Inhalative Exposure to Vanadium Pentoxide Causes DNA Damage in Workers: Results of a Multiple End Point Study

Veronika A. Ehrlich,1 Armen K. Nersesyan,2 Christine Hoelzl,1 Franziska Ferk,1 Julia Bichler,1 Eva Valic,1 Andreas Schaffer,3 Rolf Schulte-Hermann,1 Michael Fenech,4 Karl-Heinz Wagner,5 and Siegfried Knasmüller1

1Institute of Cancer Research, Department of Medicine I, Medical University of Vienna, Vienna, Austria; 2Austrian Workers Compensation Board, Vienna, Austria; 3Department of Medicine II, Medical University of Vienna, Austria; 4Commonwealth Scientific and Industrial Research Organisation, Human Nutrition, Adelaide, Australia; 5Department of Nutritional Sciences, University of Vienna, Austria

BACKGROUND: Inhalative exposure to vanadium pentoxide (V2O5) causes lung cancer in rodents.

OBJECTIVE: The aim of the study was to investigate the impact of V2O5 on DNA stability in workers from a V2O5 factory.

METHODS: We determined DNA strand breaks in leukocytes of 52 workers and controls using the alkaline comet assay. We also investigated different parameters of chromosomal instability in lymphocytes of 23 workers and 24 controls using the cytokinesis-block micronucleus (MN) cytome method.

RESULTS: Seven of eight biomarkers were increased in blood cells of the workers, and vanadium plasma concentrations in plasma were 7-fold higher than in the controls (0.31 µg/L). We observed no difference in DNA migration under standard conditions, but we found increased tail lengths due to formation of oxidized purines (7%) and pyrimidines (30%) with lesion-specific enzymes (formamidopyrimidine glycosylase and endonuclease III) in the workers. Bleomycin-induced DNA migration was higher in the exposed group (25%), whereas the repair of bleomycin-induced lesions was reduced. Workers had a 2.5-fold higher MN frequency, and nucleoplasmic bridges (NPBs) and nuclear buds (Nbuds) were increased 7-fold and 3-fold, respectively. Also, apoptosis and necrosis rates were higher, but only the latter parameter reached statistical significance.

CONCLUSIONS: V2O5 causes oxidation of DNA bases, affects DNA repair, and induces formation of MNs, NPBs, and Nbuds in blood cells, suggesting that the workers are at increased risk for cancer and other diseases that are related to DNA instability.

KEY WORDS: comet assay, cytokinesis-block micronucleus assay, DNA damage, occupational exposure, vanadium pentoxide. Environ Health Perspect 116:1689–1693 (2008). doi:10.1289/ehp.11438 available via http://dx.doi.org [Online 31 July 2008]

Vanadium pentoxide (V2O5) is used for the production of metal alloys, for the manufacturing of lithium batteries and high-pressure lamps, and for the synthesis of chemicals [International Agency for Research on Cancer (IARC) 2006]. Its annual production worldwide is in the range of 60,000 tons (IARC 2006; Woolery 1997). Occupational exposure to the oxide occurs at production sites, during processing and refining of vanadium ores and sludges, during manufacturing of vanadium-containing products, in the course of combustion of vanadium-rich fuels, and by handling of catalysts in the chemical industry (Plunkett 1987). Environmental exposure to the metal and its oxides occurs via inhalation in the vicinity of metallurgical plants or through consumption of contaminated foods (Barceloux 1999; IARC 2006). Although foods contain low concentrations, nutrition is the major source of exposure for the general population (Barceloux 1999).

The National Toxicology Program (NTP 2002) found an increase of lung adenomas and carcinomas in mice after inhalative exposure to V2O5; this was paralleled by an increased incidence of hyperplasia in lung tissue. In male rats, the number of tumors was elevated (nonsignificantly), whereas in females no increase was found (IARC 2006; NTP 2002; Resz et al. 2003). These findings led to a reevaluation of the metal oxide (IARC 2006) and to its classification as a group 2B (“possible human”) carcinogen. One of the problems encountered in the risk assessment of V2O5 is the lack of human data. According to IARC (2006), inhalative exposure to V2O5 in vanadium plants exists worldwide, and several hundred workers may be affected; furthermore, workers of other industries may also be exposed. The occupational exposure limit in Austria for V2O5 in air is 0.05 mg/m3 (Bundesminister für Wirtschaft und Arbeit 2003). The Senate Commission of the German Research Foundation [Deutsche Forschungsgemeinschaft (DFG)] decided to suspend the maximum allowed concentration of V2O5 in workplace air because of its suspected carcinogenicity (DFG 2006). In the United States, the National Institute for Occupational Safety and Health and the American Conference of Governmental Industrial Hygienists established an occupational exposure limit of 0.05 mg/m3 air (Occupational Safety and Health Administration 2006). Measurements of air concentrations in vanadium factories yielded values in the range of 0.02–5 mg/m3 (IARC 2006). Results of in vitro and animal studies indicate that the oxide causes formation of reactive oxygen species (Ingram et al. 2003, 2007; Wang et al. 2003; Zhang et al. 2001) and aneugenic effects (Migliore et al. 1993; Ramirez et al. 1997; Zhong et al. 1994) and interferes with DNA synthesis and repair (IARC 2006). Because DNA damage and aneugenic processes are known to play a role in the onset of human cancer (Duesberg et al. 2005; Pitot 1986), evidence of genetic damage in exposed humans would support the assumption of increased cancer risks. At present, only one study on the influence of occupational exposure to V2O5 on DNA stability has been published; in that study, Ivancsits et al. (2002) investigated DNA migration in leukocytes using the standard single-cell gel electrophoresis (SCGE) assay. The authors observed no indication of damage and found no elevation in the frequencies of sister chromatid exchanges (SCEs) or the concentration of 8-hydroxy-2’-deoxyguanosine in leukocytes. Lener et al. (1998) found no SCEs or chromosomal aberrations in blood cells of children living in the vicinity of a vanadium production site.

Our goal in the present study was to comprehensively evaluate the impact of inhalative V2O5 exposure on genetic stability. We monitored DNA migration in leukocytes of workers and matched controls with the standard SCGE assay, and we monitored oxidized bases using lesion-specific enzymes (Collins et al. 1993). Furthermore, we measured the sensitivity toward bleomycin (BLEO) and the repair of lesions induced by this cytostatic agent (Rajaee-Bebbahani et al. 2001; Schmezer et al. 2001; Wei et al. 2005). BLEO sensitivity was initially monitored in chromosomal aberration assays (Hsu et al. 1989; Szekely et al. 2003) and later in SCGE experiments (Schmezer et al. 2001).

Additionally, we conducted cytokinesis-block micronucleus cytome (CBMN Cyt) assays with lymphocytes. This test is widely used for the detection of DNA damage in...
humans (Fenech 2007). Micronuclei (MNs), which are formed as a consequence of chromosome breakage and/or aneuploidy (Fenech and Morley 1985), correlate with the incidence of cancer in humans (Bonassi et al. 2007). Also, we evaluated the frequencies of nucleoplasmic bridges (NPBs) and nuclear buds (Nbuds) in lymphocytes. NPBs are assumed to occur when centromeres of dicentric chromosomes are pulled to the opposite poles of the cell at anaphase and provide a measure of chromosome rearrangements (Fenech 2006). Therefore, NPBs give direct evidence of genome damage resulting from misrepaired DNA breaks, which cannot be detected when MNs are scored as the only end point. Nbuds form as a consequence of gene amplification (Fenech 2006). Amplified DNA is selectively localized at specific sites of the nucleus and eliminated through recombinogenic events during the S-phase of mitosis (Shimizu et al. 1998, 2000).

Other parameters included in the present study were necrosis and apoptosis, and the nuclear division indices (Fenech 2006). We measured plasma concentrations of folate and vitamins B6 and B12 in both groups, because deficiencies of these micronutrients may increase MN levels (Fenech et al. 1997). Furthermore, we determined the plasma vanadium levels of the participants.

Materials and Methods

Study groups. We used the SCGE assay to study DNA migration in whole-blood leukocytes from 52 vanadium production workers exposed to V2O5 by inhalation and from 52 nonexposed controls (jail wardens). Additionally, we analyzed lymphocytes of 24 workers and 23 controls using CBMN Cyt experiments. We collected data concerning age, weight, height, and smoking status with a questionnaire (Table 1).

Workers are exposed to vanadium dust during the entire shift (8 hr) and are required to wear protective masks and gloves. The study was approved by the Ethics Committee of the Medical University of Vienna. After obtaining informed consent, we collected blood (2 × 10 mL) in heparin and EDTA tubes (Vacutainer-Systems, Becton Dickinson, Plymouth, UK). We stored blood samples at 4°C and transported them to the Institute of Cancer Research at the Medical University of Vienna. Whole blood was centrifuged at 623 × g for 10 min at 14°C (Sigma Laboratory Centrifuge 4K15; Sigma Chemical Co., St. Louis, MO, USA) and plasma was collected, aliquoted, and stored at −80°C. We conducted experiments between October 2004 and May 2005.

Exposure assessment. We determined vanadium concentrations in plasma using graphite furnace atomic absorption spectrometry, as described by Ivancsits et al. (2002), with Zeeman background correction at a temperature of 2,450°C. The calibration curves for vanadium were in the range of 0–40 µg/L, and the detection limit was 0.3 µg/L (5.9 nmol/L).

Measurements of vitamins B6 and B12 and folate. We determined pyridoxal phosphate (the active form of vitamin B6) in plasma by HPLC using commercial assays (ImmunoLogic Diagnostics, Bensheim, Germany) as described by Majchrzak et al. (2006). We measured vitamin B12 and folate in plasma using commercial radioimmunoassays (IUL Instruments GmbH, Königswinter, Germany) (Majchrzak et al. 2006).

Leukocyte isolation and BLEO treatment for SCGE assays. We used the protocol of Buschini et al. (2002) for leukocyte isolation: EDTA-anticoagulated blood was maintained in erythrocyte-lysis buffer (155 mM NH4Cl, 5 mM KHCO3, 0.005 mM Na2EDTA, pH 7.4) at 37°C for 5 min; centrifuged (200 × g, 5 min at 4°C), washed twice in phosphate-buffered saline (PBS; pH 7.4); and resuspended in RPMI 1640 (Sigma-Aldrich Chemie GmbH, Munich, Germany) without serum (Buschini et al. 2002). We used a modified version of the protocol for BLEO sensitivity and DNA repair measurements (Schmezer et al. 2001). Leukocytes were treated with 10 µg/mL BLEO sulfate (Nippon Kayaku Co. Ltd., Tokyo, Japan) at 37°C for 30 min. To terminate the treatment, cells were washed twice in PBS (pH 7.4). To measure DNA repair capacity, we incubated a second batch of BLEO-treated cells for 15 min (37°C) before lysis and electrophoresis. Cell viability was monitored with trypan blue (Lindl and Bauer 1994). Viability of all samples was ≥ 90%.

Alkaline SCGE (comet) assay. We performed the SCGE assay according to the guidelines of Tice et al. (2000). Briefly, cells pellets were mixed with 60 µL 0.5% low-melting-point agarose (Invitrogen Life Technologies Ltd., Paisley, Scotland) transferred to precoated (1.5% normal-melting-point agarose) glass slides, and sealed with a coverslip. The slides were placed on ice for 5 min to allow solidification of the agarose. After removing the coverslip, we immersed the slides in lysis solution (2.5 M NaCl, 100 mM Na2EDTA, 10 mM Tris, 1% Triton X, 10% dimethyl sulfoxide, pH 10.0) and kept them at 4°C for ≥ 1 hr. To prevent DNA damage, lysis and all subsequent steps were conducted under red light. After lysis, we placed the slides on a horizontal gel electrophoresis unit (C.B.S. Scientific, Solana Beach, CA, USA) and allowed the DNA to unwind at 4°C in alkaline electrophoresis buffer (300 mM NaOH, 1 mM Na2EDTA, pH ≥ 12.5) for 20 min. Electrophoresis was performed at 25 V and 300 mA for 20 min; then the slides were covered with neutralization buffer (0.4 M Tris base, pH 7.5, 4°C) for 2 × 8 min and air dried. The coded slides were stained with ethidium bromide (20 µg/mL) and evaluated under a fluorescence microscope (Nikon 623012; Nikon, Tokyo, Japan). We used an automated analysis system (Helma and Uhl 2000) to acquire images, compute the integrated intensity profile for each cell, estimate the comet cell components, and evaluate derived parameters. For each experimental end point we analyzed three cultures, and we measured 50 randomly captured cells from each slide. To quantify DNA damage, tail lengths and tail moments were determined.

Determination of oxidized purines and pyrimidines. We used a modified version of the protocol published by Collins et al. (1993) to measure endogenous formation of oxidized DNA bases. Formamidopyrimidine glycosylase (FPG) and endonuclease III (ENDO III) were provided by K. Angelis (Czech Academy of Sciences, Prague, Czech Republic).

After lysis of the cells, the slides were washed in enzyme buffer (0.1 M KCl, 40 mM HEPES, 0.5 mM Na2EDTA, 0.2 mg/mL bovine serum albumin, 4°C, pH 8) for 2 × 8 min. Subsequently, agarose-embedded cells were covered with 50 µL enzyme buffer or with enzyme solutions (1.0 µg/mL). Gels were sealed with a coverslip and incubated at 37°C in the dark for 45 min (ENDO III) or 30 min (FPG). Subsequently, the slides were transferred into alkaline buffer for unwinding, and electrophoresis was performed. CBMN Cyt experiments. We isolated lymphocytes according to the protocol of Fenech (2000). Briefly, Histopaque 1077 (Sigma) was overlaid with RPMI 1640–diluted blood and centrifuged at 318 × g at 14°C for 30 min. Subsequently, the cell layer was removed, resuspended in RPMI, washed twice, and centrifuged (318 × g, 14°C, 10 min).

### Table 1. Distribution of selected characteristics in exposed subjects and controls.

| Variable               | Controls (n = 52) | Exposed (n = 52) | p-Value |
|------------------------|------------------|-----------------|---------|
| Age (years)            | 38.0 (32.00–44.50) | 43.0 (38.50–49.50) | > 0.05  |
| Body mass index        | 27.0 (24.00–29.50) | 26.0 (25.00–27.50) | > 0.05  |
| Cigarettes/day (no.)   | 5.0 (0.00–17.00)  | 7.0 (0.00–21.00)  | > 0.05  |
| Vanadium (µg/L)        | 0.3 (0.24–0.39)   | 2.2 (1.54–3.89)   | < 0.0001|
| Folate (µg/L)          | 4.7 (3.04–6.40)   | 3.4 (2.45–4.205)  | > 0.05  |
| Vitamin B6 (ng/mL)     | 18.2 (14.38–21.55) | 19.8 (14.06–36.94) | 0.016   |
| Vitamin B12 (ng/L)     | 665.0 (428.5–866.5) | 968.0 (662.5–1215.0) | 0.013   |

All study subjects were male Caucasians and nonvegetarians. Values shown are median (25th–75th percentiles).
We performed the CBMN Cyt test using the cytochalasin B technique described by Fenech (2007). We determined MNs, Nbuds, and NPBs, as well as apoptotic and necrotic cells, in samples from 24 workers and 23 controls using the cytochrome approach (Fenech 2007). Slides were evaluated by one of the authors (V.A.E.) who was trained in the laboratory of M.F. at Commonwealth Scientific and Industrial Research Organisation (Human Nutrition, Adelaide, South Australia), which has developed the CBMN Cyt assay and was centrally involved in the standardization of scoring criteria via the International Collaborative Project on Micronucleus Frequency in Human Populations (Human Micronucleus Project 2008). For each participant, we prepared lymphocyte cultures in duplicate. Each culture contained 10^6 cells in 750 µL RPMI 1640 supplemented with 100 µU/mL penicillin, 100 µg/mL streptomycin, 10% fetal bovine serum (Sigma), 2.0 µMol/L l-glutamine, and 30 µg/mL phytomitomagglutinin (Invitrogen, Carlsbad, CA, USA). Cultures were incubated at 37°C in a humidified atmosphere containing 5% CO2. Forty-four hours after mitogen stimulation, we added cytochalasin B (final concentration, 4.0 µg/mL; Sigma-Aldrich, St. Louis, MO, USA) to block cell division. The stimulation, we added cytochalasin B (final concentration, 4.0 µg/mL; Sigma-Aldrich, St. Louis, MO, USA) to block cell division. The total incubation time of the cultures was 72 hr. Subsequently, the cells were harvested and used to prepare slides, which were air dried, fixed, and stained with Diff Quick (Dade Behring, Deerfield, IL, USA). From each participant, we scored 2,000 binucleated cells using a light optical microscope (Microphot-FXA; Nikon) (Fenech et al. 2003).

**Statistical analysis.** We performed statistical analyses using Statistica 5.0 software (StatSoft Inc., Tulsa, OK, USA). The results are presented as medians and 25th–75th percentiles. All p-values were two-tailed, and we considered differences to be significant at p ≤ 0.05. We used the Mann-Whitney U-test for comparisons between exposed subjects and controls. We used Spearman’s correlation coefficients to test univariate relationships between different variables.

**Results**

**Demographics of the study populations.** The characteristics of participants are summarized in Table 1. Age and body mass index did not differ between the groups. All individuals were males and nonvegetarians. The control group included 11 smokers and the exposed group included 12. The concentrations of folate were similar in both groups, whereas levels of vitamins B6 and B12 were higher (8.1% and 31%, respectively) in the exposed group, possibly due to intake of supplements. Vanadium levels were 7-fold higher in the plasma of the exposed group.

**DNA migration in leukocytes.** Table 2 summarizes the results of the SCGE experiments. We found no difference of DNA migration between exposed individuals and control subjects when the assay was carried out under standard conditions (which reflect formation of single- and double-strand breaks), but we observed increased tail lengths and tail moments in all other end points in the workers. Formation of oxidized purines (detected with FPG) was elevated by 7%, and the formation of pyrimidines (by ENDO III treatment) was enhanced by 33%. Furthermore, the sensitivity of the cells toward BEO-induced DNA damage was higher (25%) in the exposed individuals. When we monitored DNA migration after a repair phase (15 min incubation after BEO treatment), we observed a decrease of tail lengths (50%) and tail moments (57%) in the controls, whereas in the workers we observed no reduction of tail lengths and a less pronounced decrease of tail moments (30%).

**Frequencies of end points measured with the CBMN Cyt assay.** Data on the frequencies of MNs, NPBs, Nbuds, apoptosis, and necrosis rates, as well as the division indices [nuclear division index (NDI) and nuclear division cytoxicity index (NDCI)] are summarized in Table 2. The number of MNs was 2.5-fold higher in the workers, and the numbers of NPBs and Nbuds were substantially increased (7-fold and 3-fold, respectively).

To find out whether V2O5 exposure and smoking cause synergistic effects, we compared the frequencies of MNs and Nbuds in smokers and nonsmokers to those in the corresponding controls. In nonexposed individuals, the levels (means ± SD) of MNs, NPBs, and NPBs were similar in smokers and nonsmokers respectively (MNs, 1.77 ± 1.62 and 1.50 ± 1.86; NPBs, 0.33 ± 0.39 and 0.36 ± 0.32; NPBs, 0.46 ± 0.81 and 0.45 ± 0.61). In workers, the MN rates were lower in smokers (4.17 ± 2.95 and 8.08 ± 4.54), whereas the frequencies of NPBs and Nbuds were more or less identical (NPBs, 6.83 ± 3.04 and 7.35 ± 2.50; Nbuds, 3.58 ± 2.76 and 3.30 ± 1.41).

The frequencies of necrotic and apoptotic cells were elevated in the workers by 55% and 50%, respectively; for the latter parameter the difference was not significant. The last two rows of Table 3 show the nuclear division indices. The division rates were similar in both groups, regardless of whether we considered the number of viable cells only (NDI) or also dead cells (NDCI). These results indicate that metal exposure has no significant impact on the proliferative capacity of viable or total lymphocytes, respectively.

**Discussion and Conclusions**

In this article we present the results of the first comprehensive study on the impact of occupational exposure to airborne V2O5 on DNA stability. For seven of eight end points, we found significant differences between exposed workers and controls.

The most important observation is the 2.5-fold higher frequency of MNs in lymphocytes of the exposed individuals (Table 3), because a prospective study showed that MNs in peripheral blood lymphocytes are a valid

### Table 2. DNA migration in leukocytes of workers and unexposed subjects using different end points of the comet assay.

| Test condition | Measure | Controls (n = 52) | Exposed (n = 52) | Δ (%) | p-Value |
|---------------|---------|------------------|-----------------|-------|---------|
| Standard conditions | TL | 4.3 (4.07–4.64) | 4.1 (3.89–4.44) | -5 | > 0.05 |
| | TM | 2.4 (2.27–2.79) | 2.3 (2.16–2.58) | -4 | > 0.05 |
| | FPG | 4.5 (4.21–4.80) | 4.8 (4.14–5.53) | +7 | 0.0236 |
| | ENDO III | 2.5 (2.34–2.62) | 2.7 (2.27–3.24) | +6 | 0.0007 |
| | TM | 7.3 (6.71–8.17) | 9.7 (7.84–12.08) | +33 | 0.0019 |
| | Tim 3.5 (3.24–3.89) | 3.8 (3.18–6.03) | +9 | 0.0023 |
| | BEO | TL | 13.2 (11.93–15.58) | 16.5 (13.55–23.56) | +25 | < 0.0001 |
| | TM | 8.7 (7.55–11.72) | 12.2 (10.07–19.58) | +40 | < 0.0001 |
| BEO + DNA repair | TL | 6.6 (6.02–7.20) | 8.08 (7.08–9.08) | +200 | < 0.0001 |
| | TM | 3.7 (3.41–4.20) | 8.3 (7.10–11.39) | +24 | < 0.0001 |

Abbreviations: Δ, difference between exposed subjects and controls; TL, tail length (µm); TM, tail moment. Values shown are median (25th–75th percentiles).

**Table 3. Frequencies of micronucleated lymphocytes and total numbers of MNs, NPBs, Nbuds, and apoptotic and necrotic cells per 2,000 binucleated cells in workers and unexposed subjects.**

| End point | Controls (n = 23) | Exposed (n = 24) | Δ (%) | p-Value |
|-----------|------------------|-----------------|-------|---------|
| Total no. of MNs | 2.0 (1.00–4.00) | 5.0 (2.50–9.00) | +150 | 0.0132 |
| Micronucleated cells | 2.0 (1.00–4.00) | 5.0 (2.50–8.00) | +150 | 0.0132 |
| NPBs | 1.0 (0.00–2.00) | 7.0 (5.00–9.00) | +600 | < 0.0001 |
| Nbuds | 1.0 (0.00–1.00) | 3.0 (2.00–5.00) | +200 | < 0.0001 |
| Necrotic cells (%) | 13.5 (10.7–16.5) | 20.9 (19.4–25.9) | +55 | < 0.0001 |
| Apoptotic cells (%) | 3.0 (1.1–10.7) | 4.5 (3.6–5.2) | +50 | > 0.05 |
| NDI | 1.9 (1.81–1.94) | 1.9 (1.86–1.96) | 0 | > 0.05 |
| NDCI | 1.7 (1.64–1.76) | 1.7 (1.61–1.70) | 0 | > 0.05 |

Δ, difference between exposed subjects and controls. Values shown are median (25th–75th percentiles).

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biomarker for predicting an increased cancer risk in humans (Bonassi et al. 2007). Furthermore, a smaller study showed strong predictability of the MN biomarker in lymphocytes for cardiovascular disease risk (Murgia et al. 2007). Because the levels of folate and vitamins \( B_6 \) and \( B_12 \) were equal or higher in the workers (Table 1), we can exclude that the increased MN levels in this group are due to deficiencies of these micronutrients.

\( V_2O_5 \) and other vanadium compounds induce MNs in bone marrow cells of rodents (Leopardi et al. 2005; Sun 1996). Furthermore, DNA-damaging properties of vanadium compounds have also been found in human and animal cells in vitro (Ivancsits et al. 2002; Kleinsasser et al. 2003; Migliore et al. 1993; Ramirez et al. 1997; Rojas et al. 1996; Roldan and Altamirano 1990; Zhong et al. 1994).

In the present study, we observed an increase of other end points of chromosomal stability (NPs and Nbdus) in the workers. The induction of NPs and Nbdus was even more pronounced than the increase in MN levels (Table 3), and it is notable that a recent case–control study on smokers showed that elevated NPB and Nbd frequency are more strongly associated with lung cancer risk than are MN rates (El-Zein et al. 2006).

We found no synergistic effect between smoking and \( V_2O_5 \) exposure in the present study. Bonassi et al. (2003) conducted a meta-analysis concerning the impact of smoking on MN frequencies among subjects in occupational and environmental surveys. They concluded that only nonexposed heavy smokers exhibit a significant increase of MNs, whereas the MN frequency was not influenced by smoking among individuals exposed to genotoxic agents. In the latter group, even slightly reduced MN levels were found in smokers compared with nonsmokers (Bonassi et al. 2003).

The only marker that was not elevated in the workers in the present study was DNA migration monitored in SCGE experiments under standard conditions, which reflect endogenous DNA damage such as single- and double-strand breaks and apurinic sites (Tice et al. 2000). This observation is in agreement with results of an earlier investigation (Ivancsits et al. 2002).

In vitro studies have shown that the metal oxide induces DNA migration in blood cells under standard conditions, but the doses required to cause effects were substantially higher than the plasma concentrations in the workers in the present study (Ivancsits et al. 2002; Kleinsasser et al. 2003; Rojas et al. 1996). Furthermore, results obtained in SCGE experiments with rodents indicate that vanadium compounds cause DNA migration in inner organs (Altamirano-Lozano et al. 1996, 1999; Leopardi et al. 2005; Villani et al. 2007).

It is notable that the basal levels for tail lengths are quite low in the present study. Our values are in a range similar to those found in other laboratories (Grossi et al. 2008; Undeger and Basaran 2005; Yoshida et al. 2006; Zhang et al. 2008) and also within the historical range of our laboratory (Bichler et al. 2007; Hoelzl et al. 2008; Steinkellner et al. 2005). Because we monitored the cell viability before electrophoresis, we can exclude that cell damage accounts for the low values. Furthermore, the results obtained with tail moment in the present study are essentially identical to those from tail length measurements.

Endogenous formation of oxidized DNA bases (shown by treatment with the repair endonucleases FPG and ENDO III) is higher in the exposed group. Earlier investigations found that vanadium-treated cells generate hydrogen peroxide and superoxide radicals (Huang et al. 2000; Shi and Dalal 1992; Ye et al. 1999; Zhang et al. 2001). These vanadium-induced radicals were shown to cause damage to macromolecules and lipid peroxidation (Donaldson et al. 1985), and it is conceivable that they account for the oxidative damage that we observed in the present study.

It is noteworthy that a number of human studies found increased sensitivity toward DNA damage by BLEO in individuals who are at increased risk for different types of cancer (Schmezer et al. 2001). In the present study, we found a strong difference between workers and controls after BLEO treatment and a 15-min repair phase (Table 2). Although the tail lengths decreased by 50% in the controls, we saw no reduction in workers after the repair phase, and the tail moments were decreased to a higher extent in the controls. This indicates that the metal oxide interacts detrimentally with DNA repair processes. Ivancsits et al. (2002) found severe inhibition of BLEO-induced repair by \( V_2O_5 \) in SCGE experiments with lymphocytes in vitro, whereas the repair of ultraviolet C–induced lesions was less affected. These findings can be taken as an indication that \( V_2O_5 \) inhibits pathways that are required for the repair of BLEO-induced lesions (homologous recombination repair, nonhomologous end joining, and base excision repair) (Schlad-Bartuski et al. 2002; Schmezer et al. 2001; Wei et al. 2005) and, to a lesser extent, nucleotide excision repair, which is required to repair ultraviolet C–induced lesions (Wei et al. 2005).

As shown in Table 3, we found that the frequencies of necrotic cells were higher in the workers. Also, programed cell death (apoptosis) was increased, but this effect did not reach significance. Induction of apoptosis by vanadium compounds has also been observed in earlier in vitro studies (Au et al. 2006; Huang et al. 2000; Lampronti et al. 2005; Ray et al. 2006; Wang et al. 2003; Ye et al. 1999), and it has been postulated that activation of mitogen-activated protein kinases by reactive oxygen species and/or by an oxidant-independent pathway may play a role (Chien et al. 2006; Choi et al. 2003; Huang et al. 2000; Luo et al. 2003).

Statistical analyses of the different end points showed no correlations between formation of oxidized purines and pyrimidines and MN induction \((p = 0.3992 \text{ and } 0.4679, \text{ respectively})\), which indicates that different mechanisms are involved. Although oxidative damage of DNA bases caused by \( V_2O_5 \) is probably due to release of reactive oxygen species, the formation of MNs may be due to its aneugenic properties, which have been found in in vitro studies (Galli et al. 1991; Roldan and Altamirano 1990; Zhong et al. 1994) and are apparently caused by disturbances of microtubule assembly (Ramirez et al. 1997).

We also failed to find correlations between vanadium plasma levels and formation of oxidized purines \((p = 0.2340)\), pyrimidines \((p = 0.2895)\), and MN frequencies \((p = 0.1571)\). Earlier studies with workers exposed to metals other than vanadium indicate that polymorphisms in repair genes have a strong impact on MN formation (Iarmarccovai et al. 2006; Mateuca et al. 2005, 2008), and it is conceivable that they also play a role in the case of the effects caused by \( V_2O_5 \).

Overall, our results show that inhalative exposure to \( V_2O_5 \) increases the levels of oxidized bases and of MN, NPB, and Nbd frequencies in blood cells and affects their DNA repair capacity. It is notable that vanadium levels similar to or even higher than those found in our study have been detected in workers of other vanadium industries and in welders (Altamirano-Lozano et al. 1999; Huang et al. 2000; Ivancsits et al. 2002; Shi and Dalal 1992; Villani et al. 2007). Because the aforementioned parameters are causally related to diseases including cancer, our findings strongly suggest that more protective measures and periodical monitoring of the workers are required. Furthermore, the current exposure levels should be reduced to avoid health risks due to vanadium-induced DNA instability.

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