection.

Material and methods: This study included fifty-six patients with invasive squamous cervical cancer and sixty control patients with hysteromyoma. Enzyme-linked immunosorbent assay was performed to detect cervical oxidative stress and serum PRX3. The expression of PRX3 and oncoprotein E6 of HPV16 or HPV18 was examined in cervical cancer tissues by immunohistochemistry. Western blot was applied to detect the expression of PRX3 and E6 in cervical cancer cell lines including CaSki, HeLa, and C33A.

Results: Patients with cervical cancer showed higher serum PRX3 than control patients with hysteromyoma. Levels of oxidative markers in cervical cancer tissues were elevated as compared to normal cervical epithelia. PRX3 expression was upregulated in cervical cancer tissues and the upregulation was positively associated with the expression of E6 of HPV16 or HPV18. The association was confirmed in HPV-containing cervical cancer cell lines including CaSki and HeLa.

Conclusions: Our results indicated a positive response of PRX3 to HPV-induced oxidative stress. Serum PRX3 might be a potential indicator of active amplification of high-risk HPV in cervical cancer cells.

Key words: cervical cancer, human papillomavirus, oxidative stress, peroxiredoxin, reactive oxygen species.

Introduction

As a kind of antioxidant enzyme, peroxiredoxin 3 (PRX3) is mainly located in mitochondria and reduces hydrogen peroxide into water with its two cysteines. By controlling mitochondrial reactive oxygen species (ROS), PRX3 is involved in diverse cellular processes such as growth, differentiation, apoptosis, and malignancy [1–3]. According to previous reports, the expression of PRX3 was upregulated in cervical cancer [4, 5]. Strong cytoplasmic staining of PRX3 was observed in the surrounded cells of cervical cancer nests, which was consistent with the pattern of Ki67 as a marker of active cell proliferation [5]. However, the precise mechanism for the upregulation of PRX3 is not well known.

The development of cervical cancer is causally linked to high-risk human papillomavirus (HR-HPV), and the integration of HPV oncogenes (e.g. E6 and E7) into the genome of host cells is considered to be a key step towards carcinogenesis. Previous studies indicated that oxidative stress was involved in the integration process. The oncogenes of HR-HPV induced ROS production and subsequent DNA damage of host cells, promoting genomic instability of the cells and the integration of HPV DNA [6, 7]. In addition, an increase in oxidative DNA damage facilitated HPV genome amplification in cancer cells [8]. Since alteration of PRX3 was involved in the persistent HPV infection and cervical carcinogenesis [9], the present study was conducted to investigate the link between PRX3 and HR-HPV in cervical cancer.

Material and methods

Patients and samples

This study was approved by the Ethics Committee of Yantai Affiliated Hospital of Binzhou Medical University (No. 2017-007R) and was conducted between October 2017 and December 2019. After informed consent from all patients, we collected blood samples from patients with squamous cervical cancer 3 days before and after the operation. Samples from patients with hysteromyoma were collected at the same time and were used as controls. General information including age, body mass index (BMI), and blood pressure (BP) was recorded before blood sampling. Fasting blood sugar (FBS), fasting insulin (FINS), total cholesterol (TC), and triglyceride (TG) were routinely examined in the blood.

Examination of HPV infection in cervixes

The status of HPV infection in cervical tissues was examined using a HPV genotyping test kit (HybridBio Company, Hong Kong, China). Briefly, DNA was extracted from cervical tissues and amplified by PCR. The PCR products were then hybridized on a HPV genotyping membrane that contained probes corresponding to 23 HPV subtypes (HR-HPVs: 16, 18, 31, 33, 35, 39, 45, 51, 52, 53, 56, 58, 59, 66, 68, 73, 82; low-risk HPVs: 6, 11, 42, 43, 81, 83). Biotin was used as a positive control and distilled water as a negative control. The membrane was visualized through NBT/BCIP to determine the status of HPV infection.

Detection of oxidative markers in cervixes

Oxidative markers including oxidative DNA adduct 8-hydroxy-2-deoxyguanosine (8-OHdG) and glycoxidation product carboxymethyl lysine

(CML) were detected in cervical cancer tissues and control samples by enzyme-linked immunosorbent assay (ELISA). The Highly Sensitive 8OHdG ELISA kit was purchased from Japan Institute for the Control of Aging (Fukuroi 437-0122, Japan), and the Human CML ELISA kit was purchased from Kamiya Biomedical Company (Seattle, WA 98168 USA). For 8OHdG detection, DNA was extracted from tissues and digested in distilled water. 50 μ l of the sample or standard and 50 μ l of the primary antibody were added to each well. After fully mixing, the plates were covered and incubated at 4°C overnight. The contents of the plates were poured off and the plates were washed thoroughly. 100 μ l of secondary antibody was added to each well and incubated at room temperature for 1 h. The contents of the plates were poured off and the plates were washed thoroughly. 100 μ l of the reconstituted enzyme substrate was added to each well and incubated at room temperature for 15 min. Finally, 100 μ l of the reaction terminating solution was added and the absorbance was read at 450 nm. The amount of 8OHdG in each sample was determined using a standard curve by plotting absorbance vs. concentration of standards. For CML detection, tissues were rinsed in ice-cold PBS to remove excess blood and weighed before homogenization. The homogenates were centrifuged for 15 min at 1500 \times g and the supernatants were used for the assay. 100 μ l of calibrators or samples were added to each well of the antibody pre-coated plate, and 100 μ l of PBS was added to the blank control well. 10 μ l of balance solution was dispensed into 100 μ l specimens and mixed thoroughly. 50 μ l of conjugate was added to each well (except the blank control well) and incubated at 37°C for 1 h. After removal of incubation mixture and washing five times, 50 μ l Substrate A and 50 μ l of Substrate B were added to each well. After incubation at 37°C for 15 min, 50 μ l of stop solution was added to each well and the absorbance was read at 450 nm. A calibration curve was constructed by plotting the average absorbance for each calibrator on the horizontal (X) axis against the concentration on the vertical (Y) axis, and the concentration of samples was calculated from the calibration curve.

Detection of serum PRX3

Blood samples from the subjects were centrifuged at 1500 \times g and 4°C for 10 min for serum separation. We performed ELISA to detect serum PRX3 by following the manufacturer's instructions (NeoScientific Company, Cambridge, MA, USA). Except for the pre-treatment of samples, the detection procedure was basically the same as described in the above section. Each sample was

assayed in triplicate and measured at the absorbance of 450 nm on a spectrophotometer.

Immunohistochemistry

The cervical tissues were processed to be formalin-fixed and paraffin-embedded blocks. Slides were incubated with mouse monoclonal antibodies against human PRX3 and E6 of HPV16 or HPV18 respectively (1 : 1000 dilution according to the suggestion of the manufacturer Abcam, Cambridge, UK). Immunostaining scores were evaluated by two pathologists according to the product of positive cell percentage and staining intensity: < 5% – 0, 5–20% – 1, 21–50% – 2, 51–75% – 3, > 75% – 4; no color – 0, light yellow – 1, yellow – 2, deep yellow – 3.

Detection of PRX3 and E6 expression in cervical cancer cell lines

Cervical cancer cell lines including CaSki, HeLa, and C33A were purchased from the Cell Bank of Chinese Academy of Sciences (Shanghai, China). These cells were cultured at 37°C in a humidified incubator with 5% CO₂ and were passaged at a 1 : 4 split ratio. Cells at passage 3 were re-cultured in six-well plates with a density of 1 × 10⁶ cells/ml. Seventy-two hours later, total protein was extracted from the cells to detect the expression of PRX3 and E6 of HPV16 or HPV18 respectively by Western blot (WB). The primary antibodies against PRX3 and E6 were purchased from Abcam, Cambridge, UK. GAPDH was used as the internal control. Cells from each well of the culture plate were considered as one sample and we detected twelve samples for each cell line. The intensities of the signals were analyzed with a Gel-Pro Analyzer and presented as integrated optical density (IOD).

Statistical analysis

Serum PRX3 was compared by *t* test. Analysis of variance was used to compare the expression of PRX3 and E6 in cervical cancer cell lines, while Pearson’s correlation coefficient was applied to analyze the association between the expression of PRX3 and E6. *P* < 0.05 was considered to be statistically significant.

Results

In the present study, we included fifty-six patients with invasive squamous cervical cancer and sixty control subjects with hysteromyoma. The FIGO stages of the cancer patients were 39 for stage I and 17 for stage II, among which 10 cases showed cancer positivity in pelvic lymph nodes. The patients did not receive chemotherapy or radiotherapy before the operation. As indicated in Table I, there was no significant difference between cancer patients and controls in relation to age, BMI, BP, FBS, FINS, TC, and TG.

Status of HPV infection and oxidative stress in cervical cancer patients

As determined by HPV genotyping test kit, all the tumor samples were infected with HR-HPVs, among which forty-eight samples were positive for HPV16, four samples were positive for HPV18, and four samples were positive for HPV33. No relationship was found between HPV infection and BMI. All of the control patients were negative for HPVs. The levels of 8-OHdG were 7.5 ± 1.2 ng/ml in cancer tissues and 2.9 ± 0.3 ng/ml in controls (cancer vs. control: *t* = 8.20, *p* < 0.001), while the levels of CML were 116.9 ± 5.7 µg/ml in cancer tissues and 52.5 ± 4.3 µg/ml in controls (cancer vs. control: *t* = 10.89, *p* < 0.001).

Serum levels of PRX3 in cervical cancer patients

We performed ELISA to detect the levels of serum PRX3 in cervical cancer patients and controls with hysteromyoma. Pre-operative serum PRX3 was 30.2 ± 4.6 ng/ml in cervical cancer patients and 14.6 ± 2.1 ng/ml in controls (cancer patients vs. controls: *t* = 2.320, *p* = 0.026). Further analysis showed higher serum PRX3 in node positive patients than in negative cases (*t* = 2.110, *p* = 0.035). Interestingly, serum levels of PRX3 in cancer patients decreased to 17.0 ± 1.3 ng/ml 3 days after the operation, which was comparable to that of control patients (15.3 ± 1.1 ng/ml). In addition, we did not note a significant association between serum PRX3 and clinical or biochemical parameters including BMI, BP, FBS, FINS, TC, and TG (data not shown).

Table I. Comparison of clinical and biochemical data between cervical cancer patients and controls

Parameter	n	Age [years]	BMI	BP [mm Hg]		FBS [mmol/l]	FINS [mU/l]	TC [mmol/l]	TG [mmol/l]
				SBP	DBP				
CC	56	47.6 ± 5.8	22.4 ± 3.1	119 ± 7.5	83 ± 8.0	4.8 ± 1.5	10.2 ± 3.3	4.8 ± 0.9	1.0 ± 0.8
Control	60	46.8 ± 3.5	22.7 ± 3.7	121 ± 5.4	81 ± 7.8	5.1 ± 1.5	10.7 ± 3.6	5.2 ± 0.9	1.2 ± 0.8
<i>t</i>		1.515	1.436	1.401	1.319	0.185	1.510	1.445	1.850
<i>P</i> -value		0.136	0.172	0.167	0.193	0.853	0.131	0.155	0.064

BMI – body mass index, BP – blood pressure, CC – cervical cancer, DBP – diastolic blood pressure, FBS – fasting blood sugar, FINS – fasting insulin, SBP – systolic blood pressure, TC – total cholesterol, TG – triglyceride.

Expression of PRX3 and E6 of HPV16 or HPV18 in cervical cancer tissues

As shown in Figures 1 A, B, PRX3 was extensively expressed in the cytoplasm of normal cervical epithelia, but the staining intensity was much weaker as compared to that of cervical cancer cells. The immunostaining score was 3.6 ± 0.9 in normal cervical epithelia and 10.6 ± 1.4 in cancer cells ($F = 2.819$, $p = 0.000$). In addition, the E6 of HPV16 or HPV18 was expressed in nearly all cervical cancer cells as illustrated in Figures 1 D, F

respectively. The staining intensity for HPV18 E6 was slightly weaker than that of HPV16 E6, but the immunostaining scores did not reach a significant difference between the two subtypes. Pearson's correlation analysis showed a positive association between the expression of PRX3 and HPV16 E6 ($r = 0.532$, $p = 0.031$) or HPV18 E6 ($r = 0.515$, $p = 0.041$). The staining scores of either PRX3 or HPV16 E6 were not associated with clinicopathologic parameters including stage, cell grade, and lymph nodes (Table II).

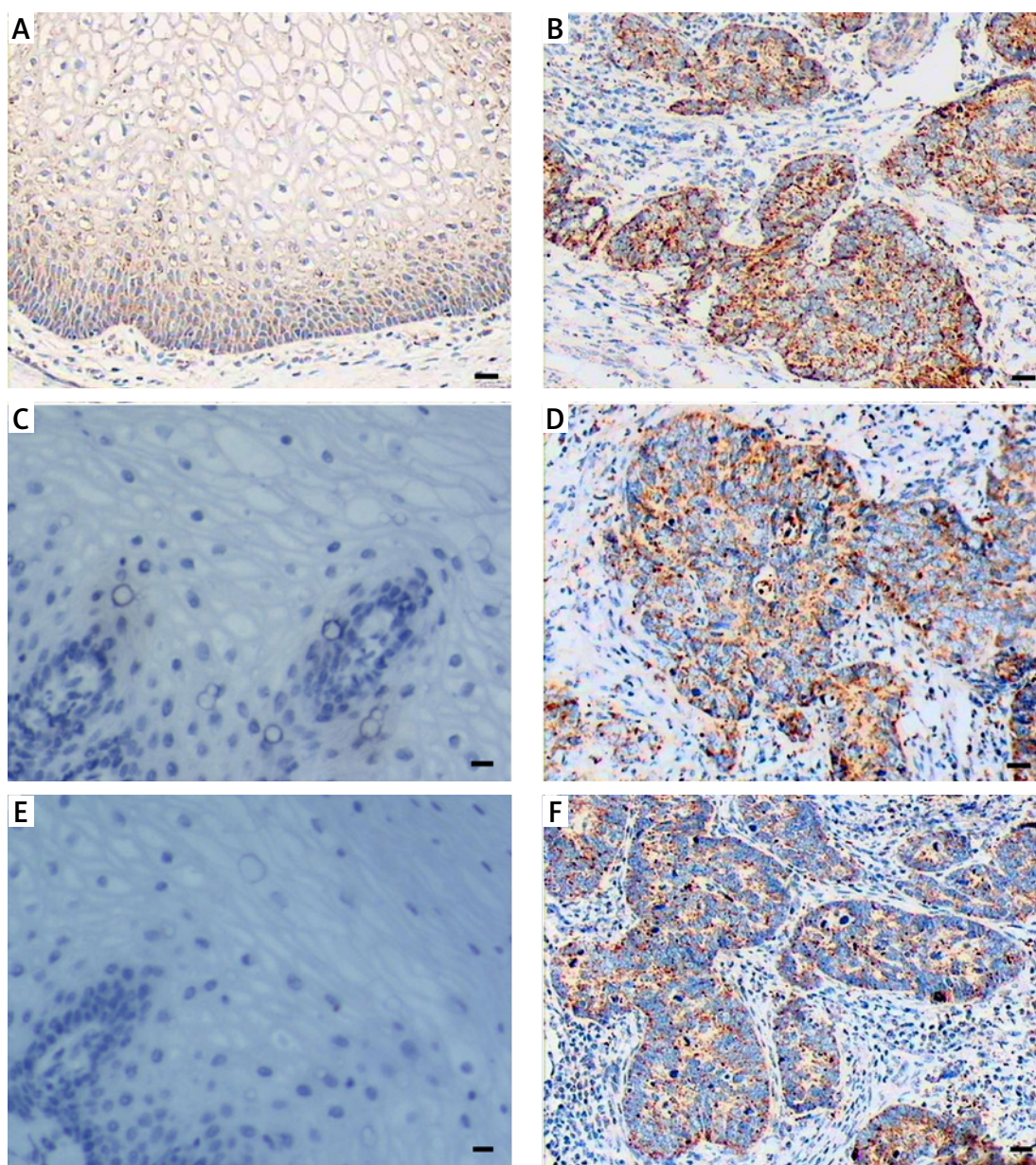


Figure 1. Representative results of immunohistochemistry for PRX3 and E6 of HPV16 or HPV18. **A** – PRX3 was extensively expressed in normal cervical epithelia. **B** – Intensive staining was observed in cervical cancer cells. **C** – Negative control of HPV16 E6 staining in normal cervical epithelia. **D** – The E6 of HPV16 was expressed in nearly all cervical cancer cells. **E** – Negative control of HPV18 E6 staining in normal cervical epithelia. **F** – The E6 of HPV18 was expressed in nearly all cervical cancer cells but the intensity was weaker than that of HPV16 E6. Scale bar = 50 μ m; original magnification $\times 200$

Expression of PRX3 and E6 of HPV16/HPV18 in cervical cancer cell lines

To confirm the association between HPV amplification and PRX3 expression, we detected the expression of PRX3 and E6 of HPV16/HPV18 in cervical cancer cell lines, including CaSki, HeLa, and C33A. As indicated in Figures 2 A, B, the expression of PRX3 was significantly increased in HR-HPV-containing cells (CaSki and HeLa) as compared to that in HPV-negative C33A cells. Figure 2 C shows the mean IOD values of PRX3 and HPV16/HPV18 E6 in each cancer cell line. In addition, the expression of PRX3 was positively associated with that of E6 of HPV16 ($r = 0.651, p = 0.027$) or HPV18 ($r = 0.560, p = 0.031$).

Discussion

In the present study, we demonstrated upregulation of PRX3 by HR-HPV in cervical cancer cells, suggesting an important role of PRX3 in the re-

sponse to HR-HPV induced oxidative stress. Since the serum PRX3 was increased in cervical cancer patients, we suppose that serum PRX3 might be a potential indicator of HR-HPV amplification.

The production of ROS is increased because of active proliferation of cervical cancer cells [10]. Another factor that contributes to the ROS production might be the amplification of HR-HPV [11], which induces oxidative stress, as indicated in the present study. Increased ROS production and development of oxidative stress lead to oxidation of proteins and peroxidation of membrane lipids [12]. Based on a review of previous literature, we suppose that the upregulation of PRX3 might be transcriptional regulation by genes including c-Myc, nuclear factor E2-related factor 2 (Nrf2), forkhead box transcription factor FOXO3A or FOXM1, and small non-coding microRNAs (miRNAs). According to previous reports, infection with HPV16 is closely associated with c-Myc amplification [13, 14], which plays an important role in

Table II. Association of staining scores of PRX3 or HPV16 E6 with clinicopathologic parameters

Parameter	Stages		Cell grades			Lymph nodes	
	I	II	I	II	III	Positive	Negative
Cases	39	17	13	15	28	10	46
PRX3 (n = 56)	9.8 ± 1.4	11.4 ± 1.4	12.2 ± 1.6	10.5 ± 1.5	9.5 ± 1.2	12.3 ± 1.6	8.9 ± 1.2
r	0.247	0.207	-0.417	-0.352	-0.383	0.32	0.313
P-value	0.224	0.292	0.06	0.084	0.058	0.104	0.146
HPV16 E6 (n = 48)	8.5 ± 1.1	9.6 ± 1.5	8.8 ± 1.5	9.1 ± 1.1	9.3 ± 1.3	9.6 ± 1.4	8.6 ± 1.2
r	0.110	0.170	-0.325	-0.372	0.393	0.296	0.192
P-value	0.593	0.415	0.122	0.538	0.071	0.142	0.339

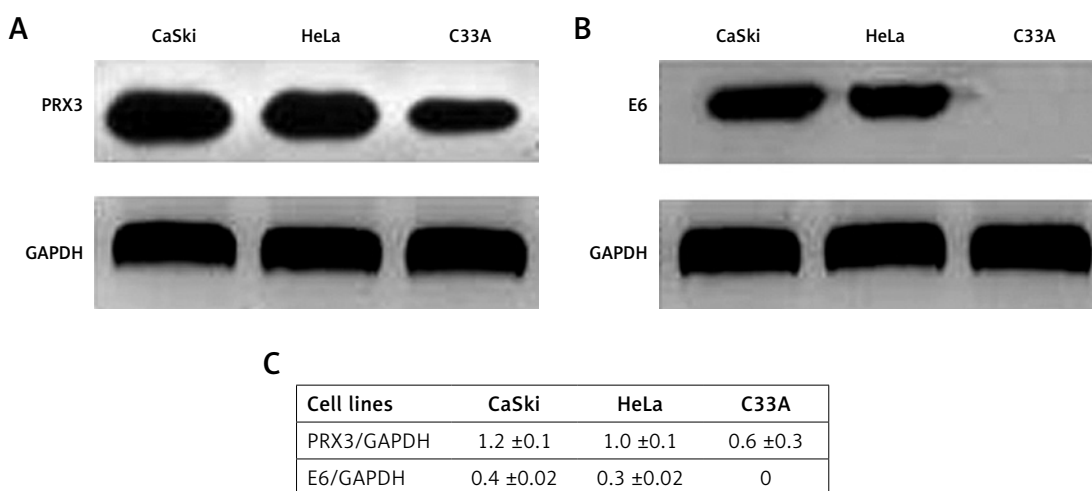


Figure 2. Expression of PRX3 and E6 as determined in cervical cancer cell lines. **A** – WB analysis for PRX3 indicated that the signals of CaSki and HeLa cells were stronger as compared to that of C33A cells. **B** – WB analysis for E6 of HPV16 and HPV18 in CaSki and HeLa cell lines, respectively. **C** – Analysis of variance showed higher expression of PRX3 in CaSki and HeLa cells than in C33A cells. CaSki vs. C33A: $F = 3.480, p = 0.002$; HeLa vs. C33A: $F = 2.901, p = 0.006$

cervical cancer development and progression [15]. Since PRX3 is a downstream target of c-Myc [3], the interaction of c-Myc and PRX3 might be a candidate pathway for the upregulation of PRX3. In lung squamous cell carcinoma, PRX3 is overexpressed in an Nrf2 dependent manner, which indicates a potentially important role of the Nrf2-PRX3 pathway in the tumor [16]. Chiribau *et al.* reported that FOXO3A mediated PRX3 expression, resulting in resistance to oxidative stress in human cardiac fibroblasts [17], which was confirmed by other researchers in pheochromocytoma cells, cardiac cells, and endothelial cells [18–20]. In addition, a study conducted by Cunniff *et al.* reported the co-existence of cytoplasmic FOXM1 and mitochondrial PRX3 in mesothelioma cells [21]. In endometrial cancer stem cells, increased expression of PRX3 was induced by FOXM1 for the maintenance of mitochondrial function and cell survival [22]. MicroRNAs (miRNAs) are a class of small non-protein-coding molecules. Up to now, miR-23b and miR-383 have been reported to control cell growth and cancer progression by negatively regulating PRX3 expression in several types of solid tumors [23, 24].

The serum levels of PRX3 in cancer patients have been investigated in recent years. For example, serum PRX3 is significantly higher in patients with hepatocellular carcinoma and could be used as a biomarker for the diagnosis and prognosis of the disease [25, 26]. However, the alteration of serum PRX3 in cervical cancer patients has not been reported up to now. Here we presented an increase of serum PRX3 in cervical cancer patients, especially in patients with cancer-positive lymph nodes. The serum PRX3 had no significant association with either the staining scores of PRX3 in cancer tissues or clinicopathologic features of patients, unlike in patients with hepatocellular carcinoma [25, 26]. The biological characteristics of cervical cancer are quite different from those of hepatocellular carcinoma. Cancer cells in the cervix grow locally and invade mainly the surrounding tissues or organs. Patients in the present study were at stages I and II, in which distant metastasis was absent. In contrast, the liver is an organ with abundant blood supply and blood-based metastasis is a common occurrence in hepatocellular carcinoma. Therefore, the oxidative stress in liver cancer can be reflected in circulating blood. In the present study, we did not find a significant association between serum PRX3 and biochemical parameters such as FBS and FINS. The present results were consistent with our previous reports that PRX3 responded mainly to oxidative stress induced by a rapid increase of glucose and insulin [27, 28].

In conclusion, we demonstrated the link between PRX3 and HR-HPV in cervical cancer cells. Upregulation of PRX3 reflects a positive response of PRX3 to the oxidative stress in cervical cancer

cells induced by HR-HPV amplification and cell growth. Further studies are needed to elucidate the underlying mechanism for the interaction between PRX3 and HR-HPV.

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Hou-Li Liu and Xiao-Juan Sun contributed equally to this work.

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Ethical approval

This study was approved by the Ethics Committee of Yantai Affiliated Hospital of Binzhou Medical University. The reference number is 2017-007R and the date of approval is 2017-9-26. Informed consent was obtained from all individual participants included in the study.

Conflict of interest

The authors declare no conflicts of interest.

References

1. Matsushima S, Ide T, Yamato M, et al. Overexpression of mitochondrial peroxiredoxin-3 prevents left ventricular remodeling and failure after myocardial infarction in mice. *Circulation* 2006; 113: 1779-86.
2. Chua PJ, Lee EH, Yu Y, et al. Silencing the peroxiredoxin III gene inhibits cell proliferation in breast cancer. *Int J Oncol* 2010; 36: 359-64.
3. Wonsey DR, Zeller KI, Dang CV. The c-Myc target gene PRDX3 is required for mitochondrial homeostasis and neoplastic transformation. *Proc Natl Acad Sci USA* 2002; 99: 6649-54.
4. Kim K, Yu M, Han S, et al. Expression of human peroxiredoxin isoforms in response to cervical carcinogenesis. *Oncol Rep* 2009; 21: 1391-6.
5. Hu JX, Gao Q, Li L. Peroxiredoxin 3 is a novel marker for cell proliferation in cervical cancer. *Biomed Rep* 2013; 1: 228-30.
6. Williams VM, Filippova M, Filippov V, Payne KJ, Duerksen-Hughes P. Human papillomavirus type 16 E6* induces oxidative stress and DNA damage. *J Virol* 2014; 88: 6751-61.
7. Chen Wongworawat Y, Filippova M, Williams VM, Filippov V, Duerksen-Hughes PJ. Chronic oxidative stress increases the integration frequency of foreign DNA and human papillomavirus 16 in human keratinocytes. *Am J Cancer Res* 2016; 6: 764-80.
8. Marullo R, Werner E, Zhang H, Chen GZ, Shin DM, Doetsch PW. HPV16 E6 and E7 proteins induce a chronic oxidative stress response via NOX2 that causes genomic instability and increased susceptibility to DNA damage in head and neck cancer cells. *Carcinogenesis* 2015; 36: 1397-406.
9. Safaiean M, Hildesheim A, Gonzalez P, et al. Single nucleotide polymorphisms in the PRDX3 and RPS19 and

- risk of HPV persistence and cervical precancer/cancer. *PLoS One* 2012; 7: e33619.
10. Li L, Zhang YG, Chen CL. Anti-apoptotic role of peroxiredoxin III in cervical cancer cells. *FEBS Open Bio* 2013; 3: 51-4.
 11. Lewitowicz P, Nasierowska-Guttmejer A, Rokita W, et al. HPV genotyping and p16/Ki-67 test significantly improve detection rate of high-grade cervical squamous intraepithelial lesion. *Arch Med Sci* 2019; 16: 87-93.
 12. Kaczmarczyk-Sedlak I, Folwarczna J, Sedlak L, et al. Effect of caffeine on biomarkers of oxidative stress in lenses of rats with streptozotocin-induced diabetes. *Arch Med Sci* 2019; 15: 1073-80.
 13. Abba MC, Laguens RM, Dulout FN, Golijow CD. The c-myc activation in cervical carcinomas and HPV 16 infections. *Mutat Res* 2004; 557: 151-8.
 14. Veldman T, Liu X, Yuan H, Schlegel R. Human papillomavirus E6 and Myc proteins associate in vivo and bind to and cooperatively activate the telomerase reverse transcriptase promoter. *Proc Natl Acad Sci USA* 2003; 100: 8211-6.
 15. McMurray HR, McCance DJ. Human papillomavirus type 16 E6 activates TERT gene transcription through induction of c-Myc and release of USF-mediated repression. *J Virol* 2003; 77: 9852-61.
 16. Kim YS, Lee HL, Lee KB, et al. Nuclear factor E2-related factor 2 dependent overexpression of sulfiredoxin and peroxiredoxin III in human lung cancer. *Korean J Intern Med* 2011; 26: 304-13.
 17. Chiribau CB, Cheng L, Cucoranu IC, Yu YS, Clempus RE, Sorescu D. FOXO3A regulates peroxiredoxin III expression in human cardiac fibroblasts. *J Biol Chem* 2008; 283: 8211-7.
 18. Jeong HJ, Jeong HW, Song SS, et al. Upregulation of peroxiredoxin III in the hippocampus of acute immobilization stress model rats and the Foxo3a-dependent expression in PC12 cells. *Cell Mol Neurobiol* 2011; 31: 1041-6.
 19. Olmos Y, Sanchez-Gomez FJ, Wild B, et al. SirT1 regulation of antioxidant genes is dependent on the formation of a FoxO3a/PGC-1alpha complex. *Antioxid Redox Signal* 2013; 19: 1507-21.
 20. Liu MH, Zhang Y, He J, et al. Upregulation of peroxiredoxin III in doxorubicin-induced cytotoxicity and the FoxO3a-dependent expression in H9c2 cardiac cells. *Exp Ther Med* 2015; 10: 1515-20.
 21. Cunniff B, Benson K, Stumpff J, et al. Mitochondrial-targeted nitroxides disrupt mitochondrial architecture and inhibit expression of peroxiredoxin 3 and FOXM1 in malignant mesothelioma cells. *J Cell Physiol* 2013; 228: 835-45.
 22. Song IS, Jeong YJ, Seo YJ, et al. Peroxiredoxin 3 maintains the survival of endometrial cancer stem cells by regulating oxidative stress. *Oncotarget* 2017; 8: 92788-800.
 23. He HC, Zhu JG, Chen XB, et al. MicroRNA-23b downregulates peroxiredoxin III in human prostate cancer. *FEBS Lett* 2012; 586: 2451-8.
 24. Li KK, Pang JC, Lau KM, et al. MiR-383 is downregulated in medulloblastoma and targets peroxiredoxin 3 (PRDX3). *Brain Pathol* 2013; 23: 413-25.
 25. Shi L, Wu LL, Yang JR, et al. Serum peroxiredoxin3 is a useful biomarker for early diagnosis and assessment of prognosis of hepatocellular carcinoma in Chinese patients. *Asian Pac J Cancer Prev* 2014; 15: 2979-86.
 26. Ismail S, Mayah W, Battia HE, et al. Plasma nuclear factor kappa B and serum peroxiredoxin 3 in early diagnosis of hepatocellular carcinoma. *Asian Pac J Cancer Prev* 2015; 16: 1657-63.
 27. Liu HL, Li TT, Yu AQ, et al. Plasma level of peroxiredoxin 3 in patients with polycystic ovarian syndrome. *BMC Endocr Disord* 2019; 19: 32.
 28. Wang L, Hao JM, Yu AQ, et al. The association of plasma peroxiredoxin 3 with insulin in pregnant women. *Biochem Biophys Res Commun* 2019; 508: 805-10.