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Rapid method for determination of DNA repair capacity in human peripheral blood lymphocytes amongst smokers

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Abstract

Background: DNA repair capacity is an important determinant of susceptibility to cancer. The hOGG1 enzyme is crucial for repairing the 8-oxoguanine lesion that occurs either as a byproduct of oxidative metabolism or as a result of exogenous sources such as exposure to cigarette smoke. It has been previously reported that smokers with low hOGG1 activity had significantly higher risk of developing lung cancer as compared to smokers with high hOGG1 activity.

Methods: In the current study we elucidate the association between plasma levels of 8-OHdG and the OGG1 repair capacity. We used the commercially available 8-OHdG ELISA (enzyme-linked immunosorbent assay), the Comet assay/FLARE hOGG1 (Fragment Length Analysis by Repair Enzymes) assay for quantification of the levels of 8-OHdG and measured the constitutive, induced and unrepaired residual damage, respectively. We compared the DNA repair capacity in peripheral blood lymphocytes following H₂O₂ exposure in 30 lung cancer patients, 30 non-, 30 former and 30 current smoker controls matched by age and gender.

Results: Our results show that lung cancer cases and current smoker controls have similar levels of 8-OHdG lesions that are significantly higher compared to the non-smokers controls. However, lung cancer cases showed significantly poorer repair capacity compared to all controls tested, including the current smokers controls. After adjustment for age, gender and family history of smoking-related cancer using linear regression, we observed a 5-fold increase in risk of lung cancer associated with high levels of residual damage/reduced repair capacity. Reduced OGG1 activity could be expected to be a risk factor in other smoking-related cancers.

Conclusion: Our study shows that the Comet/FLARE assay is a relatively rapid and useful method for determination of DNA repair capacity. Using this assay we could identify individuals with high levels of residual damage and hence poor repair capacity who would be good candidates for intensive follow-up and screening.

Background

DNA repair is a ubiquitous defense mechanism that is critical to maintaining the integrity of the genome. Several prior studies have reported associations between reduced repair capacity and cancer development [1-3]. Base excision repair (BER) is the main pathway involved in the repair of oxidative DNA damage and recent studies have indicated a correlation between reduced BER capacity and oncogenesis [4,5]. The oxidative DNA damage induced by reactive oxygen species (ROS) may

lead to single- or double-strand breaks, point and frame-shift mutations and larger-scale chromosome abnormalities [6]. ROS can be formed endogenously as a result of xenobiotic metabolism by cytochrome P450 enzymes, by redox cycling of hormone metabolites or exposure to other environmental factors [7].

Smoking is an inducer of oxidative stress which results in ROS-induced DNA damage [8] in the form of DNA adducts. This may be of particular importance in lung cancer where increased oxidative DNA damage coupled with reduced BER may play an important role in modifying the disease risk [9]. It is well known that only a small percentage of smokers develop lung cancer [10]

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suggesting that individual DNA repair capacity may play a significant role in the carcinogenic process.

The 8-oxoguanine DNA glycosylase (hOGG1) is one of the key DNA repair enzymes involved in the BER pathway in humans [11,12]. It recognizes the 8-oxoguanine modifications from both nuclear and mitochondrial DNA [13]. Among the more than 30 different products of modified DNA and RNA by oxidative damage [14], 8-Hydroxy-2'-deoxyguanosine (8-OHdG) is the most extensively studied induced lesion. 8-OHdG lesion causes G→T and A→C transversions [15] that have been reported as the sites of spontaneous oncogene expression and ultimately cancer manifestation [16,17]. Deletion of the hOGG1 gene was shown to be associated with accumulation of 8-OHdG lesion and increase in mutational risk [18]. OGG1 deficiency in yeast was shown to result in a spontaneous mutator phenotype [19]. Collins et al observed a lack of repair activity on a DNA substrate containing 8-OHdG protein extracts from homozygous OGG1 knockout mouse embryonic fibroblasts [20]. In contrast, mammalian cells over-expressing OGG1 repair 8-OHdG more rapidly after toxic insult with oxidants [21]. Increased levels of 8-OHdG has been associated with an increased risk of lung cancer among smokers [22]. In a recent study, Paz-Elizur et al evaluated the enzymatic OGG1 activity among smoking lung cancer cases and controls. The authors reported that the OGG1 activity was significantly decreased in lung cancer cases, and that the risk of developing lung cancer in smokers with low OGG1 activity was significantly higher compared to smokers with high OGG1 activity [23]. To date, the association between the levels of 8-OHdG and OGG1 repair capacity has not been determined.

In the current study we hypothesized that an inverse association exists between the levels of 8-OHdG and OGG1 activity. Furthermore, we hypothesized that lung cancer patients have higher levels of 8-OHdG reflecting suboptimal OGG1 repair capacity when compared to healthy controls. To test these hypotheses, we quantified the levels of 8-OHdG and measured ROS-induced DNA damage and repair in peripheral blood lymphocytes of lung cancer cases and controls. We used the commercially available 8-OHdG ELISA (enzyme-linked immunosorbent assay) to measure the levels of 8-OHdG in plasma and the Comet assay and hOGG1 FLARE (Fragment Length Analysis by Repair Enzymes) assay to measure the constitutive, induced and unrepaired residual damage by measuring DNA strand breaks and abasic sites. The Comet assay is used to detect DNA damage (including single-strand breaks, double-strand breaks and alkali-labile sites) [24,25] as well as to determine interindividual variation in DNA repair capacity through the measurement of residual DNA damage after

exposure to mutagens [26,27]. The hOGG1 FLARE is a modified Comet assay that employs 8-oxoguanine DNA glycosylase to recognize and nick the DNA specifically at the 8-OHdG lesion thus resulting in strand breaks [28]. Cells not exposed to hOGG1 serve as a baseline for naturally occurring DNA breaks; while cells exposed to hOGG1 contain both naturally occurring breaks in DNA, as well as breaks caused by hOGG1 at places containing unrepaired 8-OHdG adducts [29]. Our results show that lung cancer cases and current smoker controls had higher levels of 8-OHdG compared to the former and non-smokers controls. Similarly, our results demonstrated that baseline DNA damage is comparable between lung cancer cases and current smoker controls but is significantly higher than that observed among former smokers and non-smokers controls. Further investigation revealed that lung cancer cases showed significantly poorer repair capacity than that observed in current smoker controls suggesting impairment of adduct removal. In conclusion, our study shows that the Comet/FLARE assay is a relatively rapid and useful method for determination of DNA repair capacity and that adduct removal by OGG1 is a mechanism that might be significantly impaired among smokers who develop lung cancer. Using this assay we could potentially identify individuals with high levels of residual damage and hence poor repair capacity. Identification of the small percentage of susceptible smokers with poor repair would allow for better prevention interventions.

Methods

Study population

A total of 120 participants were included in the study. Thirty smoker lung cancer patients were recruited from The University of Texas M. D. Anderson Cancer Center as part of an ongoing lung cancer case-control study. The patients had newly diagnosed, histologically confirmed lung cancer with no prior history of treatment (radiation or chemotherapy) upon enrollment in the study. Controls (n = 90) were matched to cases on age (+/-5 years) and gender and were recruited from the Kelsey-Seybold Clinics in Houston, Texas. The control participants consisted of 30 non-smokers, 30 former smokers and 30 current smokers with no previous history of cancer. After informed consent was obtained, a personal interview was conducted and information regarding socio-demographics, smoking history, alcohol consumption, occupational exposures, diet and family history of cancer was collected. There were no age and gender restrictions for study eligibility. The institutional review boards at both M. D. Anderson Cancer Center and Kelsey-Seybold Clinic approved this study. After providing informed consent, 10-mL blood was collected from all study participants.

Peripheral blood lymphocyte isolation

Peripheral blood and lymphocytes were isolated using standard Ficoll-Histopaque method. Briefly, 10 ml of whole blood from each subject was drawn into heparinized tubes, layered over 10 ml of Histopaque -1077 (Sigma Aldrich Co., St. Louis, MO) at room temperature and centrifuged at 1500 rpm for 30 min. The mononuclear cells were removed from the interphase, washed twice with phosphate-buffered saline pH 7.2 and centrifuged at 1500 rpm for 10 min. Cells were suspended in 1 ml of RPMI 1640 medium and the cell membrane integrity was determined by Trypan Blue solution 0.4% and adjusted to a concentration of 1×10^6 cells/ml. Duplicate lymphocyte cultures were prepared for each study subject. Lymphocytes were then resuspended in RPMI 1640 medium supplemented with 1% L-glutamine, 100 U/ml of penicillin and of streptomycin (Gibco Invitrogen Corp., Grand Island, NY), 10% fetal calf serum, 2% PHA (Murex Biotech Ltd., Dartford, England, UK) and cultured in 24-well microplates for 24 h at 37°C in 5% CO₂ prior to treatment.

Establishment of hydrogen peroxide (H₂O₂) treatment conditions

We used H₂O₂ as the ROS-inducing agent in vitro. In order to establish the optimal conditions for exposure and repair kinetics, peripheral blood lymphocytes were isolated from 20 healthy controls. A dose-response curve was generated using in vitro treatment of lymphocytes with H₂O₂ (at a range of concentrations between 20-100 μM) for 15 min. Results from the dose response curve indicated that the 60 μM concentration, lead to a cell viability >85% and DNA damage that was significantly higher than the negative control ($P < 0.001$) (Additional file 1, Figure S1A).

The cells concentration was adjusted to 3×10^4 cells/ml and suspended in RPMI 1640 medium without FBS, supplemented with 2 mM L-glutamine, 100 U/ml of penicillin/streptomycin and 20% Alamar Blue cell assay™ (BioSource International Inc., Camarillo, CA, USA) to determine metabolic activity/cell viability of the cells. The cell viability was measured for each time of treatment with a spectrophotometer (Tecan, Genios Pro) at 535 nm excitation and 590 nm emission. Using Pearson's correlation we estimated the correlation between the most common parameters reported in the literature to determine damage by the Comet assay namely: the tail extent moment (TEM) and the tail length (TL). Our results indicated a pair-wise correlation of $R^2 = 0.94-0.96$ ($P < 0.001$) and therefore in the following sections we reported our findings using the TEM only.

Optimization of recovery time for H₂O₂-induced damage/repair

A normal lymphoblastoid cell line (AH, [cat#GM14520], Coriell Cell Repositories, Camden, NJ, USA) was used

to determine the optimal incubation time for recovery of in vitro H₂O₂-induced damage. Significant DNA damage TEM was observed after 15 min H₂O₂-treatment as compared to the untreated cells (mean + SD = $0.89 + 0.18$ versus $0.47 + 0.07$ respectively, $P = 0.002$). After 1 h incubation at 37°C to allow the cells to recover, a significant reduction in TEM was observed as compared to the H₂O₂-treated cells (mean ± SD = 0.65 ± 0.11 versus 0.89 ± 0.18 respectively, $P = 0.007$); however the damage was still significantly higher than that observed in the untreated cells (mean + SD = $0.65 + 0.11$ versus $0.47 + 0.07$ respectively, $P = 0.01$). Incubation of the cells for 2 h at 37°C showed a further reduction in the level of measured TEM when compared to the 1 h incubation (mean ± SD = 0.51 ± 0.05 versus 0.65 ± 0.11 , respectively); however these differences were not statistically significant and we therefore decided to use the one-hour incubation in our subsequent experiments in order to accommodate the further experimental procedures that included a further incubation with the hOGG1 enzyme followed by denaturation and electrophoresis (Additional file 1, Figure S1B).

Measurement of 8-OHdG levels

Baseline serum levels of 8-OHdG was measured using the commercially available 8-OHdG ELISA Kit (BIOXY-TECH® OXIS International Inc., Foster City, CA, USA) following manufacturer's recommendations. This kit is a competitive ELISA for quantitative measurement of 8-OHdG in tissue, serum or plasma resulting from oxidative damage to DNA.

Comet/FLARE® assay

Collins et al [20] first introduced this technique to study BER pathway. The principal of this assay depends on incubating cell extracts with a substrate of DNA containing specific lesions. Incubation of the substrate DNA with the specific enzymes allows for the uncovering of specific damaged bases. Treating lysed, immobilized cells with a DNA glycosylase, converts the base to an alkali sensitive site, followed by DNA unwinding that allows additional damage to be recognized. This leads to the introduction of breaks that are measured using the single-cell gel electrophoresis (Comet) assay. On the basis of the initial results, the lymphocytes were treated with a concentration of 60 μM H₂O₂ in a total volume of 200 μM RPMI at 4°C for 15 min. Cells were then incubated at 37°C for 1 h to allow the cells to recover. The cell suspension was then aspirated from the wells and washed with PBS pH 7.2. Cells were then embedded in a layer of low melting point agarose (LMA) at 37°C prepared in PBS and the mixtures were immediately transferred to the FLARE®-slides (Trevigen Inc.). Duplicate slides were prepared from each well. Slides were

immersed in lysis solution (2.5 M NaCl, 100 mM EDTA pH 10, 10 mM Tris base, 1% sodium lauryl sarcosinate, 1% Triton X-100 and 1% DMSO) (Trevigen Inc.) for 1 h at 4°C. The slides were washed in FLARE buffer (250 mM HEPES-KOH pH 7.4, 2.5 M KCl and 250 mM EDTA) at room temperature three times over a 15 min period. 100 µl of hOGG1 (1:500) was then added to each sample and incubated at 37°C for 40 min (following manufacturer's recommendations) to allow for further cell recovery. The DNA of the nuclei in the agarose gels was denatured in electrophoresis buffer pH 12.1 (3 M NaCl, 500 mM EDTA) for 30 min at 4°C. Electrophoresis was performed in alkaline solution pH 13 applying 300 mA, 25 V for 30 min at 4°C. The slides were then placed in cold methanol for 20 min, dried and stored in a slide box at room temperature. At time of analysis, the slides were hydrated in cold water at 4°C for 20 min and stained with ethidium bromide solution (2 µg/mL). Slides were coded in order to blind the treatment conditions from the scorer and 100 random cells/sample were analyzed (50 cells/duplicate slide) using an epi-fluorescence microscope (Nikon) equipped with Comet IV[®] software (Perceptive Imaging, Haverhill, Suffolk, UK). The mean values of the TEM in each cell were computed for each experimental group.

Statistical analysis

Descriptive statistics were used to characterize the cases and controls. Analysis of variance (ANOVA) was used to compare age and other continuous variables and the Chi-square test was used to compare the categorical variables. For the FLARE assay, the Percent Residual DNA Damage (PRD) was calculated using the following equation:

$$PRD = \frac{X_{H_2O_2} - OGG - X_{H_2O_2} \times X_{H_2O_2}}{X_{H_2O_2} - OGG} \times 100$$
 Where $X_{H_2O_2}$ is the mean TEM values for each study subject after treatment with H₂O₂, while $X_{H_2O_2} + OGG$ is the mean TEM values for each study subject after H₂O₂ treatment and further incubation with OGG1 enzyme for removal of additional residual damage. Linear regression analysis was conducted to adjust the TEM H₂O₂ + OGG1 by TEM H₂O₂. The residual values from the regression defined as Residual Damage Linear Regression (RDLR) among the cases and controls were computed using ANOVA. RDLR was dichotomized using the 75th percentile of the controls and estimates of lung cancer risk (OR and 95% CL) for RDLR adjusted for age, sex and family history of a smoking related cancer were determined.

Results

Demographics and the study population

The demographic characteristics of the 30 cases and 90 controls are summarized in Table 1. No significant

differences were observed between cases and the controls did not differ significantly in terms of age, gender or ethnicity. Fifty-two percent of the patients self-reported a family history of smoking-related cancer, compared with 40% of the current, 59% of the former and 50% of the non-smoker controls ($P = 0.07$). Cases had on an average smoked cigarettes for 44.23 years, compared with 42.70 and 35.37 years for current and former smoker controls ($P > 0.05$ and $P < 0.001$, respectively); however both cases and controls (current and former groups) smoked about the same number of cigarettes per day (mean number of cigarettes per day \pm SE for cases = 29.2 ± 2.5 and 24.4 ± 1.75 and 35.0 ± 3.3 for current and former smokers respectively, $P = 0.10$).

Measurement of 8-OHdG levels at baseline by ELISA

Among controls, the current and former smoker controls showed significantly higher levels as compared to non-smoker controls (mean \pm SE, 1.99 ± 0.55 ng/ml, 1.68 ± 0.99 ng/ml versus 0.83 ± 0.62 respectively, $P < 0.001$). There was no significant difference between the level of 8-OHdG among the former and current smokers controls ($P = 0.28$). Lung cancer cases had over 2-fold higher levels of 8-OHdG than the non-smoker controls (mean \pm SE, 2.09 ± 0.91 ng/ml versus 0.83 ± 0.62 , $P < 0.001$); however the difference was not significant when compared to former and current smoker controls ($P = 0.88$ and $P = 0.33$, respectively).

Measurement of DNA damage at baseline by Comet assay

Table 2 shows the extent of baseline DNA damage measured as TEM. Among controls, the non- and former smokers had significantly lower DNA damage than the current smoker controls (mean \pm SE = 0.26 ± 0.11 and 0.23 ± 0.09 versus 0.54 ± 0.15 , $P < 0.001$). There was no significant difference between the level of baseline DNA damage among the non- and former smokers controls ($P = 0.95$).

Comparing cases and controls, the baseline levels of DNA damage was significantly different between groups (0.57 ± 0.12 versus 0.35 ± 0.19 , $P < 0.001$). However, when the controls were subdivided by smoking status, the levels of TEM were significantly different between lung cancer cases and non- and former smoker controls only ($P < 0.001$). There was no difference between the level of baseline DNA damage among the current smoker controls and the lung cancer cases (mean \pm SE = 0.54 ± 0.15 versus 0.57 ± 0.12 respectively, $P = 0.83$).

Measurement of H₂O₂-induced DNA damage by Comet assay

Level of H₂O₂-induced DNA damage after an hour incubation (to allow cellular intrinsic repair of damage)

Table 1 Comparison of demographic and clinical factors among cases and controls

	Cases		Controls		P-value
	N = 30	Current Smokers N = 30	Former Smokers N = 30	Non-Smokers N = 30	
Age (mean ± SE)	62.7 ± 0.57	63.5 ± 0.84	61.9 ± 0.69	58.6 ± 0.98	0.27
Gender: N (%)					
Male	15(50.0)	20(66.7)	21(70.0)	20(66.7)	
Female	15(50.0)	10(33.3)	9(30.0)	10(33.3)	0.58
Ethnicity					
Anglo	30(100.0)	29(96.7)	25(83.3)	30(100.0)	
Non-Anglo	0(0.0)	1(3.3)	5(16.7)	0(0.0)	0.24
No. of Cigarettes/day Mean ± SE	29.2 ± 2.5	24.4 ± 1.75	35.0 ± 3.3	0.0 ± 0.0	<0.001
Years smoked Mean ± SE:	44.23 ± 1.39	42.70 ± 1.46	35.37 ± 1.58	0.0 ± 0.0	<0.001
Family History Cancer					
N (%)					
No	14(48.3)	15(60.0)	12(41.4)	15(50.0)	
Yes	15(51.7)	10(40.0)	17(58.6)	15(50.0)	0.07

P-values from Chi-square test for association (categorical) or ANOVA (continuous)

is shown in Table 2. The level of TEM in all three groups of controls was significantly different with mean + SD = 3.2 + 0.82; 5.8 + 0.31 and 10.2 + 0.93 for non-former and current smoker controls, respectively. Lung cancer cases had over 6-, 3- and 1-fold higher levels of induced DNA damage than the non-, former and current smoker controls, respectively. The lung cancer cases had the highest level of DNA damage followed by the current smokers (mean + SD = 19.5 + 0.41 versus 10.2 + 0.93, $P < 0.001$). The non-smokers exhibited the least damage (mean + SD = 3.2 + 0.82 versus 5.8 + 0.31, 10.2 + 0.93, $P < 0.001$, for former and current smoker controls, respectively).

Measurement of DNA repair and residual capacity by FLARE assay

To elucidate the underlying mechanisms underlying the differences in response between the lung cancer cases and the current smoker controls, we used the FLARE assay to measure the repair capacity between the 2 groups. The TEM measured by the FLARE assay is the end result of the H₂O₂-induced damage as well as a manifestation of the removal of additional 8-oxoG lesions recognized by the hOGG1 enzyme. Thus the FLARE can be used to measure the residual damage

that has not been removed during the initial one hour recovery after H₂O₂ treatment. Lung cancer cases had over 2-fold higher levels of DNA damage than the current smoker controls (mean ± SD = 44.6 ± 0.89 versus 18.4 ± 1.37, $P < 0.001$). The residual DNA damage among cases and current smoker controls are summarized in Table 3. Comparing cases and current smokers controls revealed significantly lower levels of residual DNA damage in the controls ($P < 0.001$), thus reflecting more efficient repair capacity compared to the cases.

Figure 1 shows the range of calculated DNA repair capacity in response to H₂O₂ treatment (Figure 1A) and H₂O₂ + hOGG1 (Figure 1B) among cases and current smokers controls using the 75th percentile of the controls as the cutoff point. Minimal variation in repair was observed among the cases, with about 90% and 100% of the cases falling into the poor repair group (<20%) in response to H₂O₂ treatment alone and H₂O₂ + hOGG1 treatments, respectively. Only 1% of the cases showed a repair capacity over 80%. In contrast, the current controls had a wide range of repair, with 76% of the control subjects having a repair capacity of >80% in response to the H₂O₂ treatment and H₂O₂ + hOGG1 treatments. Table 2 shows the distribution of residual damage among cases and current controls and determination of

Table 2 Distribution of baseline and H₂O₂-induced DNA damage among cases and controls

Tail Extent Moment Mean ± SE	Controls			Cases	P-value
	Non-Smokers	Former Smokers	Current Smokers	Current Smokers	
Baseline DNA damage	0.26 ± 0.11	0.23 ± 0.09	0.54 ± 0.15	0.57 ± 0.12	<0.001
H ₂ O ₂ - induced DNA damage	3.2 ± 0.82	5.8 ± 0.31	10.2 ± 0.93	19.5 ± 0.41	<0.001

Table 3 Distribution of DNA damage and percent residual DNA damage among cases and current smoker controls

Tail Extent Moment Mean ± SE	Cases	Controls	P-value	Adjusted OR	95% CL
	Current Smokers	Current Smokers			
H ₂ O ₂ + OGG1	44.6 ± 0.89	18.4 ± 1.37	<0.001		
Percent Residual Damage (PRD)	237.4 ± 8.7	180.8 ± 50.7	<0.001		
Residual Damage Logistic Regression (RDLR)					
≤75th N (%)	13(43.3)	41(75.9)			
>75th N (%)	17(56.7)	13(24.1)			
	5.07	1.80-14.24		5.07	1.80-14.24
		Referent			

P-value from t-test; TEM RDLR dichotomized by 75th percentile of the controls. Adjusted by age, sex and family history of a smoking-related cancer

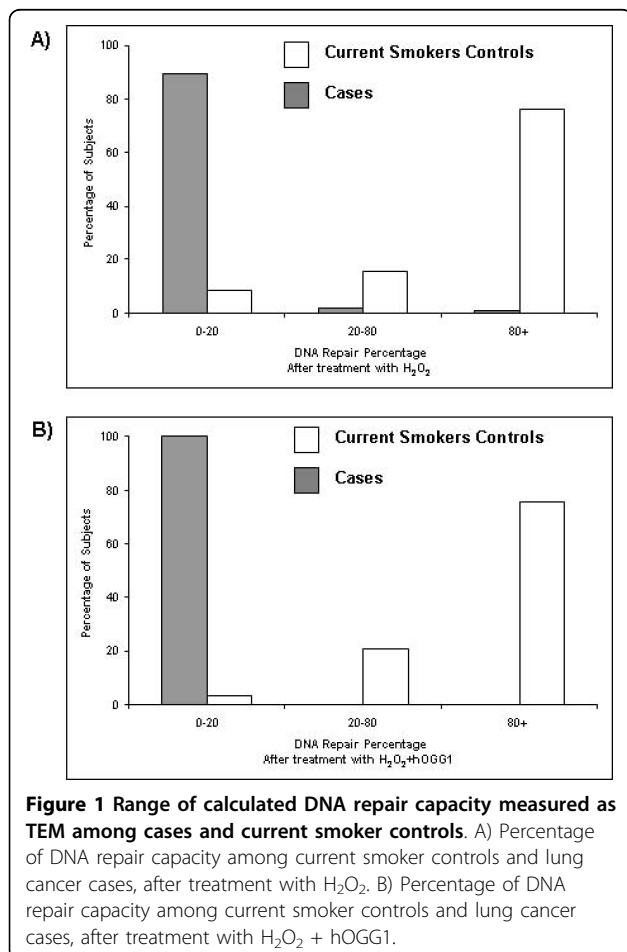
risk for lung cancer. After adjustment for age, gender and family history of a smoking-related cancer using linear regression there was a 5-fold increase in risk of lung cancer associated with high levels of residual damage or reduced repair capacity (OR = 5.07; 97% CL = 1.80-14.24).

Discussion

DNA damage generated by ROS produced as byproducts of cellular metabolism has been proposed as a key factor in mutagenesis and cancer process. Superoxide (O₂⁻), hydrogen peroxide (H₂O₂), and the hydroxyl radical (OH⁻) are mutagens produced as byproducts of normal metabolism in mitochondria and by ionizing radiation [7]. ROS can induce single-strand breaks and several types of DNA base damage, including fragmented, ring-opened forms and oxidized aromatic derivatives. Among many repair pathways, the BER pathway is the most important cellular protection mechanism responding to oxidative DNA damage induced by ROS [5]. Therefore, increased oxidative DNA damage and reduced BER capacity may play an important role in lung carcinogenesis.

OGG1 is a critical component of the BER pathway required for the removal of 8-OHdG lesion from DNA exposed to ROS. The impaired repair activity of OGG1 protein was suggested to be a factor contributing to the high somatic mutation rate in human cells, since the accumulation of 8-OHdG cause mutations by mis-pairing with adenine during replication [11-14]. It has also been reported that lower enzymatic levels of OGG1 in peripheral lymphocytes correlated with an increased risk of lung cancer among smokers [23]. In our current study, we elucidate the association between plasma levels of 8-OHdG and the OGG1 repair capacity among lung cancer cases and controls.

Our results demonstrated that baseline DNA damage is similar among lung cancer cases and current smoker controls and is significantly higher than that observed among the former and non-smokers controls. These findings are consistent with other studies showing higher levels of constitutive DNA damage in the form of spontaneous chromosomal aberrations and elevated endogenous single-stranded breaks [30-32] among patients compared to controls. Results from the baseline serum levels of 8-OHdG followed a similar pattern to



that of the Comet assay where lung cancer cases and current smoker controls had the highest levels of 8-OHdG as compared to the former smokers; however, these differences were not significant. Interestingly, the levels of 8-OHdG were significantly lower among non-smoker controls as compared to the lung cancer cases and current smoker controls. These findings are consistent with studies that have reported that smoking-induced DNA adducts persist for a long time and could still be detectable long after quitting [33,34]. Moreover, numerous reports have associated radicals with smoking-related oxidative stress and DNA damage [23,35,36] since the metabolic activation process leads to the formation of DNA adducts, which are carcinogen metabolites bound covalently to DNA, usually at guanine or adenine. If DNA adducts escape cellular repair mechanisms and persist, they may lead to miscoding, resulting in a permanent mutation. Using a linear regression model we elucidated whether the increase in 8-OHdG levels in cancer patients is related to the increase in residual damage by hOGG1 FLARE among smokers. A significant correlation, ($P < 0.02$) was found between high levels of 8-OHdG levels at baseline and high levels of residual damage.

Lung cancer is a logical disease for evaluating oxidative damage and the role of free radicals because the etiologic agents for lung cancer are tobacco carcinogens that are known to damage DNA. We therefore used H_2O_2 as the ROS-inducing agent in lymphocytes of lung cancer patients and controls in order to measure differential sensitivity (reflected by level of induced DNA damage) and variation in repair capacity. H_2O_2 undergoes a Fe (II)-mediated Fenton reaction in the cell, resulting in the formation of the highly reactive hydroxyl radical and other reactive oxygen species which can induce strand breaks in the DNA [37]. Several studies have shown that the repair of single-strand breaks in normal cells is generally rapid, occurring within one hour from exposure [38,39] which is in agreement with results from our DNA repair time course (Additional file 1, Figure S1B). In our experiments, treatment of the cells with H_2O_2 followed by a one-hour incubation (to allow strand breaks repair) showed that the level of induced DNA damage was significantly different in all four groups studied, with 3.36- and 1.76-fold increases in detected DNA damage in the cancer cases and current smoker controls respectively as compared to former smoker controls. These results indicate that the peripheral lymphocytes from the three comparison groups may have differential sensitivity to H_2O_2 with the lung cancer patients being more sensitive than the current and former smoker controls.

In addition to the increase in sensitivity to H_2O_2 -induced damage, the lung cancer cases may have a poorer

repair capacity than either group of current smoker controls. Several case-control studies have reported that low DNA repair capacity is an independent risk factor for several types of cancers [27,40,41]. Likewise, several studies have shown similar increases in DNA damage and decreases in repair efficiency in response to *in vitro* treatment with chemical mutagens [42-44]. In the current study, H_2O_2 treatment followed by a one-hour incubation with hOGG1 enzyme allowed the quantification of the level of residual 8-oxoG in DNA and determination of the repair capacity in the two groups. Our results showed that current smoker controls had a significantly lower level of residual DNA damage compared to the cases. After adjustment for age, gender and family history of a smoking-related cancer using linear regression there was a 5-fold increase in risk of lung cancer associated with high levels of residual damage/reduced repair capacity (OR = 5.07; 97% CL = 1.80-14.24). Reduced OGG1 activity could be expected to be a risk factor in other smoking-related cancers. However, given the abundance of 8-oxoguanine and the suspected role of oxidative stress in cancer, reduced OGG1 activity might be associated with the risk of some other cancers as well. The excision of 8-oxoguanine residues by OGG1 protects against aberrant adenine-cytosine and guanine-thymine conversions [15] that can lead to heritable mutagenesis, particularly in non-proliferative cells in which lesions accumulate by cell division [45]. OGG1 expression therefore preserves genomic integrity, and it has been shown that *Ogg1*-deficient mice experience enhanced incidences of mutations and tumor formation [46].

Furthermore, our results showed that the current smoker controls had a relatively wide range of repair capacity (Figures 1A and 1B) as compared to the cases, the majority of which fell in the poor repair category. Reduced repair capacity of a particular repair protein may only be revealed when expressed in adverse cellular conditions under oxidative stress. The major risk factor for lung cancer is tobacco smoke and a suboptimal DNA repair could play a crucial role in the susceptibility of the disease. Using the FLARE assay, we were able to identify the small percentage of current smoker controls with poor repair who would be good candidates for intensive follow-up and screening.

Conclusion

In summary, results from this study suggest that high levels of 8-OHdG are correlated with high levels of oxidative DNA residual damage and suboptimal OGG1 repair capacity all of which were predominantly seen in the lung cancer case group. A limitation of our study is the relatively small sample size of the study groups. However, given the striking differences in damage and repair capacities between groups, it is unlikely that the

effect is due to the sample size. Large prospective studies are therefore warranted in which robust functional assays such as the ones used in our study could contribute to the identification of individuals. Such an approach, which may be extended to include additional DNA repair pathways, may provide an effective strategy for the prevention of tobacco-related cancers. The simplicity, rapidity and sensitivity of the FLARE assay make it a valuable tool for screening and possibly for prioritizing potential cases for intensive surveillance.

Additional material

Additional file 1: Supplementary results. Figure S1: A) Mean cell viability in response to different concentrations of H₂O₂. B) Repair over time in a normal cell-line after 60 μM treatment with H₂O₂.

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Authors' contributions

REZ was responsible for conceiving and designing the study as well as analyzing and interpreting the data and writing the different stages of the manuscript. CMM was responsible for the Comet/FLARE experiments and contributed in the data analysis and drafting the manuscript. AC was responsible for the ELISA assays and contributed in the data analysis. MRS and AG were responsible for coordination of the subject recruitment for the study and participated in data interpretation and manuscript preparation. CEJ was responsible for data analysis and contributed to manuscript preparation. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.
Financial competing interests:

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