

Genotoxic and carcinogenic risks associated with the dietary consumption of repeatedly heated coconut oil

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Repeated heating of vegetable oils at high temperatures during cooking is a very common cooking practice. Repeated heating of edible oils can generate a number of compounds, including polycyclic aromatic hydrocarbons (PAH), some of which have been reported to have carcinogenic potential. Consumption of these repeatedly heated oils can pose a serious health hazard. The objectives of the present study were to evaluate the genotoxic and carcinogenic risks associated with the consumption of repeatedly heated coconut oil (RCO), which is one of the commonly consumed cooking and frying medium. The PAH were analysed using HPLC in fresh CO, single-heated CO (SCO) and RCO. Results revealed the presence of certain PAH, known to possess carcinogenic potential, in RCO when compared with SCO. Oral intake of RCO in Wistar rats resulted in a significant induction of aberrant cells ($P < 0.05$) and micronuclei ($P < 0.05$) in a dose-dependent manner. Oxidative stress analysis showed a significant ($P < 0.05$) decrease in the levels of antioxidant enzymes such as superoxide dismutase and catalase with a concurrent increase in reactive oxygen species and lipid peroxidation in the liver. In addition, RCO given alone and along with diethylnitrosamine for 12 weeks induced altered hepatic foci as noticed by alteration in positive (γ -glutamyl transpeptidase and glutathione-S-transferase) and negative (adenosine triphosphatase, alkaline phosphatase and glucose-6-phosphatase) hepatospecific biomarkers. A significant decrease in the relative and absolute hepatic weight of RCO-supplemented rats was recorded ($P < 0.05$). In conclusion, dietary consumption of RCO can cause a genotoxic and preneoplastic change in the liver.

Coconut oil: Polyaromatic hydrocarbons: Altered hepatic foci: Genotoxicity

Nutrition and diet as environmental factors and determinants of growth and body composition can contribute to the risk of human cancers^(1–3). Evidence pertaining to the role of dietary factors in carcinogenesis comes from both epidemiological studies and laboratory experiments. Dietary fat has been known to cause cancer of different organs⁽⁴⁾. Vegetable oils are a major source of fat intake, and have been used as a common medium for cooking since ages. The practice of repeated heating of edible oils is associated with adverse health effects⁽⁵⁾. High-temperature cooking techniques such as grilling and pan frying are associated with increased risk of carcinogenicity⁽⁶⁾ and genotoxicity^(7,8). Repeatedly heated fat possesses mutagenic potential⁽⁹⁾, and increase in mutagenicity is proportionally related to its frying time⁽¹⁰⁾. A number of genotoxic responses caused by the dietary use of heated cooking oils have also been reported^(7,11). Studies from different sources have indicated that the heating of the vegetable oils at high temperatures leads to the formation of compounds having the potential to cause cancers^(8,12,13). The heat processing of the oils at high temperatures generates food

contaminants such as polycyclic aromatic hydrocarbons (PAH), a family of toxic and mild to potent carcinogenic chemicals^(14,15). The epidemiological study done by Lopez-Abente *et al.*⁽¹⁶⁾ suggested that individuals exposed to PAH are at the risk of developing cancers. Diet is the major non-occupational source of PAH for non-smokers⁽¹⁷⁾, with meat, meat products, cereals, oils and fats being the principal sources⁽¹⁸⁾.

Epidemiological studies have indicated that a high intake of saturated fat increases the risk of cancers⁽¹⁹⁾. Coconut oil (CO), also known as coconut butter, is a tropical oil extracted from copra (the dried inner flesh of coconuts) of *Cocos nucifera* (Family: Arecaceae), which is used as a cooking medium mainly in the southern parts of India and other South-East Asian countries. The CO contains predominantly medium-chain TAG, with 86.5% SFA, 5.8% MUFA and 1.8% PUFA. The SFA of CO could have an effect on atherosclerosis not only beyond their role in affecting plasma lipoproteins, but also through increased production of inflammatory cytokines in the arterial walls⁽²⁰⁾. High-saturated fat

Abbreviations: AHF, altered hepatic foci; AlkPase, alkaline phosphatase; ATPase, adenosine triphosphatase; B(a)P, benzo(a)pyrene; CO, coconut oil; DEN, diethylnitrosamine; FCO, fresh CO; G6Pase, glucose-6-phosphatase; GGT, γ -glutamyl transpeptidase; Gr, groups; GST-P, glutathione-S-transferase placental type; LPO, lipid peroxidation; PAH, polycyclic aromatic hydrocarbons; RCO, repeatedly heated CO; ROS, reactive oxygen species; SCO, single-heated CO.

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diet is related to oxidative stress⁽²¹⁾, and has a promoting effect on pancreatic carcinogenesis in azaserine-treated rats⁽²²⁾.

In order to delineate the impact of repeatedly heated CO (RCO) use on genotoxicity and/or carcinogenicity, it is important to determine the presence of toxic chemicals which are ingested through repeatedly heated saturated fat-rich diets. Thus, the present study has been designed to investigate the presence of PAH mixture in RCO using HPLC. Furthermore, the potential of RCO consumption to induce genotoxicity, oxidative stress and carcinogenicity in Wistar rats was also investigated.

Materials and methods

Chemicals

Dimethyl sulphoxide, *n*-hexane, cyclohexane, anhydrous sodium sulphate, dichloromethane, acetonitrile and silica gel were of analytical grade and purity (purchased from Merck Chemical Company, Mumbai, India). PAH standard mix was purchased from Supelco (Bellefonte, PA, USA). Diethylnitrosamine (DEN), 2-acetylaminofluorene, benzo(*a*)pyrene (B(*a*)P), γ -glutamyl-4-methoxy- β -naphthylamide, glycylglycine, fast blue BB salt (4-amino-2,5-diethoxybenzanilide diazotated zinc double salt, 4-benzoylamino-2,5-diethoxybenzenediazonion chloride (zinc chloride) salt), ATP disodium salt, sodium- β -glycerophosphate, sodium diethylbarbiturate, glucose-6-phosphate monosodium salt, B(*a*)P, colchicine, dichlorodihydrofluorescein-diacetate dye, phenazine methosulphate, 2-thiobarbituric acid, 1,1,3,3-tetra-methoxy propane, nitroblue tetrazolium, NADPH and NADH were purchased from Sigma Chemical Company (St Louis, MO, USA). Anti-glutathione-S-transferase placental type (Anti-GST-P) and goat anti-rabbit IgG peroxidase secondary antibodies were procured from Calbiochem (Darmstadt, Germany). CO (refined) used in the study was purchased from the local market (Meghdoot Oil Mills Private Limited, Mumbai, India). The rest of the chemicals that were used in the study were of analytical grade and purity, and were procured locally.

Repeatedly heated coconut oil sample preparation

The refined CO was heated (>300°C) above its smoke point (232°C; 450°F) for 30 min, and was cooled to room temperature. For RCO, the process was repeated six times, and the oil sample became viscous dark brown in colour after being repeatedly heated.

HPLC of coconut oil

Extraction and clean-up procedure for PAH was carried out using fresh CO (FCO), single-heated CO (SCO) and RCO according to the protocol of Pandey *et al.* with slight modifications. Briefly, for the extraction of PAH, 10 g of oil were added to 20 ml of *n*-hexane and extracted thrice with dimethyl sulphoxide (10 ml) followed by the addition of cold distilled water (4°C) slowly, and re-extraction was done thrice with cyclohexane (solvent was changed for specific extraction of PAH). After washing, it was concentrated and passed through a glass column (20 × 2.2 cm) containing silica gel and anhydrous sodium sulphate. The elution was carried out with a mixture of cyclohexane and dichloromethane,

and the mixture was evaporated to dryness. The final residue was dissolved in 1 ml of acetonitrile, and passed through a 0.2 μ m filter.

Quantification of the extracted PAH was carried out using an HPLC instrument (Water Associates, Inc., Milford, MA, USA) equipped with a dual pump and Rheodyne injector with a 20 μ l loop. The reversed-phase column used for the analysis was C-18 (E. Merk, Darmstadt, Germany) with a pre-column of the same type. The column was eluted at an ambient temperature (27°C) with 70:30 acetonitrile in water as the solvent at a flow rate of 1.5 ml/min, which was monitored on an UV detector at λ 254 nm. The chromatogram was recorded and processed using Waters Millennium Software (Scientific Equipment Source, Pickering, ON, Canada). The sixteen PAH selection criteria were based on (i) the number of benzene rings in PAH, (ii) PAH generally found in oils following the literature search and (iii) resolution of PAH on HPLC. The peaks of the mixture of PAH in oils were identified by comparing the retention time with that of the standard by using an UV detector for the respective fractions. Quantification of PAH was performed by comparing the integrated peak area with that of the standards⁽¹⁵⁾.

Animals and treatments

Male Wistar rats (6 weeks of age) were procured from the Indian institute of toxicology research (Lucknow, India) animal breeding colony, maintained under standard conditions and fed *ad libitum* a synthetic pellet basal diet (Ashirwad, Chandigarh, India) and water. Institutional guidelines for the care and use of animals were followed, and the ethical approval for the experiments was obtained from the institutional animal ethical clearance committee, Indian Institute of Technology Roorkee (IITR). All experimental procedures involving animals were approved by the animal ethical clearance committee, IITR.

Chromosomal aberration and micronuclei induction assays

Rats were randomly divided into two sets of eight groups (Gr) having six rats each. In Gr I, no treatment was given to the rats, whereas in Gr II, B(*a*)P (100 mg/kg body weight; intraperitoneally) was administered to the rats 24 h before killing. Each rat was administered FCO (0.5 ml) in Gr III, SCO in Gr IV (IVa: 0.25 ml and IVb: 0.5 ml), and RCO in Gr V (Va: 0.1 ml, Vb: 0.25 ml and Vc: 0.5 ml) consecutively by gavage. At the end of 1-week experimental period, rats were killed by cervical dislocation, and bone marrow was collected from their femurs.

Oxidative stress assay

Rats were divided equally into five Gr consisting of ten rats each, where Gr I, II and III rats were given treatments that were similar to those used for the genotoxic assays. SCO (0.5 ml/rat) and RCO (0.5 ml/rat) were given by gavage to other Gr, i.e. Gr IV and V, respectively. After 1 week, rats were killed, and their livers were excised and were homogenised separately in ice-cold PBS (pH 7.4) for enzymatic estimations and reactive oxygen species (ROS) determination.

Medium-term bioassay

A total of seventy-two rats were divided into six Gr. Gr I rats were fed a normal basal diet during the entire study period, i.e. 12 weeks, and they served as negative controls. Gr II, III, IV and V rats were given a single dose of DEN (200 mg/kg body weight) intraperitoneally. After 1 week of recovery period, rats of Gr II were fed 0.05% 2-acetylaminofluorene in a crushed diet, and Gr III, IV and V rats were administered 0.5 ml/rat of FCO, SCO and RCO, respectively, by gavage. Simultaneously, Gr VI rats were administered RCO (0.5 ml/rat). Rats were subjected to two-third partial hepatectomy after 3 weeks (Fig. 1). After killing, the livers were excised from each rat, cleaned, weighed and stored at -80°C until further processing. Tissue sections (10 μm) were cut from the frozen livers using a cryostat microtome (SLEE Medical GmbH, Menz, Germany), and were stained for the histochemical localisation of various markers such as GST-P, γ -glutamyl transpeptidase (GGT), glucose-6-phosphatase (G6Pase), adenosine triphosphatase (ATPase) and alkaline phosphatase (AlkPase).

Methodology

Chromosomal aberration assay

Colchicine (4 mg/kg of the body weight) was given to the rats 2 h before killing in order to arrest the metaphase stage, and a cytogenetic analysis was carried out⁽²³⁾. Briefly, the bone marrow was flushed out from both the femurs using Hank's buffered salt solution (pH 7.2) and centrifuged, and the pellet was re-dispersed in a hypotonic solution (0.56%, w/v) of KCl at 37°C to permit osmotic swelling of cells. Swollen cells were fixed in ice-cold Carnoy's fluid, placed on slides and stained with a phosphate-buffered 5% Giemsa solution. A total of

seventy-five well-spread metaphase plates per rat in each Gr were analysed for chromosomal aberrations at a magnification of $100\times$, and the mitotic index was calculated from a scan of 1000 cells per rat and classified as breaks, fragments and exchanges. The incidence of aberrant cells was expressed as the percentage of damaged cells (aberrant metaphases).

Micronuclei induction assay

The frequency of micronucleated polychromatic erythrocytes was evaluated using a modified protocol of Shukla *et al.*⁽²³⁾. The bone marrow was flushed from both the femurs using Hank's buffered salt solution, 1% (w/v) bovine serum albumin and 0.15% (w/v) EDTA (pH 7.2). Evenly spread bone marrow smears were stained using the May-Grunwald and Giemsa stain. A total of 1000 polychromatic erythrocytes were scored for each treated and control Gr.

Biochemical estimations of antioxidant enzymes

The protein content of the tissue was determined by the method of Lowry *et al.*⁽²⁴⁾ using bovine serum albumin as the standard. Cu/Zn superoxide dismutase activity was analysed as per the protocol of Kakkar *et al.*⁽²⁵⁾, and was expressed as specific activity (units/min per mg protein). The activity of catalase was analysed according to the method of Sinha⁽²⁶⁾ using hydrogen peroxide as the substrate. The enzyme activity was measured following the disappearance of hydrogen peroxide, and expressed as μmol of hydrogen peroxide consumed/min per mg of protein. Lipid peroxidation (LPO) was analysed by the method of Ohkawa *et al.*⁽²⁷⁾, and peroxides were expressed as nmol of thiobarbituric acid-reactive substance/h per mg of tissue protein using 1,1,3,3-tetramethoxy propane as the standard.

Reactive oxygen species generation

ROS production was monitored by flow cytometry (BD-LSR II, San Jose, CA, USA) using dichlorodihydrofluorescein-diacetate dye⁽²⁸⁾. The fluorescence increased due to the hydrolysis of dichlorodihydrofluorescein-diacetate to dichlorodihydrofluorescein by some non-specific cellular esterases, and its subsequent oxidation by peroxides was measured and expressed in terms of mean fluorescence intensity using the software 'CellQuest'.

Medium-term bioassay

Immunohistochemical localisation of GST-P activity was performed as per the protocol of Tatematsu *et al.*⁽²⁹⁾. The exogenous peroxidase activity in fresh sections of liver was quenched with methanol: hydrogen peroxide solution in dark, and the non-specific binding was blocked using normal goat serum. The sections were incubated with the primary anti-GST-P antibody (1:300), and then with the respective secondary antibody. The colour was developed using 3-amino-9-ethylcarbazole and counterstained with haematoxylin. The histochemical activities of GGT, G6Pase, ATPase and AlkPase in rat liver were detected by the method of Shukla & Arora⁽³⁰⁾. Briefly, to detect GGT activity, the sections were incubated in a medium containing γ -glutamyl-4-methoxy- β -naphthylamide

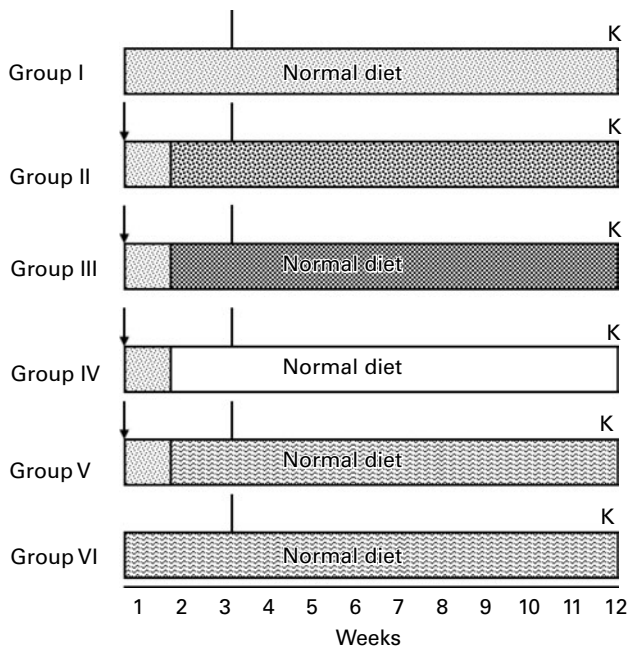


Fig. 1. Treatment schedule for medium-term bioassay. (□), Untreated; (↓), diethylnitrosamine (DEN); (■), 2-acetylaminofluorene (AAF) 0.05% mixed in crushed diet; (▣), fresh coconut oil (FCO) 0.5 ml/animal by gavage; (▢), single-heated coconut oil (SCO) 0.5 ml/animal by gavage; (▤), repeatedly heated coconut oil (RCO) 0.5 ml/animal by gavage; (|), partial hepatectomy; K, killed.

Table 1. The amount ($\mu\text{g}/\text{kg}$) of polyaromatic hydrocarbons (PAH) in fresh (FCO), single-heated (SCO) and repeatedly heated coconut oil (RCO) samples

(Mean values with their standard errors of three independent set of analyses)

PAH	Coconut oil ($\mu\text{g}/\text{kg}$)					
	FCO		SCO		RCO	
	Mean	SE	Mean	SE	Mean	SE
Naphthalene	2.98	0.10	0.31	0.01	25.98	3.20
Acenaphthylene	7.03	0.50	12.2	1.40	5.66	1.01
Fluorene + acenaphthene	1.03	0.70	0.92	0.01	2.05	0.03
Phenanthrene	6.50	0.40	9.57	1.10	4.87	0.09
Anthracene	0.37	0.01	11.23	1.70	4.25	0.10
Fluoranthene	0.78	0.04	2.75	0.08	12.09	2.08
Pyrene	0.59	0.03	0.88	0.01	12.38	3.10
Benzo(a)anthracene + chrysene	1.56	0.10	1.2	0.05	14.45	2.90
Benzo(k)fluoranthene	1.44	0.06	2.03	0.10	14.87	1.90
Benzo(b)fluoranthene	1.34	0.08	2.32	0.20	14.68	2.20
Benzo(a)pyrene	1.33	0.05	1.68	0.10	18.85	2.70
Dibenzo(a,h)anthracene	2.86	0.10	21.12	2.80	15.22	1.90
Indeno(1,2,3cd)pyrene + Benzo(g,h,i)perylene	2.38	0.10	0.048	0.001	14.66	2.10
Total	30.19		66.25		160.01	

dissolved in dimethyl sulphoxide, glycylglycine and fast blue BB salt, and were transferred into a copper sulphate solution, washed and mounted. To detect G6Pase activity, sections were incubated in a medium containing lead nitrate and glucose-6-phosphate at 37°C for 30 min, and were developed in yellow ammonium sulphide. The brownish black deposits of lead sulphide indicated the active site of G6Pase activity. To detect ATPase activity, the sections were incubated in a medium containing lead nitrate, magnesium chloride and ATP for 60 min at room temperature, and were developed in yellow ammonium sulphide. Brown precipitates indicated the site of enzyme activity. For the localisation of AlkPase activity, the sections were incubated in a freshly

prepared medium containing sodium β -glycerophosphate, sodium diethylbarbiturate, calcium chloride and magnesium sulphate. After incubation, sections were transferred to cobalt nitrate solution and developed in yellow ammonium sulphide. The black deposits of cobalt sulphide indicated the site of enzyme activity.

The activity of biomarker enzymes was observed with reference to the area and count of altered hepatic foci (AHF) in each Gr using Leica QWin500 image analysis software (Leica Mikrosysteme Vertrieb GmbH, Wetzlar, Germany). The image analysis was performed for each slide in triplicates with at least ten fields in each slide. The foci were scored only if their diameter was >0.20 mm.

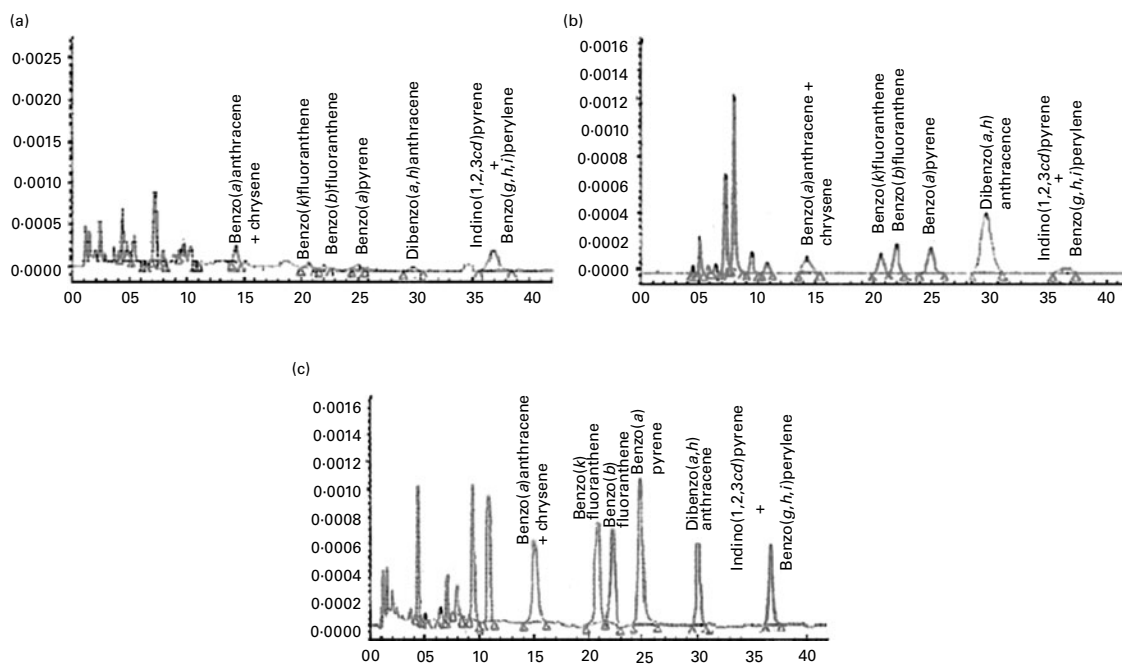
**Fig. 2.** HPLC chromatograms (autoscaled) showing the presence of polyaromatic hydrocarbons in coconut oil: (a) fresh coconut oil, (b) single-heated coconut oil and (c) repeatedly heated coconut oil.

Table 2. Effects of repeatedly heated coconut oil treatment on mitotic index, chromosomal aberration and micronuclei induction in Wistar rats

(Mean values with their standard errors of ten rats)

Gr†	Mitotic index		No. of aberrant cells (%)				Total no. of aberrant cells		MNPCE/1000 PCE	
	Mean	SE	Breaks	Fragments	Exchanges	Multiple damages	Mean	SE	Mean	SE
I	5.0	0.6	2.42	2.01	0.43	1.14	6.0	0.4	4.2	0.4
II	1.4	0.1	7.62	7.45	4.43	7.70	27.2	2.5	9.8	0.7
III	4.8	0.3	2.36	2.04	0.21	1.49	6.1	0.7	4.4	0.4
IVa	4.6	0.5	2.38	2.24	0.22	1.60	6.4	0.7	4.6	0.3
IVb	4.5	0.5	2.43	2.12	0.25	2.00	6.8	0.9	4.8	0.8
Va	3.0**	0.3	3.48	2.74	1.52	5.56	13.3*	1.2	7.4*	0.7
Vb	2.2**	0.1	4.62	3.91	1.55	5.62	15.7*	1.3	8.5*	0.5
Vc	2.1**	0.2	4.87	4.01	2.13	5.49	16.5*	1.2	8.7*	0.9

Gr, groups; MNPCE, micronucleated polychromatic erythrocytes; PCE, polychromatic erythrocyte; B(a)P, benzo(a)pyrene.

Mean values were significantly different for induction and suppression over untreated Gr I: * $P < 0.05$, ** $P < 0.05$.

† Gr I, untreated; Gr II, B(a)P (100 mg/kg body weight) treated; Gr III, fresh coconut oil (0.5 ml/rat) treated; Gr IV, single-heated coconut oil (IVa: 0.25 ml/rat and IVb: 0.5 ml/rat) treated; Gr V, repeatedly heated coconut oil (Va: 0.1 ml/rat, Vb: 0.25 ml/rat and Vc: 0.5 ml/rat) treated.

Statistical analysis

The statistical analysis for different parameters between different Gr was evaluated using Student's *t* test; $P < 0.05$ was considered significant for all the assays.

Results

Presence and amounts of polycyclic aromatic hydrocarbons in repeatedly heated coconut oil

On the basis of their molecular weight, carcinogenicity and ring sizes, PAH have been categorised into two classes: 'light PAH', such as naphthalene, acenaphthylene, fluorene + acenaphthene, phenanthrene, anthracene, fluoranthene and pyrene, and 'heavy PAH', such as benzo(a)anthracene + chrysene, benzo(k)fluoranthene, benzo(b)fluoranthene, B(a)P,

dibenzo(a,h)anthracene, indeno(1,2,3cd) pyrene and benzo(g,h,i)perylene⁽³¹⁾. Higher amount of PAH was present in RCO (160.0 $\mu\text{g}/\text{kg}$) than in FCO (30.2 $\mu\text{g}/\text{kg}$) and SCO (66.3 $\mu\text{g}/\text{kg}$) (Table 1 and Fig. 2). The amount of light PAH was 19.3 $\mu\text{g}/\text{kg}$ (64 % of the total PAH) in FCO extract, 37.9 $\mu\text{g}/\text{kg}$ (57 % of the total PAH) in SCO extract and 67.3 $\mu\text{g}/\text{kg}$ (42 % of the total PAH) in RCO extract. RCO showed an increase in the concentrations of naphthalene (9-fold), fluoranthene (16-fold) and pyrene (21-fold) among the light PAH than FCO. The quantity of heavy PAH in the extracts prepared from FCO, SCO and RCO was 10.9 $\mu\text{g}/\text{kg}$ (36 % of the total PAH), 28.4 $\mu\text{g}/\text{kg}$ (43 % of the total PAH) and 92.7 $\mu\text{g}/\text{kg}$ (58 % of the total PAH), respectively. Apparently, out of the total content of PAH in the oil extracts, the amount of heavy PAH was more in RCO than in FCO and SCO. RCO showed an increase in the concentrations of

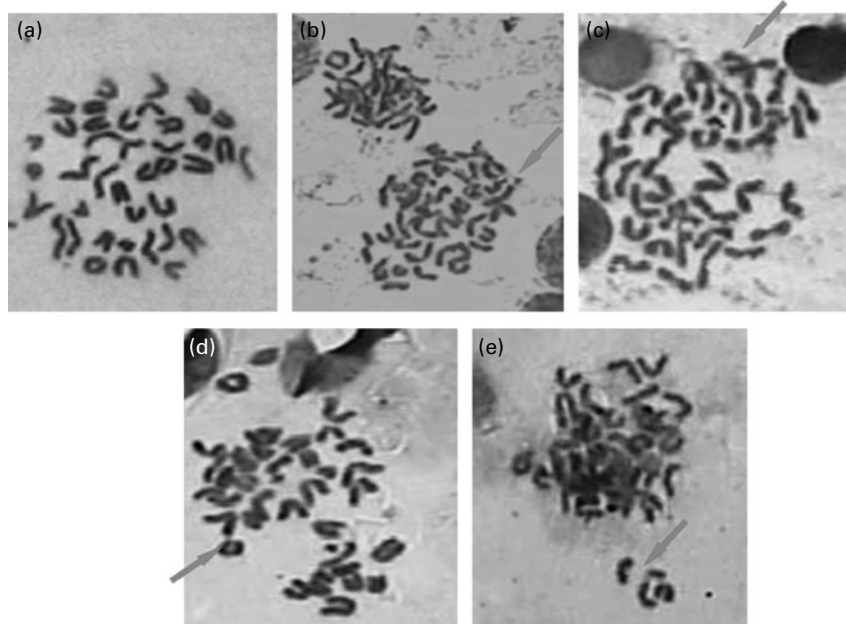


Fig. 3. Representative pictures (100 \times) showing chromosomal aberrations induced by repeatedly heated coconut oil (0.5 ml/rat) in bone marrow of Wistar rats (a) no aberration; (b) exchange; (c) break; (d) ring and (e) fragment.

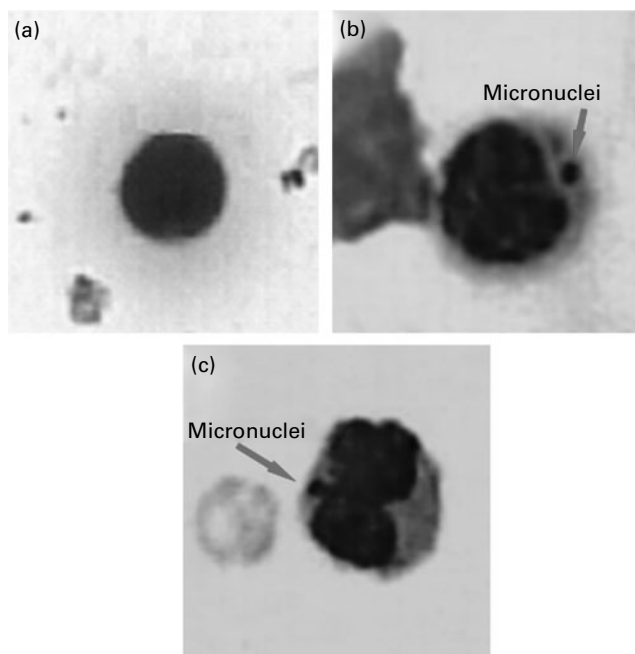


Fig. 4. Representative pictures showing micronuclei (100 ×) in (a) untreated; (b) benzo(a)pyrene-treated (100 mg/kg body weight) and (c) repeatedly heated coconut oil-treated (0.5 ml/rat) groups.

benzo(a)anthracene + chrysene (9-fold), benzo(k)fluoranthene (10-fold), benzo(b)fluoranthene (11-fold), B(a)P (14-fold), dibenzo(a,h)anthracene and indeno(1,2,3cd)pyrene + benzo(g,h,i)perylene (6-fold) compared with FCO (Fig. 2(c)).

Effects of repeatedly heated coconut oil repeatedly heated coconut oil on induction of micronuclei and chromosomal aberrations

Administration of RCO caused a higher incidence of micronuclei and chromosomal aberrations (Table 2 and Figs. 3 and 4) in rats. An induction in micronuclei of up to 2.3-fold was found in the positive control (B(a)P administered) Gr.

A significant induction in the percentage of aberrant cells of up to 2.2-, 2.3- and 2.4-fold was observed in the RCO-administered Gr Va, b and c, respectively. Mitotic index was also found to decrease up to 1.6-fold in Gr Va, 2.3-fold in Gr Vb and 2.4-fold in Gr Vc. A dose-dependent increase ($P < 0.05$) in the incidence of micronucleated polychromatic erythrocytes/1000 polychromatic erythrocyte was recorded in the RCO-administered rats (Table 2). SCO administration increased aberration, micronuclei induction and cytotoxicity up to a marked level compared with the FCO treatment and controls ($P < 0.05$).

Effects of repeatedly heated coconut oil on induction of oxidative stress

A significant decrease in the levels of antioxidant enzymes catalase (65.1%) and superoxide dismutase (45.6%) were found in B(a)P-administered Gr II than in Gr I (Table 3). The activity of catalase (55.0%) and superoxide dismutase (28.0%) also declined in RCO-fed rats ($P < 0.05$). A considerable induction in LPO levels was observed in B(a)P (80%)- and RCO (40%)-administered rats compared with the untreated controls (Table 3). However, in FCO (Gr III)- and SCO (Gr IV)-treated Gr, no significant changes in the studied parameters were observed ($P > 0.05$).

The determination of intracellular ROS generation in terms of the mean fluorescence intensity of dichlorodihydrofluorescein-diacetate dye indicated 94.66 (SE 1.5) in the untreated Gr, which significantly increased up to 176.68 (SE 6.6) in the B(a)P-fed Gr, 147.29 (SE 10.2) in the RCO-fed Gr and 129.38 (SE 3.4) in the SCO-fed Gr.

Effects of repeatedly heated coconut oil on induction of altered hepatic foci

Rats administered DEN + 2-acetylaminofluorene (Gr II) exhibited a significant ($P < 0.05$) decrease in both body weight (135.5 (SE 21.5) v. control 210.0 (SE 21.5)) and relative liver weight (2.84 (SE 0.3) v. control 3.34 (SE 0.4)) compared with Gr I rats. RCO along with DEN also decreased

Table 3. Effects of repeatedly heated coconut oil treatment on levels of antioxidant enzymes and lipid peroxidation (LPO) in Wistar rats (Mean values with their standard errors of ten rats)

Gr‡	Cu/Zn superoxide dismutase (unit/min per mg protein)†		Catalase (μ m/min per mg protein)†		LPO (nm TBARS/mg protein)†	
	Mean	SE	Mean	SE	Mean	SE
I	0.068	0.003	195.12	11.7	3.20	0.2
II	0.037* (45.6%)	0.001	68.06* (65.1%)	4.7	5.76** (80.0%)	0.3
III	0.063 (7.3%)	0.002	178.30 (8.6%)	12.5	3.36 (5.14%)	0.2
IV	0.062 (9.4%)	0.003	174.64 (10.5%)	12.2	3.59 (12.2%)	0.3
V	0.049* (27.94%)	0.002	87.85* (54.98%)	5.7	4.47** (39.68%)	0.3

Gr, groups; TBARS, thiobarbituric acid-reactive substance.

Mean values significantly decrease in the level of antioxidant enzymes over untreated Gr I and significantly increase in LPO of rats over untreated Gr I: * $P < 0.05$, ** $P < 0.05$.

† Numbers given in the parentheses indicate the percentage change over untreated Gr I.

‡ Gr I, untreated; Gr II, B(a)P (100 mg/kg body weight) treated; Gr III, fresh coconut oil (0.5 ml/rat) treated; Gr IV, single-heated coconut oil (0.5 ml/rat) treated; Gr V, repeatedly heated coconut oil (0.5 ml/rat) treated.

Table 4. Altered hepatic foci-inducing effects of repeatedly heated coconut oil treatment in Wistar rats in terms of the percentage area (foci/mm²) (Mean values with their standard errors of ten rats)

Gr†	Treatment	Alkaline phosphatase†		ATP†		Glucose-6-phosphatase†		Glutathione-S-transferase†		γ-Glutamyl transpeptidase†	
		Mean	SE	Mean	SE	Mean	SE	Mean	SE	Mean	SE
I	Untreated	2.04	0.05	1.92	0.02	2.44	0.05	1.90	0.04	2.58	0.30
II	DEN + 2-AAF	0.43 (79.0%) ↓ §	0.01	0.29 (85%) ↓ §	0.06	0.18 (93%) ↓ §	0.03	4.11 (116.0%) ↑ *	0.80	5.30 (105.0%) ↑ *	0.60
III	DEN + FCO	2.01 (1.5%)	0.21	1.85 (3.6%)	0.10	2.41 (1.2%)	0.01	2.08 (9.4%)	0.10	2.70 (4.6%)	0.30
IV	DEN + SCO	1.92 (5.9%)	0.15	1.76 (8.3%)	0.06	2.26 (7.4%)	0.01	2.11 (11.2%)	0.50	2.82 (9.3%)	0.20
V	DEN + RCO	0.87 (57.0%) ↓ §	0.09	1.00 (48%) ↓ §	0.11	1.32 (46%) ↓ §	0.14	3.27 (71.0%) ↑ *	0.34	3.92 (52.0%) ↑ *	0.40
VI	Saline + RCO	1.44 (29.4%) ↓ §	0.20	1.38 (28%) ↓ §	0.18	1.72 (29.5%) ↓ §	0.26	2.44 (28.4%) ↑ *	0.21	3.28 (27.0%) ↑ *	0.30

Gr, groups; DEN, diethylnitrosamine; 2-AAF, 2-acetylaminofluorene; FCO, fresh coconut oil; SCO, single-heated coconut oil; RCO, repeatedly heated coconut oil.

* Mean values were significantly different in induction (↑) in the level of positive biomarkers with respect to Gr I ($P < 0.05$).

† Numbers given in the parentheses indicate the percentage change over to Gr I.

‡ Gr I, untreated; Gr II, DEN (200 mg/kg body weight) + 2-AAF (0.05% in crushed diet) treated; Gr III, DEN (200 mg/kg body weight) + FCO (0.5 ml/rat) treated; Gr IV, DEN (200 mg/kg body weight) + SCO (0.5 ml/rat) treated;

§ Gr V, DEN (200 mg/kg body weight) + RCO (0.5 ml/rat) treated; Gr VI, RCO (0.5 ml/rat) treated.

§ The symbol ↓ represents significant suppression in the level of negative biomarkers.

Table 5. Altered hepatic foci-inducing effects of repeatedly heated coconut oil treatment in Wistar rats in terms of the count of foci/mm² (Mean values with their standard errors of ten rats)

Gr†	Treatment	Alkaline phosphatase†		ATP†		Glucose-6-phosphatase†		Glutathione-S-transferase†		γ-Glutamyl transpeptidase†	
		Mean	SE	Mean	SE	Mean	SE	Mean	SE	Mean	SE
I	Untreated	48.4	5.2	924.5	94.5	286.3	29.5	348.2	35.8	252.5	25.3
II	DEN + 2-AAF	7.8 (84.0%) §	0.9	220.6 (76.0%) §	23.3	86.6 (70.0%) §	9.5	636.4 (83.0%) *	62.2	479.3 (89.0%) *	50.3
III	DEN + FCO	44.6 (7.8%)	5.2	892.6 (3.4%)	10.2	263.4 (7.9%)	26.5	364.8 (4.8%)	36.5	270.3 (7.0%)	28.4
IV	DEN + SCO	42.6 (12.0%)	4.8	841.3 (9.0%)	83.7	260.5 (9.0%)	18.4	384.8 (10.5%)	36.8	280 (11.0%)	26.7
V	DEN + RCO	28.4 (41.0%) §	30.5	570.7 (38.0%) §	56.8	182.8 (36.0%) §	19.5	498.2 (43.0%) *	20.3	386.3 (53.0%) *	12.8
VI	Saline + RCO	33.3 (31.0%) §	5.5	718.5 (22.0%) §	90.3	225.6 (21.0%) §	29.1	439.5 (26.0%) *	36.6	326.6 (29.0%) *	28.2

Gr, groups; DEN, diethylnitrosamine; 2-AAF, 2-acetylaminofluorene; FCO, fresh coconut oil; SCO, single-heated coconut oil; RCO, repeatedly heated coconut oil.

* Mean values were significantly different in induction in the level of positive biomarkers with respect to Gr I ($P < 0.05$).

† Numbers given in the parentheses indicate the percentage change over to Gr I.

‡ Gr I, untreated; Gr II, DEN (200 mg/kg body weight) + 2-AAF (0.05% in crushed diet) treated; Gr III, DEN (200 mg/kg body weight) + FCO (0.5 ml/rat) treated; Gr IV, DEN (200 mg/kg body weight) + SCO (0.5 ml/rat) treated;

§ Gr V, DEN (200 mg/kg body weight) + RCO (0.5 ml/rat) treated; Gr VI, RCO (0.5 ml/rat) treated.

§ Significant suppression in the level of negative biomarkers are represented.

relative liver weight (3.00 (SE 0.4); $P < 0.05$), whereas no change in body weight was observed. FCO and SCO treatments had no effects on the body weight and liver weight of rats when compared with the treatment given to Gr I rats ($P > 0.05$).

The results revealed significant changes in positive (increase in GST-P and GGT) and negative biomarker (decrease in AlkPase, ATPase and G6Pase) levels in Gr given DEN + 2-acetylaminofluorene (Gr II), DEN + RCO (Gr V) and RCO alone (Gr VI) with respect to the untreated Gr, Gr I (Tables 4 and 5 and Fig. 5; $P < 0.05$). No alterations were observed in any positive or negative biomarkers by oral administration of either FCO (Gr III) or SCO (Gr IV) compared with the untreated controls ($P > 0.05$).

Discussion

Food that we eat may contain carcinogenic contaminants, or may become a mutagen itself after the transformation of some of its components⁽³²⁾. To date, many dietary constituents have been studied for their carcinogenic or mutagenic potential^(3,11,33). Relationship between dietary use of vegetable fats and risk of cancers has long been known, but studies exploring the mechanism for this connection are ambiguous. In this context, the present study was conducted to identify and quantify the contaminants generated in CO during the process of repeated heating to the temperatures usually used

for cooking. There are reports regarding the formation of carcinogenic compounds by the heat processing of vegetable oils^(11,12). These compounds may form DNA adducts, which have been proposed as predictive biomarkers of human cancers⁽³⁴⁾. Results of the study revealed the presence of a higher amount of total PAH in RCO (six times heated) than in SCO and FCO (Table 1). Pandey *et al.*⁽¹⁵⁾ also reported that repeated heating of edible oil at high temperatures generates greater amounts of PAH, which are dependent on the number of cycles used for heating. Moreover, RCO contained a higher amount of heavy PAH of the total PAH than FCO and SCO. Heavy PAH are reported to have significant carcinogenic and mutagenic potential⁽³⁵⁾.

As the amount of PAH was high in RCO, it is assumed that its dietary use may result in risk of genotoxicity and carcinogenicity. The assays performed to detect both chromosomal aberration and micronuclei induction in the bone marrow are considered to be able to detect early effects of genotoxicity following the exposure to mutagens and carcinogens⁽²³⁾. By using these two assays in the present study, RCO was found to enhance the incidence of aberrant cells, including breaks, fragments, exchanges and multiple chromosomal damages and micronuclei in a dose-dependent manner (Table 2). Chromosomal aberration and micronuclei induction by B(a)P⁽²³⁾ and heated cooking oils⁽³⁶⁾ are well documented. The results of the present study suggest that RCO may contain a high amount of B(a)P (Table 1).

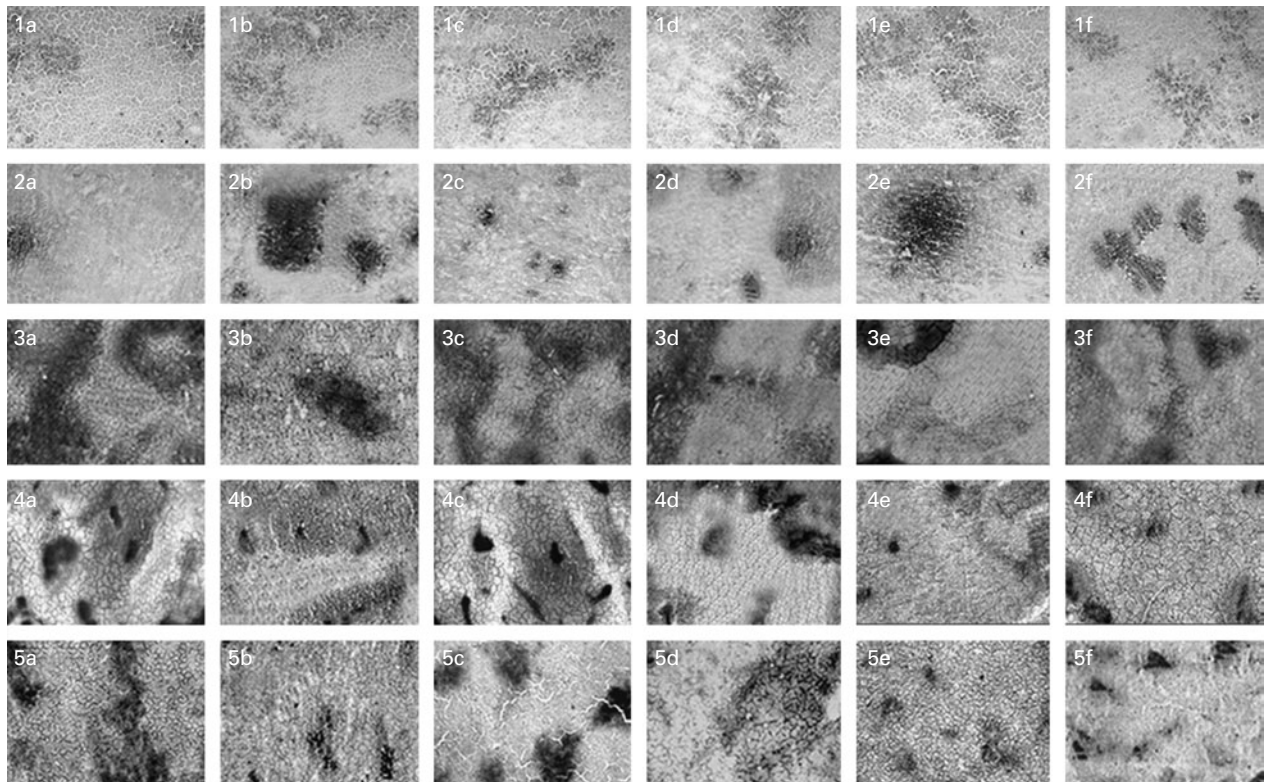


Fig. 5. Representative pictures (10 ×) for cellular localisation of the expression of liver-specific enzyme markers showing (1) γ -glutamyl transpeptidase, (2) glutathione-S-transferase, (3) ATP, (4) glucose-6-phosphatase, (5) alkaline phosphatase, where (1a–5a) represent untreated group; (1b–5b) represent diethylnitrosamine (200 mg/kg body weight) + 2-acetylaminofluorene (0.05% in crushed diet)-treated group; (1c–5c) represent diethylnitrosamine (200 mg/kg body weight) + fresh coconut oil (0.5 ml/rat)-treated group; (1d–5d) represent diethylnitrosamine (200 mg/kg body weight) + single-heated coconut oil (0.5 ml/rat)-treated group; (1e–5e) represent diethylnitrosamine (200 mg/kg body weight) + repeatedly heated coconut oil (RCO; 0.5 ml/rat)-treated group; (1f–5f) represent RCO (0.5 ml/rat)-treated group.

The prevention of oxidation is an essential process as decreased antioxidant levels may lead to cytotoxicity, mutagenicity and carcinogenicity^(37,38). There is evidence linking the metabolism of PAH with the generation of ROS and thus oxidative stress⁽³⁹⁾. Cooking oil fumes were found to induce DNA damage via ROS production⁽⁸⁾. Our results revealed a significant ($P < 0.05$) decrease in the levels of antioxidant enzymes (viz. catalase and superoxide dismutase) and increase in the levels of LPO and ROS following RCO administration (Table 3). Accordingly, the decrease in the levels of antioxidant enzymes has been observed earlier in rats fed heated and fried oil⁽⁴⁰⁾. Altered levels of antioxidant enzymes are evident in cancers⁽³⁸⁾. From the present set of observations, it has been concluded that RCO has the potential to cause oxidative imbalance by decreasing antioxidant enzyme levels and increasing ROS and LPO, which in turn may cause susceptibility towards cancer development⁽⁸⁾. Ip & Sinha⁽⁴¹⁾ reported that saturated fat induces LPO. A study done by Udilova *et al.*⁽⁴²⁾ suggested that the peroxidation of biomembranes by hydroperoxides from heated oils may contribute to enhanced risk of cancers. In another study, 1,*N*(6)-ethenodeoxyadenosine, which is determined as a marker for DNA damage caused by LPO products and a marker for oxidative stress, formed the highest adducts in the CO diet⁽⁴³⁾.

Induction of preneoplastic lesions is among the events leading to liver tumour development, and has been used to assess the carcinogenic/anticarcinogenic potential of various agents⁽³⁰⁾. A number of genotoxic responses have been reported in fresh or heat-treated foodstuffs, which are also reported to be carcinogenic in rodent bioassays. Earlier, we have shown that heated mustard oil induces AHF significantly⁽³⁰⁾. In the present study, investigations were performed for the evaluation of AHF-developing potential of RCO. A significant enhancement in the expression of positive biomarker enzymes (GGT and GST-P) and a reduction in negative markers (AlkPase, ATPase and G6Pase) were observed in the rats administered DEN + RCO and RCO alone compared with the FCO- and SCO-treated Gr (Tables 4 and 5). GST-P is used as a diagnostic marker for hepatocellular carcinoma, and is also used for the detection of other chronic liver diseases⁽⁴⁴⁾. Another study suggested that a deficiency in canalicular ATPase and G6Pase provides the best marker for the larger foci and nodules, and thus can be used to identify preneoplastic changes in the liver⁽⁴⁵⁾. Aruna *et al.*⁽⁴⁶⁾ suggested that initiation of hepatocarcinogenesis is influenced by thermally oxidised vegetable oil. The AHF-inducing activity of RCO in DEN-initiated rats may be attributed to its SFA. Repeatedly used frying oils when administered to rats have been shown to produce indications of cellular damage to the liver and kidneys⁽⁴⁷⁾. RCO leads to the induction of oxidative stress, which in turn inhibits the activity of enzymes such as G6Pase and ATPase within the cells⁽¹³⁾. Increase in ROS production is related to epigenetic toxicity⁽⁴⁸⁾, and also accounts for GST-P induction in hepatocytes⁽⁴⁹⁾. Since tumour initiation and promotion involve genetic and epigenetic pathways, respectively, it seems like RCO (Table 1) having more amounts of carcinogenic PAH and inducing AHF alone as well as with DEN works both as an initiator and as a promoter.

In conclusion, the presence of a high amount of carcinogenic PAH in RCO was responsible for oxidative stress via enhancement of ROS generation and LPO, thus leading to

the formation of preneoplastic lesions in rat liver. Therefore, dietary consumption of RCO exhibits the possible risk of carcinogenicity and genotoxicity.

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