

Expression of DNA doublestrand repair proteins in oral leukoplakia and the risk of malignant transformation

MINWEN ZHU¹, WEI LIU², LINJUN SHI¹, XUAN XIAO¹, WENYAN WU¹, LAN WU¹ and ZENGTONG ZHOU¹

Departments of ¹Oral Mucosal Diseases and ²Oral and Maxillofacial-Head and Neck Oncology, Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine, Shanghai 200011, P.R. China

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Abstract. The present study assessed the expression of the DNA doublestrand repair (DDR) proteins ATM serine/threonine kinase (ATM), checkpoint kinase 2 (CHEK2) and γ H2A histone family member X (γ H2AFX) in oral leukoplakia (OL) and evaluated their clinical significance and usefulness as biomarkers for predicting OL transformation. Retrospectively, ATM, CHEK2 and γ H2AFX protein levels were evaluated using immunohistochemical analysis in 61 OL, 33 oral squamous cell carcinoma (OSCC) and 15 normal oral mucosa tissues. OL tissues were classified into two groups according to the epithelial dysplasia pathology: The low risk dysplasia group (n=41) and the high-risk dysplasia group (n=20). The results of the present study revealed that the expression of ATM and γ H2AFX in OSCC was significantly increased compared with that in OL with low-risk dysplasia and normal oral mucosa tissues. There was no statistically significant difference in CHEK2 expression among the groups. ATM expression was correlated with that of γ H2AFX in OSCC tissue. The prognostic values of the DDR proteins and their correlation with clinical and pathological parameters were evaluated further in 99 OL patients with low risk dysplasia. Multivariate analysis revealed that increased expression of ATM and γ H2AFX was significantly associated with an increased risk of malignant transformation. Immunohistochemical analysis of ATM and γ H2AFX protein expression provided useful prognostic information on the carcinogenesis of OL. Increased ATM and γ H2AFX expression may indicate a poor prognosis.

Introduction

Oral cavity cancer is the most common head and neck cancer (1). It was estimated that 300,400 new cases of oral cavity cancer and 145,400 cases of oral cavity cancer-induced mortality have occurred in 2012 globally (2). Oral leukoplakia (OL) is one of the most common potentially malignant disorders of the oral cavity (3), with a malignant transformation rate of 17-35% (4). The prognosis and overall survival rate of patients with oral cavity cancer depend on the early detection of any lesion that may identify a patient with increased risk or with early infiltration prior to metastatic disease (3).

DNA doublestrand repair (DDR) is associated with cancer occurrence and progression (5,6). DDR activation occurs almost universally in the earliest stages of carcinogenesis (7,8). Three DDR proteins, ATM serine/threonine kinase (ATM), checkpoint kinase 2 (CHEK2) and γ H2AFX have been observed in numerous premalignant lesions and are associated with the DNA damage response (7,9-11).

ATM activates checkpoint signaling at doublestrand breaks (DSBs), following apoptosis and in response to genotoxic stresses, and thereby functions as a DNA damage sensor. ATM responds to DSBs by phosphorylating numerous substrates and may initiate DSB signaling (12).

CHEK2 is a crucial downstream target of ATM (13). Following DNA damage, ATM preferentially activates CHEK2 (14). Subsequently, activated CHEK2 modulates the activity of cell division cycle 25C, which either facilitates DNA repair or directs the cell to the apoptotic pathway (15,16). The expression of CHEK2 is aberrant in numerous human premalignant and malignant lesions (8,17,18).

H2A histone family member X (H2AFX) is a key DDR component. Within minutes of DNA damage, H2AFX is phosphorylated at its carboxyl terminus to form γ H2AFX at DSB sites (19). The formation of numerous DDR proteins requires H2AFX, indicating that H2AFX serves a key function in the early stages of DDR. H2AFX protein is phosphorylated by ATM, and the level of γ H2AFX is positively associated with the degree of DNA damage (20).

Accordingly, the present study hypothesized that alterations to ATM, CHEK2 and γ H2AFX may influence the carcinogenesis of OL. As the epithelium of OL is a useful model for monitoring abnormalities and exploring oral carcinogenesis (9), the present study evaluated the protein

Correspondence to: Dr Lan Wu or Professor Zengtong Zhou, Department of Oral Mucosal Diseases, Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine, 639 Zhizaoju Road, Shanghai 200011, P.R. China
E-mail: teana-wu@hotmail.com
E-mail: zhouzengtong@hotmail.com

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expression of ATM, CHEK2 and γ H2AFX in OL and OSCC tissues using immunohistochemistry. In addition, the present study assessed the association between the clinicopathological data and expression of these proteins, and their usefulness as biomarkers for predicting the oral carcinogenesis.

Patients and methods

Patients and collection of clinical specimens. In the present study, all patients with a clinical and pathological diagnosis of OL or OSCC at the Department of Oral Mucosal Diseases at Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine (Shanghai, China) were enrolled. The patients underwent biopsy or surgery between January 2005 and December 2014. Normal oral mucosa tissues were obtained during teeth extraction, gingivectomy or other minor surgical procedures. All the study specimens were 10% formalin-fixed for 24 h at room temperature and paraffin-embedded. Age, sex, lesion site, dietary habit, smoking history and alcohol use were also collated.

Study design. The present study was approved by the Institutional Review Board of Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine. The patients enrolled in the present study were divided into two cohorts. Cohort 1 consisted of 61 OL patients, 33 OSCC patients and 15 healthy individuals. The OL patients were classified into the low risk dysplasia group (OL low risk, n=41) and the high-risk dysplasia group (OL high risk, n=20) according to the degree of epithelial dysplasia: No/questionable/mild dysplasia (low risk) and moderate or severe dysplasia (high risk) (21). All examinations of tissues were determined by light microscope in 4 random fields (magnification, x400). A total of 33 OSCC specimens were confirmed as grade I without lymph node metastasis. The exclusion criteria for patients with OL and OSCC were as follows: (I) Any patient without an initial histopathological examination of OL and OSCC, (II) any patient treated with radiotherapy or chemotherapy prior to sampling and (III) any patient diagnosed with OL and concomitant OSCC at the first visit. The clinical characteristics of cohort 1 were summarized in Table I. Cohort 2 was based on a case-control study and included 99 patients clinically and pathologically diagnosed with OL with low risk dysplasia confirmed by the first biopsy. The inclusion criteria were as follows: (I) Patients were treated with Vita A (7.5 mg once a day for 3 months; Shanghai Donghai Pharmaceuticals Co. Ltd., Shanghai, China) and mouth rinsing (primary ingredient is gallnut containing gallic acid, 5 ml three times a day when necessary; Xinjiang Qikang Habowei Pharmaceutical Co. Ltd.) during the disease course, (II) all patients underwent two biopsies and the interval between biopsies was ≥ 3 years and (III) the lesion sites of each biopsy should remain the same. The exclusion criteria were the same as described for cohort 1. According to the results of the second biopsy, the 99 patients were classified either into the untransformed (UT) group or the malignant-transformed (MT) group.

Immunohistochemical analysis of the expression of ATM, CHEK2 and γ H2AFX. Serial tissue sections (3 μ m) from the paraffin blocks of normal oral tissues, OL and OSCC were

placed in xylene for deparaffinization and in graded alcohol dilutions (ethanol concentration was 80, 95 and 100%, respectively) for hydration. Antigen retrieval was performed with 1 mM Tris-EDTA (pH 8.0) in a 100°C water bath for 20 min and endogenous peroxidase activity was blocked with 3% hydrogen peroxide for 10 min at room temperature. Immunohistochemical analysis was performed for the sections. Primary, monoclonal antibodies against ATM (cat. no. ab78; 1:1,000), CHEK2 (cat. no. ab109413; 1:100) and γ H2AFX (cat. no. ab22551; 1:200; all Abcam, Cambridge, UK) (0.01 mol/l; pH 7.4; Wuhan Boster Biological Engineering Co., Ltd., Wuhan, China) were used for 1 h at room temperature. Following rinses with PBS three times for 10 min, a Peroxidase/DAB, K5007 EnVision™ Detection System kit (ready-to-use, Dako; Agilent Technologies, Inc., Santa Clara, CA, USA) was used to detect the primary antibodies for 1 h at room temperature, according to the manufacturer's instructions. The sections were subsequently counterstained with 4.8 mg/ml Harris Hematoxylin for 2 min at room temperature. Overall, at least three sections were stained to confirm reproducibility. The staining intensity of the cells was observed under a light microscope (Axio Scope A1; Carl Zeiss AG, Oberkochen, Germany). The mean percentage of positive cells was determined in 4 random fields (magnification, x400). To ensure pathological diagnoses were standardized, the cellular localization, intensity, and the percentage of cells with positive ATM, CHEK2 and γ H2AFX staining were assessed by two oral pathologists (Department of Oral Pathology, Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine) in a doubleblind manner and a consensus was reached in cases of discrepancy. Cell nuclear and/or cytoplasmic immunoreactivity in the epithelium was considered to indicate positive expression of ATM. Cell nuclear immunoreactivity in the epithelium was considered to indicate positive expression of CHEK2 and γ H2AFX. Positive controls for the antibodies were used according to the manufacturer's instructions. Negative control slides omitted primary antibodies. Positive staining intensity was defined as 0, 1, 2, and 3 for no staining, light yellow, yellow brown, and brown, respectively. The scoring method used for ATM and γ H2AFX was referred to by Hu *et al* (22). The positive cell percentages of 0, 1-25, 26-50, 51-75, and >75 were defined as 0, 1, 2, 3, and 4, respectively. The scoring method used for CHEK2 was a modified version of that used by Alkema *et al* (23). The positive cell percentages of 0-5, 6-25, 26-50, 51-75, and >75 were defined as 0, 1, 2, 3, and 4, respectively. The semiquantitative expression level was evaluated by multiplying the distribution and intensity score. A final score of <5 was defined as low expression of ATM and γ H2AFX, of ≥ 5 as high expression of ATM and γ H2AFX, of <7 as low expression of CHEK2, and of ≥ 7 as high expression of CHEK2.

Statistical analysis. The SPSS 19.0 software package (IBM Corp., Armonk, NY, USA) was used for all statistical analysis. ATM, CHEK2 and γ H2AFX expression levels in the normal tissue, OL low risk, OL high risk, and OSCC groups were compared using the Kruskal-Wallis test followed by Dunn's Test. The association between the expression of the proteins and clinicopathological features were assessed using

Table I. Clinical characteristics of cohort 1.

Characteristic	Normal (n=15)	OL low risk (n=41)	OL high risk (n=20)	OSCC (n=33)
Age				
Mean ± SD	44.80±15.48	56.10±11.86	57.10±10.92	56.36±13.74
Range	26-70	35-79	31-82	26-81
Sex				
Male	4	21	13	11
Female	11	20	7	12
Lesion site				
Tongue	5	18	15	26
Buccal	5	18	5	6
Gingiva	3	2	0	1
Palate	1	1	0	0
Mouth floor	0	0	0	0
Lip	1	2	0	0
Smoking history				
Never	11	22	12	24
Past and present	4	19	8	9
Alcohol intake				
Never	8	16	8	14
Past and present	7	25	12	19
Dietary habits				
Bland	10	21	14	15
Spicy	5	20	6	18

OL, oral leukoplakia; n, number of patients; OSCC, oral squamous cell carcinoma; SD, standard deviation.

a χ^2 test. Pearson correlation analysis was used to evaluate the association between protein expressions for patients in cohort 1. The Fisher's exact test was used to assess the statistical difference between the expression levels of certain proteins in cohort 2. A logistic regression model was used to evaluate the relative risk of OL malignant transformation. Ranked data were presented as percentage. All tests were two-sided. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

ATM, CHEK2 and γ H2AFX expression levels in cohort 1.

ATM was primarily expressed in the nucleus and cytoplasm in the epithelium, while CHEK2 and γ H2AFX were primarily expressed in the nucleus. Cells positive for ATM, CHEK2 and γ H2AFX expression were detected not only in the stratum basale and stratum spinosum, but also in the stratum granulosum, and in the stratum corneum of certain patients (Fig. 1). ATM expression tended to increase gradually in the normal tissue, OL low risk, OL high risk and OSCC groups during carcinogenesis ($P=0.005$; Table II). ATM expression was significantly increased in 29/33 of the samples of the OSCC group, compared with the normal tissue group ($P=0.008$). In addition, ATM expression was significantly increased in 23/41 of the samples of the OL low risk group compared with the OSCC group ($P=0.027$). In the

OL high-risk group, ATM expression in 13/20 of the samples exhibited no statistically significant difference compared with that in the other three groups, respectively. γ H2AFX expression increased in the groups as carcinogenesis progressed ($P=0.001$; Table II). A comparison of the groups revealed that there was a significant difference between the OL low risk and the OSCC groups ($P=0.014$), and between the normal tissue and the OSCC groups in γ H2AFX expression ($P=0.001$). There was no significant difference in CHEK2 expression among the four groups ($P=0.074$; Table II). The expression of ATM and γ H2AFX increased in the groups as carcinogenesis progressed. To assess this association, Pearson correlation analysis was performed. The result was statistically significant [($P=0.045$; Pearson correlation coefficient (r)=0.192)] among the groups. correlation demonstrated in the OSCC group exhibited [$r=0.383$ ($P=0.028$)]

Correlation and clinical significance of ATM and γ H2AFX expression in normal oral mucosa, OL and OSCC tissues in cohort 1.

The association between ATM and γ H2AFX expression and multiple clinical characteristics of OL in cohort 1 ($n=109$) was assessed using the χ^2 test. Multiple degrees of epithelial dysplasia revealed different expression levels of ATM ($P=0.004$; Table III). OSCC and normal tissues demonstrated the highest and lowest percentage, respectively, of ATM expression among the tissues. No correlation was demonstrated between ATM expression and the other clinical characteristics:

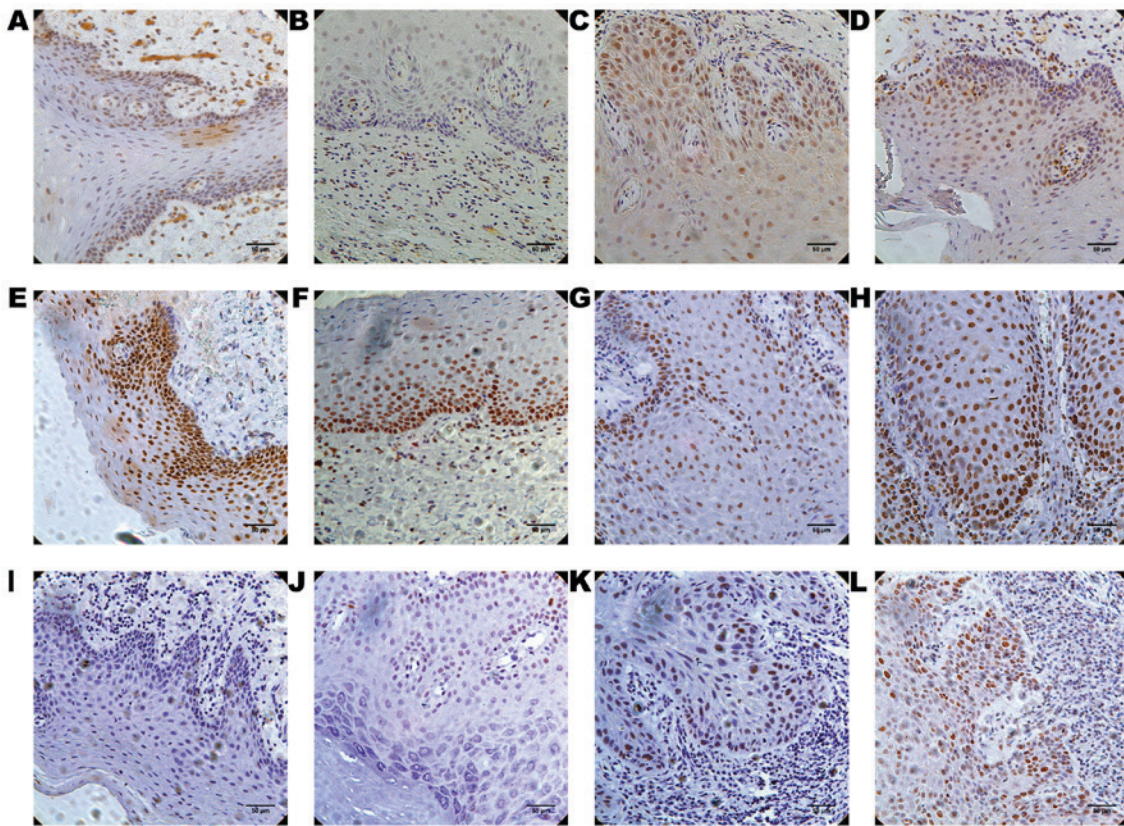


Figure 1. Expression of ATM, CHEK2 and γ H2AFX in the four tissue groups (magnification x400). (A) Expression of ATM in normal oral tissue. (B) Expression of ATM in OL low risk tissue. (C) Expression of ATM in OL high risk tissue. (D) Expression of ATM in OSCC tissue. (E) Expression of CHEK2 in normal oral tissue. (F) Expression of CHEK2 in OL low risk tissue. (G) Expression of CHEK2 in OL high risk tissue. (H) Expression of CHEK2 in OSCC tissue. (I) Expression of γ H2AFX in normal oral tissue. (J) Expression of γ H2AFX in OL low risk tissue. (K) Expression of γ H2AFX in OL high risk tissue. (L) Expression of γ H2AFX in OSCC tissue. ATM, ATM serine/threonine kinase; CHEK2, checkpoint kinase 2; γ H2AFX, γ H2A histone family member X; OL, oral leukoplakia; OSCC, oral squamous cell carcinoma.

Age, sex, lesion site, dietary habits, smoking history and alcohol use. γ H2AFX expression was also associated with the degree of epithelial dysplasia ($P=0.001$; Table IV). OSCC and normal tissues revealed the highest and lowest percentage, respectively, of γ H2AFX expression among the tissues. No clinical characteristics revealed association with γ H2AFX expression.

Identifying candidate DDR proteins in cohort 1. The selection criteria for candidate DDR proteins were as follows: The expression of candidate DDR proteins increased or decreased successively with respect to increasing grades of carcinogenesis, with significance set at $P<0.05$ and assessed using the Kruskal-Wallis test. According to the results of the present study, not only did none of the three proteins assessed demonstrate decreased expression as carcinogenesis progressed but ATM and γ H2AFX expression increased as carcinogenesis progressed. ATM and γ H2AFX expression levels were also significantly different between the normal tissue and the OSCC groups, and between the OL low risk and the OSCC groups. Therefore, ATM and γ H2AFX expression was assessed further and validated in the case-control study.

ATM and γ H2AFX expression in cohort 2. Cohort 2 consisted of 99 patients with OL low risk. On the basis of the second biopsy, these patients were assigned to the UT group ($n=81$) or the MT group ($n=18$). Table V summarizes the characteristics

of these patients. High expression of ATM was demonstrated in 54 of the 99 patients (54.5%). Increased expression of ATM was detected in 40 of the 81 (49.4%) patients in the UT group and in 14 of the 18 (77.8%) patients in the MT group ($P=0.037$; Fig. 2). In addition, 29 of the 99 patients (29.3%) demonstrated high expression of γ H2AFX. Increased expression of γ H2AFX was detected in 19 of the 81 (23.5%) patients in the UT group and in 10 of the 18 (55.6%) patients in the MT group ($P=0.01$; Fig. 2).

High expression of ATM as an independent factor for OL malignant transformation in cohort 2. To evaluate the risk of OL malignant transformation, clinicopathological parameters, and ATM and γ H2AFX expression were assessed using logistic regression (Table VI). In the univariate analysis, age, sex, lesion site, smoking history, and alcohol use were not significant risk factors for transformation in cohort 2, and high expression of ATM and γ H2AFX was associated with a 3.59-fold [95% confidence interval (CI), 1.09-11.83; $P=0.036$] and a 4.08-fold (95% CI, 1.41-11.80; $P=0.009$), increase in the risk of malignant transformation, respectively. In the multivariate analysis, high expression of ATM and γ H2AFX was also significantly associated with an increased risk of malignant transformation. The adjusted odds ratio for malignant transformation was 4.29 for high ATM expression (95% CI, 1.22-15.07; $P=0.023$) and 4.79 for high γ H2AFX expression (95% CI, 1.56-14.73; $P=0.006$).

Table II. ATM, CHEK2 and γ H2AFX expression in cohort 1 (n=109).

Expression	Total (n)	Normal control (n)	OL low risk (n)	OL high risk (n)	OSCC (n)	P-value
ATM						0.005
Low	38	9	18	7	4	
High	71	6	23	13	29	
CHEK2						0.074
Low	20	1	12	1	6	
High	89	14	29	19	27	
γ H2AFX						0.001
Low	79	15	33	15	16	
High	30	0	8	5	17	

ATM, ATM serine/threonine kinase; CHEK2, checkpoint kinase 2; γ H2AFX, γ H2A histone family member X; n, number of patients; OL, oral leukoplakia; OSCC, oral squamous cell carcinoma.

Table III. Association between ATM expression and clinicopathological features in cohort 1 (n=109).

Clinicopathological feature	Total (n)	Low ATM expression (n, %)		High ATM expression (n, %)		P-value
Age, years						0.866
\leq 60	70	24	34.3	46	65.7	
$>$ 60	39	14	35.9	25	64.1	
Sex						0.711
Male	49	18	36.7	31	63.3	
Female	60	20	33.3	40	66.7	
Lesion site						0.345
Tongue	64	20	31.3	44	68.8	
Non-tongue	45	18	40.0	27	60.0	
Dietary habits						0.333
Bland	62	24	38.7	38	61.3	
Spicy	47	14	29.8	33	70.2	
Smoking history						0.261
Never	68	21	30.9	47	69.1	
Past and present	41	17	41.5	24	58.5	
Alcohol intake						0.345
Never	45	18	40.0	27	60.0	
Past and present	64	20	31.3	44	68.8	
Epithelial dysplasia						0.004
Normal	15	9	60.0	6	40.0	
OL low risk	41	18	43.9	23	56.1	
OL high risk	20	7	35.0	13	65.0	
OSCC	33	4	12.1	29	87.9	

Non-tongue denotes buccal, gingiva, palate, mouth floor, or lip. ATM, ATM serine/threonine kinase; n, number of patients; OL, oral leukoplakia; OSCC, oral squamous cell carcinoma.

Discussion

To the best of our knowledge, the present study is the first to evaluate ATM, CHEK2 and γ H2AFX expression in patients

with OL with multiple degrees of epithelial dysplasia and to assess the functions of these proteins in predicting the risk of OSCC in two independent cohorts using immunohistochemical analysis. ATM serves a key function in the DNA

Table IV. Association between γ H2AFX expression and clinicopathological features in cohort 1 (n=109).

Clinicopathological feature	Total (n)	Low γ H2AFX expression (n, %)		High γ H2AFX expression (n, %)		P-value
Age, years						0.905
≤ 60	70	51	72.9	19	27.1	
> 60	39	28	71.8	11	28.2	
Sex						0.834
Male	49	36	73.5	13	26.5	
Female	60	43	71.7	17	28.3	
Lesion site						0.056
Tongue	64	42	65.6	22	34.4	
Non-tongue	45	37	82.2	8	17.8	
Dietary habits						0.402
Bland	62	43	69.4	19	30.6	
Spicy	47	36	76.6	11	23.4	
Smoking history						0.9
Never	68	49	72.1	19	27.9	
Past and present	41	30	73.2	11	26.8	
Alcohol intake						0.14
Never	45	36	80.0	9	20.0	
Past and present	64	43	67.2	21	32.8	
Epithelial dysplasia						0.001
Normal	15	15	100.0	0	0.0	
OL low risk	41	33	80.5	8	19.5	
OL high risk	20	15	75.0	5	25.0	
OSCC	33	16	48.5	17	51.5	

Non-tongue denotes buccal, gingiva, palate, mouth floor, or lip. γ H2AFX, γ H2A histone family member X; n, number of patients; OL, oral leukoplakia; OSCC, oral squamous cell carcinoma.

DSB-induced signaling cascade. Tumorigenic events that occur early in the progression of major types of human cancer activate ATM-regulated cell cycle checkpoints and thereby an inducible barrier that inhibits tumor progression and genetic instability (7,24). A previous study has demonstrated that ATM expression was increased in certain types of cancer tissue compared with that in benign tumorous lesions and normal tissues (22). Other previous studies have suggested that ATM potentially represents a promising indicator for hyperplasia and cancer, and may serve as a useful marker for identifying patients with poor prognosis (25,26). Raynaud *et al* (10) demonstrated that the difference in ATM activation between normal and precancerous tissues was not significant, though ATM expression differed significantly between precancerous and cancerous tissues. The results of the aforementioned studies support those of the present study.

He *et al* (9) indicated that ATM protein expression was higher in OL compared with that in normal oral tissue, but demonstrated no significant difference between OL and OSCC tissues in ATM protein expression. The present study dynamically observed the activation of the DNA damage signaling pathway in normal oral mucosa, OL (low risk dysplasia and high risk dysplasia) and OSCC tissues in a large population. In contrast to the results demonstrated by

He *et al* (9) those demonstrated in the present study revealed that ATM expression gradually increased as OL progressed to OSCC. In addition, the present study revealed a significant difference in ATM expression between OL with low risk dysplasia and OSCC. In the present study, ATM expression correlated with the degree of epithelial dysplasia during carcinogenesis and age, sex, lesion site, dietary habits, smoking history, and alcohol use were not significant factors in the expression of ATM. Therefore, the results of the present study indicated that ATM was activated in oral precancerous lesions and served a function in the early stages of oral carcinogenesis.

One of the proteins phosphorylated following DNA damage, a process initiated by ATM, is H2AFX, which, in phosphorylated form (γ H2AFX), functions as a specific indicator for the presence of DSBs (27). Increased γ H2AFX expression may result in increased radiosensitivity (28). Multiple studies have revealed that γ H2AFX expression is increased in certain types of cancer and their premalignant lesions (8,22,29,30), which supports the results of the present study. Overexpression of γ H2AFX may represent an independent prognostic indicator of a poor overall patient survival rate (31,32). In contrast to the results of the present study, Chou *et al* (33) reported that γ H2AFX expression was increased in dysplastic epithelium and significantly

Table V. Clinical characteristics of cohort 2.

Characteristic	UT (n=81)	MT (n=18)
Age, years		
Mean \pm SD	56.53 \pm 11.27	54.83 \pm 11.63
Range	31-82	35-71
Sex		
Male	44	8
Female	37	10
Lesion site		
Tongue	44	13
Buccal	32	3
Gingiva	2	1
Palate	1	0
Mouth floor	0	0
Lip	2	1
Dietary habits		
Bland	50	11
Spicy	31	7
Smoking history		
Never	50	11
Past and present	31	7
Alcohol intake		
Never	33	6
Past and present	48	12
ATM expression		
Low	41	4
High	40	14
γ H2AFX expression		
Low	62	8
High	19	10

SD, standard deviation; ATM, ATM serine/threonine kinase; γ H2AFX, γ H2A histone family member X.

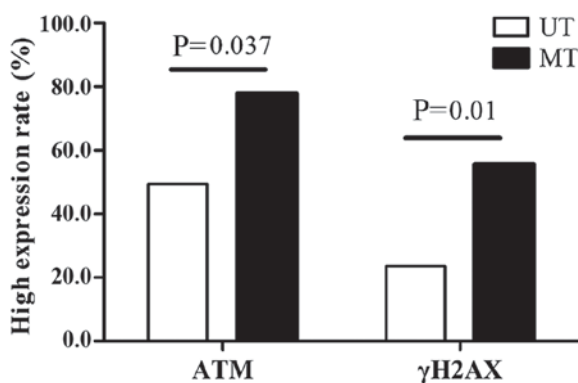


Figure 2. Frequency of increased ATM and γ H2AFX expression in oral leukoplakia low risk tissue. ATM, ATM serine/threonine kinase; γ H2AFX, γ H2A histone family member X; UT, untransformed; MT, malignant-transformed.

decreased in OSCC tissue. The results of the present study revealed that γ H2AFX expression increased in OSCC tissue

Table VI. Logistic regression analysis of the potential risk of oral cancer.

Characteristic	OR (95% CI)	P-value
Univariate analysis		
Age	1.36 (0.48-3.82)	0.560
Sex	0.67 (0.24-1.88)	0.449
Lesion site	2.19 (0.71-6.70)	0.171
Dietary habits	1.03 (0.36-2.93)	0.961
Smoking history	1.03 (0.36-2.93)	0.961
Alcohol intake	1.38 (0.47-4.03)	0.562
High ATM expression	3.59 (1.09-11.83)	0.036
High γ H2AFX expression	4.08 (1.41-11.80)	0.009
Multivariate analysis		
High ATM expression	4.29 (1.22-15.07)	0.023
High γ H2AFX expression	4.79 (1.56-14.73)	0.006

OR, odds ratio; CI, confidence interval; ATM, ATM serine/threonine kinase; γ H2AFX, γ H2A histone family member X.

with increasing disease severity; this discrepancy between the aforementioned and present study may be due to the difference in the OSCC tumor differentiation grade selected. In the present study, similar to ATM expression, γ H2AFX expression correlated with the degree of epithelial dysplasia, according to the results of the associations between clinicopathological features and γ H2AFX expression. Detecting γ H2AFX expression may help to evaluate precancerous oral cavity lesions and monitor cancer progression.

CHEK2 serves a key function in inhibiting cell cycle progression in response to the DNA damage pathway (34). In multiple types of solid tumor, CHEK2 expression was decreased compared with that in normal tissues (35-37). CHEK2 expression in oral precancerous lesions is rarely assessed. In the present study, the expression of CHEK2 protein altered during oral carcinogenesis. There were no significant differences between any two groups of the four during carcinogenesis. Based on these conflicting results, the present study suggested that: (I) Aberrant CHEK2 protein may be functionally defective and regulated by other, unknown upstream proteins during DDR; (II) CHEK2 protein expression is regulated differently depending on the type of carcinoma; (III) the ATM-CHEK2 pathway may not be associated with oral carcinoma or precancerous lesions or (IV) more complex signaling pathways may participate in the DNA damage response.

Pearson correlation analysis of cohort 1 demonstrated that γ H2AFX and ATM expression was correlated ($P=0.045$; $r=0.192$), particularly in OSCC tissue ($P=0.028$; $r=0.383$). Therefore, the present study suggested that the ATM- γ H2AFX pathway contributes to the DNA damage response.

For cohort 2, the present study assessed the prognostic value of ATM and γ H2AFX expression and evaluate whether the prognostic value was independent of clinicopathological factors. Univariate and multivariate analysis revealed that increased ATM and γ H2AFX expression was significantly

associated with an increased risk of transformation ($P < 0.05$). The results of the present study indicated that increased ATM and γ H2AFX expression served as an independent predictor of carcinogenesis. However, age, sex, lesion site, dietary habits, smoking history, and alcohol use were not revealed to be significant risk factors for OL malignant transformation in the cohort 2, a result that reflects that of multiple previous studies (38-40).

To conclude, the results of the present study suggested that ATM and γ H2AFX expression in OL tissue was associated with oral cancer progression. Immunohistochemical staining of ATM and γ H2AFX may represent a promising technique for the early identification and risk evaluation of OSCC in patients with precancerous oral lesions. Further studies are required to assess the function of ATM and γ H2AFX in oral carcinogenesis, including for grade I, II and III OSCC. Further study of the mechanisms underlying DNA damage and response in OL tissue is also required.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

LW, ZZ and WL conceived and designed the study. MZ, LS, XX and WW performed the experiments. MZ analyzed the data and wrote the manuscript. LW and ZZ reviewed and edited the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The present study was approved by the Institutional Review Board of Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine.

Consent for publication

The subjects or parent/guardian that participated in the present study provided written informed consents for publication.

Competing interests

The authors declare that they have no competing interests.

Authors' information

LW, Department of Oral Mucosal Diseases, Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine, committee member of the Oral Mucosal Disease Professional Committee of Shanghai Stomatological Association; committee member of Chinese and Western medicine professional committee of Chinese Stomatological Association.

ZZ, Department of Oral Mucosal Diseases, Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine, former chairman of the oral mucosal disease professional committee of Chinese Stomatological Association; former chairman of the oral mucosal disease professional committee of Shanghai Stomatological Association.

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