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## Mini Review

## The role of reactive oxygen species (ROS) and cytochrome P-450 2E1 in the generation of carcinogenic etheno-DNA adducts

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## ABSTRACT

Exocyclic etheno-DNA adducts are mutagenic and carcinogenic and are formed by the reaction of lipidperoxidation (LPO) products such as 4-hydroxynonenal or malondialdehyde with DNA bases. LPO products are generated either *via* inflammation driven oxidative stress or *via* the induction of cytochrome P-450 2E1 (CYP2E1). In the liver CYP2E1 is induced by various compounds including free fatty acids, acetone and ethanol. Increased levels of CYP2E1 and thus, oxidative stress are observed in the liver of patients with non-alcoholic steatohepatitis (NASH) as well as in the chronic alcoholic. In addition, chronic ethanol ingestion also increases CYP2E1 in the mucosa of the oesophagus and colon. In all these tissues CYP2E1 correlates significantly with the levels of carcinogenic etheno-DNA adducts. In contrast, in patients with non-alcoholic steatohepatitis (NASH) hepatic etheno-DNA adducts do not correlate with CYP2E1 indicating that in NASH etheno-DNA adducts formation is predominately driven by inflammation rather than by CYP2E1 induction. Since etheno-DNA adducts are strong mutagens producing various types of base pair substitution mutations as well as other types of genetic damage, it is strongly believed that they are involved in ethanol mediated carcinogenesis primarily driven by the induction of CYP2E1.

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## Introduction

Oxidative stress is an important mechanism in the pathogenesis of many diseases including cancer. The generation of reactive oxygen species (ROS) with consecutive DNA damage is an initial step in carcinogenesis induced by inflammatory processes. During

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inflammation ROS is generated among others through various cytokines, but also through other mechanisms such as the induction of cytochrome P4502E1 (CYP2E1) as demonstrated following chronic alcohol consumption [1,2]. This review will focus on the effect of alcohol on CYP2E1 and its role in ROS formation, but major emphasis will be led on the generation of carcinogenic exocyclic etheno-DNA adducts as a consequence of the reaction between lipidperoxidation products generated by ROS and DNA bases following ethanol administration *in vitro* and *in vivo*. It will be shown that these etheno-DNA adducts following chronic ethanol consumption are of major importance with respect to ethanol mediated carcinogenesis in the liver and in other tissues.

### Inflammation, oxidative stress and DNA damage

Chronic inflammation induced by various agents including viruses and bacteria is associated with an increased cancer risk due to tissue damage and genetic instability [3–7]. Oxidative stress with the generation of ROS may occur in chronic infection and inflammation primarily due to the generation of nitric oxide (NO), superoxide anion ( $O_2^-$ ) and other ROSs by macrophages and neutrophils that infiltrate the inflamed tissue [8,9].

Activated inflammatory cells in various tissues including the liver in turn induce oxidant generating enzymes such as NADPH oxidase, inducible nitric oxide synthetase (iNOS), xanthine oxidase (XO) and myeloperoxidase (MPO) [10,11]. In such conditions ROS and reactive nitrogen species (RNS) are generated. As a consequence ROS and RNS can damage DNA, RNA, lipids and proteins through nitration and oxidation resulting in an increased mutation load [10,11].

Furthermore, cytokines are released in inflammatory tissues which not only activate the above mentioned enzymes to create ROS and RNS, which also activate NFκB a nuclear transcription factor which among others stimulates cyclooxygenase 2 (COX2), lipoxigenase (LOX), and iNOS [4,10–12]. Upregulation of iNOS, COX2, and LOX results also in an overproduction of ROS and RNS [10].

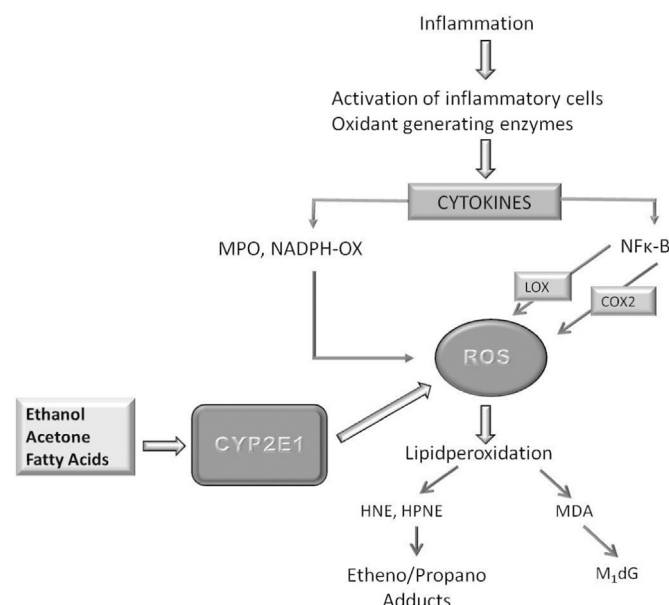
iNOS catalyses nitric oxide (NO) generation which reacts with oxygen to produce  $N_2O_3$  a strong nitrosating compound which deaminates DNA bases and react with secondary amines to form N-nitrosoamines which are highly carcinogenic [10].

Another reaction with  $O_2$  leads to peroxyinitrite with the formation of 8-nitroguanine. Peroxyinitrite also results in single strand breakage of DNA [7,10].

COX2 catalyse the conversion of arachidonic acid (AA) to prostaglandins is inducible by various factors including NFκB, cytokines and tumour promoters and may influence apoptosis, angiogenesis, tumour invasion, but also the generation of oxidative stress [12,13]. An upregulation of COX2 has been shown in familial adenomatous polyposis (FAP) and in the Apc Min mouse model which resembles FAP and this was associated with a highly significant increase in various etheno-DNA lesions [13–16].

LOX metabolizes AA to hydroxyeicosatetraenoic acids (HETEs) or leukotrienes. It has been shown in mouse skin carcinogenesis that LOX isoenzymes are overactivated and some of the metabolites cause chromosomal damage which was found to be inhibited by LOX inhibitors [17,18].

Thus, all the factors mentioned above lead to the generation of ROS and RNS with consequent lipid peroxidation and the production of lipidperoxidation products such as 4-hydroxynonenal (4HNE), 4-hydroxyhydroperoxy-2-nonenal (HPNE) and malondialdehyde (MDA) (Fig. 1). These lipidperoxidation products react with DNA either directly or through bifunctional intermediates to form various promutagenic exocyclic etheno-DNA adducts. Some major types are depicted in Fig. 2.



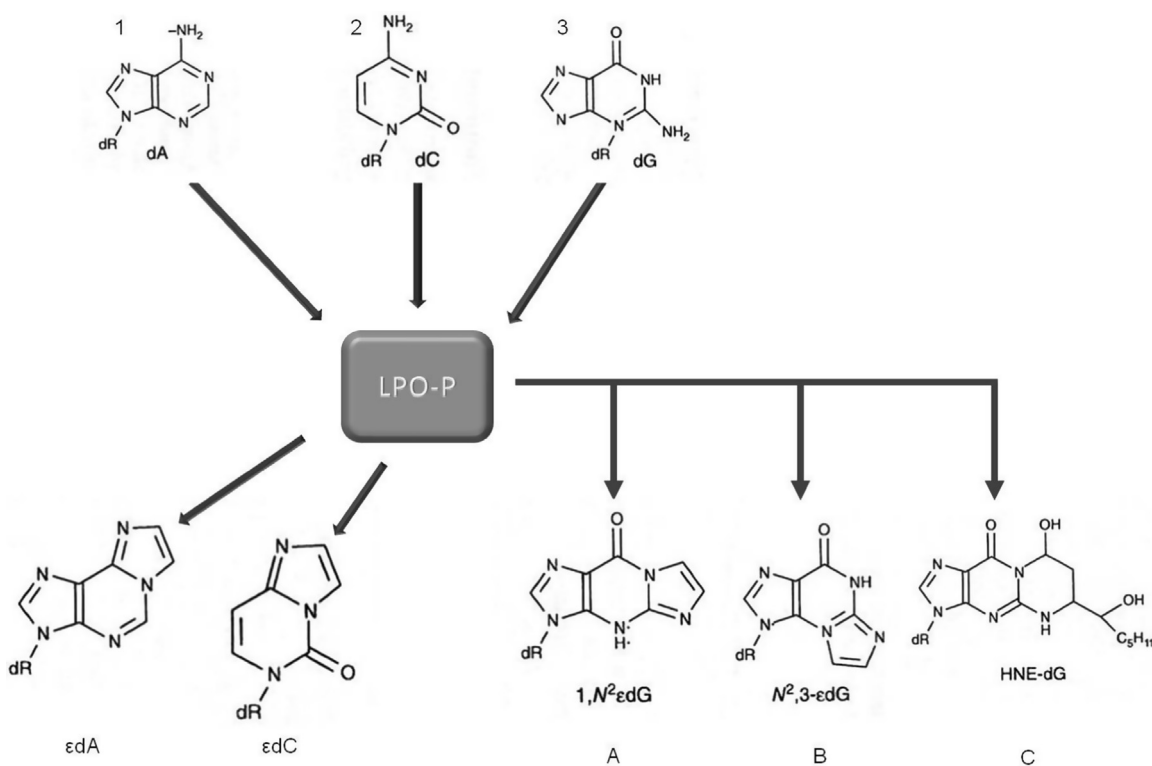
**Fig. 1.** Simplified pathophysiology of reactive oxygen species (ROS) and etheno-DNA adduct formation. Inflammation driven cytokine secretion results among others in NFκB activation and in the activation of NADPH oxidase (NADPH-Ox) as well as myeloperoxidase (MPO). NFκB is also activated by acetaldehyde, the first metabolite of ethanol oxidation. NFκB stimulates lipoxigenase (LOX), cyclooxygenase 2 (COX2), and inducible nitric oxide synthase (iNOS). As a result ROS and reactive nitrogen species (RNS) are generated, which lead to lipidperoxidation with the occurrence of lipidperoxidation products such as 4-hydroxynonenal (4-HNE), 4-hydroxyhydroperoxy-2-nonenal (HPNE), and malondialdehyde (MDA). These adducts react with DNA bases to form exocyclic etheno/propane-DNA adducts. Chronic alcohol consumption results in the induction of cytochrome P-4502 E1 which is involved in ethanol oxidation through the microsomal ethanol oxidizing pathway. During this reaction ROS is generated without inflammation. Other compounds such as free fatty acids or acetone also induce CYP2E1 which is especially relevant in nonalcoholic fatty liver disease (NAFLD), when the liver is loaded with fat and in patients with diabetes mellitus when acetone is generated in the liver.

LPO products derived from  $\gamma$ -linoleic acid, including HNE, a major LPO product and its electrophilic epoxy-, hydroperoxy-, and oxo-enal intermediates react with the DNA bases A, C, and G to yield *inter alia* the unsubstituted etheno-DNA adducts 1,  $N^6$ -etheno-2'-deoxyadenosine ( $\epsilon$ dA), 3,  $N^4$ -etheno-2'-deoxycytidine ( $\epsilon$ dC), 1,  $N^2$ -etheno-2'-deoxyguanosine (1,  $N^2$  $\epsilon$ dG), and  $N^2,3$ -etheno-2'-deoxyguanosine ( $N^2,3\epsilon$ dG). In addition, also substituted base adducts are formed such as HNE-dG carrying a fatty acid chain residue (Fig. 2). 2,  $N^4$ -etheno-5-methyl-2'-deoxycytidine ( $\epsilon$ 5mdC), an endogenous, hitherto unknown LPO-derived adduct was identified in the DNA of human tissue which could play a role in epigenetic mechanisms of carcinogenesis [10,19–27].

In addition, DNA can also be modified directly by ROS and RNS to 8-nitro-dG and 8-Oxo-dG [28]. All of these DNA changes have been detected in human specimens [29–34].

### The importance of etheno DNA adducts in carcinogenesis

Exocyclic etheno-DNA adducts exhibit strong mutagenic properties producing various types of base pair substitution mutations and other types of genetic damage in all organisms tested so far [35,36].  $\epsilon$ dA can lead to AT  $\rightarrow$  GC transition and AT  $\rightarrow$  TA and AT  $\rightarrow$  CG transversions [37,38].  $\epsilon$ dC can cause CG  $\rightarrow$  AT transversions and CG  $\rightarrow$  TA transition [39,40], and  $N^2,3\epsilon$ dG can lead to GC  $\rightarrow$  AT transition [40]. Incorporation of a single  $\epsilon$ dA in either DNA strand of HeLa cells showed a similar miscoding frequency and was more mutagenic than 8-oxo-dG [41].



**Fig. 2.** Generation of various etheno DNA-adducts. Deoxyadenine, deoxycytosine, and deoxyguanosine react with lipidperoxidation products to form 1N<sup>6</sup>-ethano-2'-deoxyadenosine (εdA), 3,N<sup>4</sup>-etheno-2'-deoxycytidine (εdC), and 1,N<sup>2</sup>-etheno-2'-deoxyguanosine (1,N<sup>2</sup>εdG), N<sup>2</sup>,3-etheno-2'-deoxyguanosine (N<sup>2</sup>,3-εdG), and HNE-derived 1,N<sup>2</sup>-propano-2'-deoxyguanosine adduct (HNE-dG).

Some etheno-adducts are poorly repaired in some tissues and cells supporting their biological relevance [42]. Strong support that etheno-DNA adducts play a causal role in the initiation and progression of liver carcinogenesis comes from the formation of εdA and εdC *in vivo* by the human liver carcinogen vinyl chloride [43] and by the potent multiorgan, multispecies carcinogen urethane *via* their reactive epoxy-intermediates [44]. The biological importance of etheno-DNA adducts is further stressed as they are preferentially formed in codon 249 of TP53 (which encodes p53), leading to a mutation that renders cells more resistant to apoptosis and provides them some growth advantage [45].

LPO-derived reactive products and their macromolecular interactions have been so far characterized primarily by *in vitro* studies, making it difficult, to pinpoint the main precursors and pathways involved in the generation of cancer-relevant DNA damage in human *in vivo*. For this reason earlier studies analysed in human specimens εdA and εdC as marker lesions for several other exocyclic adducts that could be formed with DNA *in vivo*, for which sensitive detection methods were not yet available.

Using ultrasensitive and specific detection methods [46], two miscoding etheno-DNA adducts εdA and εdC and also ε5mdC were unequivocally identified in humans. Samples were collected from “at-risk” patients affected by chronic inflammatory processes, persistent viral infections, iron storage- and alcohol-related diseases or exposed to inherited/acquired cancer risk factors. Adduct levels increased 10–100-fold progressively in human cancer-prone organs including liver, bile duct, oesophagus, colon and pancreas. Consistent results were also observed in rodent tumour models, that mimic human disease (for review [47]). Taken together these data incriminate LPO-derived adducts as strongly mutagenic cancer-causing lesions.

### Alcohol and oxidative stress

Chronic ethanol consumption may result in the development of alcoholic liver disease (ALD) and cancer of various sites including the liver and the upper aerodigestive tract [2,48]. One mechanism by which alcohol exerts its deleterious effects is the generation of ROS. As already pointed out the formation of ROS such as superoxide anion (O<sub>2</sub><sup>-</sup>) and hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) causes oxidative injury [1,2]. ROS as well as acetaldehyde, the first metabolite of ethanol oxidation, both activate NFκB an important transcription factor involved in carcinogenesis [48]. Inflammation driven oxidative stress including activated hepatic macrophages as observed in alcoholic hepatitis (AH) is predominantly responsible for the generation of ROS in AH [49]. Also hepatic iron overload as observed in the alcoholic increases ROS [50,51]. Furthermore, ethanol also results in the increase of iNOS with an increased production of nitric oxide and the generation of the highly reactive peroxynitrite (ONOO<sup>-</sup>) [52].

In addition, several enzyme systems are capable to produce ROS including CYP2E1 as part of the microsomal ethanol oxidizing system (MEOS) which metabolizes ethanol to acetaldehyde in the presence of oxygen and NADPH [53,54]. This system is of major importance, since it can be induced by chronic consumption of ethanol. It has been shown that CYP2E1 induction occurs already at a daily ethanol dose of 40 g and already at 1 week of consumption, which is further enhanced with time. However, an inter-individual intensity of CYP2E1 increase has been observed. Some individuals react to alcohol consumption with a striking CYP2E1 induction, while others reveal only a weak induction of CYP2E1 [55]. It is noteworthy that CYP2E1 induction by ethanol may also depend on dietary factors since medium chain triglycerides diminish CYP2E1 induction as compared to the application of long chain triglycerides in animal experiments [56].

CYP2E1 has a high rate of NADPH oxidase activity, resulting in the generation of large quantities of  $O_2^-$ ,  $H_2O_2$  and hydroxyethyl radicals [1,2,57]. Thus, CYP2E1 dependent microsomal ethanol oxidation produces ROS leading to lipidperoxidation with the generation of lipidperoxidation products such as 4-hydroxynonenal (4-HNE) and malondialdehyde (MDA) [1,2].

The role of CYP2E1 in the formation of ROS, in the progression of ALD and in ethanol mediated carcinogenesis has been clearly demonstrated [2,58–62]. Thus, the severity of ALD was significantly enhanced in CYP2E1 overexpressing mice [63], and reduced in CYP2E1 knockout mice [62]. When chlormethiazole (CMZ), a strong and specific CYP2E1 inhibitor was given in addition to an ethanol containing diet which induces ALD a significant reduction of ROS and RNS was noted in the liver of these animals [62] associated with a striking improvement of the liver disease [64]. Subsequently, oxidized DNA lesions have been found to be lower in CYP2E1 knockout mice as compared to wild type mice following chronic alcohol administration [65].

#### *CYP2E1 induction and etheno-DNA adduct generation in the liver*

##### *Effect of ethanol*

We have recently investigated the effect of CYP2E1 on lipidperoxidation products and etheno-DNA lesions in HepG2 cells overexpressing CYP2E1, and in humans with alcoholic liver disease [66]. When HepG2 cells overexpressing CYP2E1 were incubated with increasing concentrations of ethanol up to 50 mM, an increasing load of  $\epsilon$ DA and  $\epsilon$ DC could be detected as compared to control cells. This was not only a concentration dependent, but also a time dependent process. However, when 20  $\mu$ M CMZ were added to the cell culture a highly significant inhibition of the generation of etheno-DNA adducts was observed [66].

Since increased levels of  $\epsilon$ DA adducts have been observed in the hepatic nuclei of patients with ALD (Fig. 3) [30], we extended our experiments and studied liver biopsies from alcoholic patients with various severities of ALD by using immunohistology for the detection of CYP2E1 and etheno-DNA adducts. Again there was a significant correlation between CYP2E1, the lipidperoxidation product 4-HNE and  $\epsilon$ DA, as well as  $\epsilon$ DC [66].

Most recently, we have investigated CYP2E1 and exocyclic etheno-DNA adducts in a large cohort study of 97 alcoholics with non-cirrhotic ALD. All patients were liver biopsied, histologically evaluated and CYP2E1 as well as etheno-DNA adducts were immunohistologically determined. As a result a strong significant correlation between CYP2E1 and  $\epsilon$ DA ( $p=0.0001$ ) has been observed (Seitz, personal observation).

##### *Non-alcoholic fatty liver disease (NAFLD)*

CYP2E1 is not only induced by chronic ethanol ingestion, but also in NAFLD [67–69] possibly by free fatty acids and acetone [54]. Although, this induction is less pronounced as in ALD, it also has severe consequences with respect to the generation of oxidative stress. Indeed, inflammation driven oxidative stress may be an important mechanism in the progression of NAFLD [70]. We therefore determined CYP2E1 as well as  $\epsilon$ DA in liver biopsies from patients with pure non-alcoholic fatty liver and patients with NASH using immunohistology.  $\epsilon$ DA was detected in a broad range of intensity and correlated significantly with the severity of inflammation, but not with CYP2E1 (Linhart and Seitz, personal communication).

NAFLD also is an increasing health problem in children [71–73]. As reported recently, oxidative stress as measured by the hepatic expression of 8-hydroxy-2-deoxyguanosine (8-OHG), serum protein carbonyls, and circulating antibody against malondialdehyde adducted human serum albumin has been frequently found in children with NAFLD and was also found to be associated with an

increased severity of steatohepatitis [74]. Therefore, we determined  $\epsilon$ DA, and CYP2E1 in liver biopsies of children with NASH. In these studies we also could show for the first time that not only CYP2E1 was found to be increased, but also that  $\epsilon$ DA occurs. In a few of these children at an age below 15 years and the diagnosis of diabetes mellitus a striking load of  $\epsilon$ DA was found in the nuclei of their hepatocytes [75]. In contrast to ALD these adducts did not significantly correlate with CYP2E1, but rather with the state of inflammation. Thus, inflammatory driven ROS production may be predominant to explain  $\epsilon$ DA formation in patients with NAFLD.

In animal experiments, the progression of NASH is influenced by the concomitant administration of ethanol [76,77]. This has been shown in dietary induced NASH, which could be due among others to oxidative, nitrosative, and mitochondrial stress, as well as increased inflammation and cellular apoptosis [76,77].

Furthermore, in the Zucker rat, a leptin deficiency and insulin resistance genetic NASH model the administration of ethanol not only increased CYP2E1, but also  $\epsilon$ DA in a linear way [66].

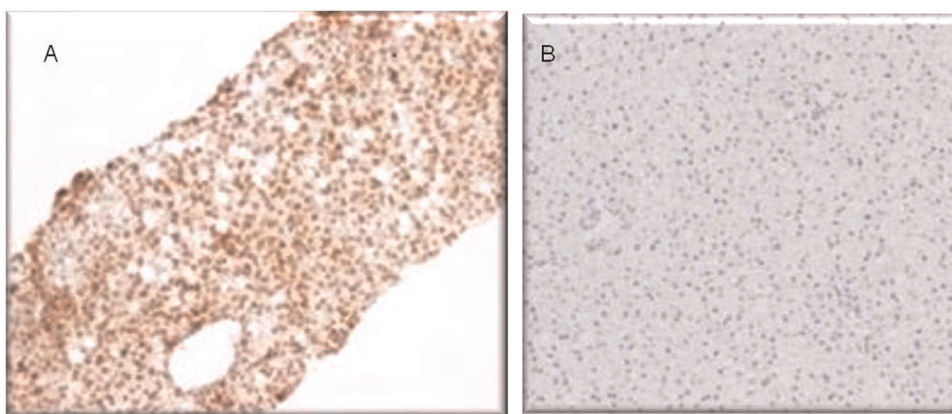
The fact that ethanol consumption in patients with NASH enhances oxidative stress possibly predominantly by a further increase in CYP2E1 associated with the generation of highly carcinogenic etheno-DNA lesions may of special interest in the context that patients with NASH have significant higher HCC risk and develop HCC in a much shorter time frame when they consume alcohol even at social levels [78].

#### *CYP2E1 induction and etheno-DNA adduct generation in the oesophagus and in the colorectal mucosa*

Chronic alcohol consumption is a major risk factor for undefined cancer [2,79]. Various mechanisms may mediate carcinogenesis including the genotoxic effect of acetaldehyde and oxidative stress [2,79–81]. As discussed above for the liver, ethanol may also exert its carcinogenic effect in other tissues among others via the induction of CYP2E1 and the generation of carcinogenic etheno-DNA adducts. Therefore, we investigated if such effects can also be observed in the human oesophagus [82]. We studied undefined biopsies of 37 patients with upper aerodigestive tract cancer and heavy alcohol consumption of more than 100 g on average per day as well as 16 controls without tumours (12 teetotallers and 4 subjects with a maximum of 25 g ethanol/day). CYP2E1, etheno-DNA adducts and Ki67 as a marker for cell proliferation were determined immunohistologically in the undefined mucosa adjacent to the tumour. Chronic alcohol ingestion resulted in a significant induction of CYP2E1 which correlated with the amount of alcohol consumed. Furthermore, a significant correlation between CYP2E1 and the generation of the carcinogenic exocyclic etheno-DNA adducts  $\epsilon$ DA and  $\epsilon$ DC was observed. Etheno-DNA adducts also correlated significantly with cell proliferation, which was especially enhanced in patients who both drank and smoked. The results showed clearly again a correlation between CYP2E1 and etheno-DNA adducts. In contrast to the liver this induction correlated significantly with the amount of ethanol ingested [82].

More recently, we also investigated immunohistologically the effect of ethanol on colorectal CYP2E1 and etheno-DNA adducts in colorectal biopsies from 31 alcoholics and 15 non-drinking controls (Linhart and Seitz, personal communication). Again we found a significant correlation between the two parameters. It is interesting that chronic ethanol consumption using a Lieber DeCarli diet resulted in a hyperproliferation of the colorectal mucosa with an extension of the proliferative compartment towards the lumen of the crypt, which is a first step in carcinogenesis of this tissue [83]. A similar observation was made in patients with heavy alcohol consumption [84]. The administration of vitamin E, a radical scavenger, however, reduced the proliferative rate significantly emphasizing indirectly that most likely oxidative stress





**Fig. 3.** Immunohistochemistry of edA in two liver biopsies from patients with alcoholic liver disease (A) and control patients with a normal liver (B). The brownish colour shows edA. This adduct occurs in the nuclei of the hepatocytes and the percentage of nuclei positive hepatocytes can be counted. A significant load of etheno adducts is observed in ALD, while the control healthy liver reveals background activity only.

induced by CYP2E1 may be responsible for this regenerative behaviour [85].

#### *CYP2E1 and experimental hepatocarcinogenesis*

It has been believed for a long time that ethanol by itself is not a carcinogen rather than a co-carcinogen or a tumour promoter. However, meanwhile various animal studies have demonstrated that the administration of ethanol alone without any chemical carcinogen can result in tumours of the liver [86], the upper aerodigestive tract [87], the mammary gland [88] and the intestine [89]. Besides the fact that DNA lesions induced either by the binding of acetaldehyde to DNA [2,48] or by the reaction of DNA with ROS do occur during chronic ethanol consumption, the role of CYP2E1 in this process has not intensively investigated.

In a series of experiments we used an animal model in which a small amount of diethylnitrosamine (20 mg/kg b.wt.) was administered once to initiate hepatocarcinogenesis [90,91]. CYP2E1, inflammatory proteins, cell proliferation, protein bound 4-HNE, etheno-DNA adducts as well as 8-hydroxy-2'-deoxyguanosine (8-OHdG), retinoid concentrations and hepatic carcinogenesis were examined. Chronic ethanol ingestion for 1 month resulted in increased CYP2E1 levels and an increased nuclear accumulation of NF $\kappa$ B protein. In addition, TNF $\alpha$  expression was also enhanced associated with increased cyclin D1 expression and p-GST positive altered hepatic foci. All these changes were significantly inhibited by the concomitant administration of CMZ. Following 10 months of ethanol feeding hepatocellular adenoma were detected in ethanol fed rats only, but not in control rats. The administration of CMZ inhibited completely the formation of hepatic adenomas. In addition, 8-OHdG formation was found to be significantly increased after alcohol and almost normalized with CMZ. Although, etheno-DNA adduct formation increased following ethanol ingestion and decreased with CMZ, this effect was not significant.

More recently, Tsuchishima and co-workers produced hepatocellular carcinoma in mice without any additional insult. This process was significantly associated with the expression of CYP2E1 [92].

#### **Summary**

The most important mechanism associated with oxidative stress and the generation of ROS is chronic inflammation. During inflammation cytokines are liberated resulting in the activation of oxidant generation enzymes such as NADPH oxidase, and NF $\kappa$ B

with the activation of LOX, Cox-2 and iNOS finally leading to the formation of ROS. In addition, ROS can also be generated through CYP2E1 which is induced by chronic alcohol consumption as well as in NASH where free fatty acids as well as acetone (mostly in diabetics) induce CYP2E1. ROS leads to lipidperoxidation with the occurrence of lipidperoxidation products such as 4-HNE and MDA. Both compounds can bind to DNA forming highly carcinogenic etheno-DNA adducts. In a series of experiments we could show that a significant correlation exists between CYP2E1 levels and etheno-DNA adduct formation in cell culture, animal experiments and biopsies from patients with ALD. Since in NASH the inflammatory process predominates as compared to the induction of CYP2E1 the etheno-DNA adduct levels do not correlate with CYP2E1 but rather with the intensity of the inflammatory process.

#### **Conclusion**

Cell culture and animal experiments as well as clinical biopsy studies in patients with ALD emphasize an important role of CYP2E1 in alcohol mediated carcinogenesis in the liver, but also in other tissues. In addition, CYP2E1 seems to be a driving force in the progression of ALD. Inhibition of CYP2E1 by a nontoxic inhibitor may be a successful approach in the treatment of ALD and alcohol mediated carcinogenesis.

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