As we previously demonstrated in human hepatocellular carcinoma (HepG2) cells, ethanol at low concentration triggers the Fas apoptotic pathway. However, its role in other intracellular signaling pathways remains unknown. Therefore, the aim of the present study was to evaluate the role of low concentration of ethanol on different intracellular signaling pathways. For this purpose, HepG2 cells were treated with 1 mM ethanol for 10 min and the phosphorylation state of protein kinases was determined. In addition, the mRNA levels of transcription factors and genes associated with the Fas apoptotic pathway were determined. Our data demonstrated that ethanol-induced phosphorylation of protein kinases modulates both anti-apoptotic and pro-apoptotic mechanisms in HepG2 cells. Pro-apoptosis resulted mainly from the strong inhibition of the G-protein coupled receptor signaling pathway. Moreover, the signal transduction initiated by ethanol-induced protein kinases phosphorylation lead to increased expression of the transcription factors with subsequent expression of genes associated with the Fas apoptotic pathway (Fas receptor, Fas ligand, FADD and caspase 8). These results indicate that low concentration of ethanol exert their effect by predominant activation of pro-apoptotic events that can be divided in two phases. An early phase characterized by a rapid transient effect on protein kinases phosphorylation, after 10 min exposure, with subsequent increased expression of transcription factors for up to 6 hr. This early phase is followed by a second phase associated with increased gene expression that began after 6 hr and persisted for more than 24 hr. This information provided a novel insight into the mechanisms of action of ethanol (1mM) in human hepatocellular carcinoma cells.

Key words: Ethanol, HepG2 cells, protein kinases, signal transduction, transcription factors, gene expression

1. Introduction

Apoptosis is a highly organized form of cell death that takes place in normal physiological processes, such as development, homeostasis, tissue turnover, and immune response. Hereby, a balance between cell survival and cell death (apoptosis and necrosis) is mandatory. Apoptosis also plays an important role in different pathological conditions (i.e. cancer, autoimmune disease and neurodegeneration) including alcoholic liver disease (ALD) [1].

The toxic effect of ethanol in the liver has been extensively demonstrated in several animal and clinical studies [2-8]. The hepatotoxic effect of ethanol directly correlates with time of exposure and the applied concentration [2-5]. High concentrations of ethanol lead to necrotic cell death [6]. This results from induction of the cytochrome P4502E1 (CYP2E1) with subsequent production of reactive oxygen species [5, 7, 8]. Interestingly, ethanol at low concentration causes apoptosis preferentially [9-11]. At such low concentration, ethanol-induced apoptosis in human hepatocellular carcinoma (HepG2) cells is triggered by activation of the Fas receptor [12], a member of the tumor necrosis family, with subsequent activation of the intracellular adapter protein FADD (Fas Associated Death Domain) and caspase-8 [13, 14]. Caspase-8 represents the key component of ethanol-induced apoptosis when applied at low concentration, as confirmed by completely suppressing apoptosis after caspase-8 inhibition [12].

Fas receptor activation has been involved not only with the apoptotic process, but also with triggering of other intracellular signaling pathways [15, 16]. These include the mitogen-activated protein kinase (MAPK) pathway [13, 17-19], and the transcription factor NFκB signal cascade [20, 21]. However, the effect of low concentration of ethanol in the intracellular signaling pathways activated by the Fas receptor pathway has not been documented. Therefore, the aim of the present study was to determine the effect of ethanol exposure at low concentration (1 mM) on protein kinases phosphorylation in human hepatocellular carcinoma (HepG2) cells. In addition, the effect of protein kinases phosphorylation on the expression of transcription factors and subsequent gene expression was determined.

Low concentration of ethanol seems to have a potential therapeutic effect for the treatment of human...
hepatocellular carcinoma; however, its mechanisms of action remain to be determined. This work represent the first step in our understanding of the pathophysiological mechanisms associated with ethanol exposure at low concentration, which are needed to establish its use as a potential target for the treatment of human hepatocellular carcinoma.

2. Methods

Human hepatocellular carcinoma (HepG2) cells

HepG2 cells, derived from human hepatocellular carcinoma (obtained from Deutsches Krebsforschungszentrum Heidelberg, Germany) were seeded in 250 ml tissue culture flasks (Falcon-Becton Dickinson, Heidelberg, Germany) at 1x10⁶/ml concentration in 10 ml RPMI-1640 medium supplemented with 10% fetal bovine serum (Boehringer Mannheim, Mannheim, Germany), 100 U/ml penicillin and 100 µg/ml streptomycin (ICN Flow, Meckenheim, Germany) at 37°C in a humidified atmosphere of 7.5% CO₂. After 7 days of cell culture, the cells were harvested with 0.05% trypsin / 0.02% EDTA (Gibco BRL, Eggersheim, Germany). Cells were seeded in 6-well plates (Falcon-Becton Dickinson, Heidelberg, Germany) at concentrations of 1x10⁶/ml. Six sets of experiments were performed (n=6). Each set consist of two groups as follow: HepG2 cells treated with 1 mM ethanol and HepG2 cells without ethanol treatment as a control.

The rationale for using a low concentration of ethanol, namely 1 mM, was based on previous reported studies in which the potential therapeutic effect of such concentrations has been proposed [11]. In addition, human hepatocellular carcinoma (HepG2) cells were studied without comparing against normal hepatocytes because of the way HepG2 cells metabolize ethanol and because of the selective induction of apoptosis without necrosis observed in HepG2 cells exposed to 1 mM ethanol [9], which has not been observed in normal hepatocytes.

Assessment of protein kinases phosphorylation

For protein kinase phosphorylation studies, HepG2 cells were exposed to low ethanol concentration (1 mM; Merck, Darmstadt, Germany) for 10 min. After treatment, total cell lysates were prepared as described by Zhang et al. [22]. Briefly, cells were washed twice with ice-cold phosphate-buffered saline (PBS; Gibco BRL, Eggersheim, Germany); scraped in lysis buffer (20 mM Tris, 20 mM β-glycerophosphate, 150 mM NaCl, 3 mM EDTA, 3 mM EGTA, 1 mM Na₃VO₄, 0.5% Nonidet P-40, and 1 mM dithiothreitol); supplemented with 1 mM phenylmethanesulfonyl fluoride, 2 µg/ml leupeptin, 4 µg/ml aprotinin, and 1 µg/ml pepstatin A; and sonicated for 15 sec. Cell debris was removed by centrifugation at 1400 x g for 30 min at 4°C. Protein concentration was determined by the Bradford assay [23]. Protein kinase phosphorylation state was assessed using phosho-antibody screening KPKS-1.0 (Kinexus Bioinformatics Corporation, Vancouver, Canada). For this, the Kinetikworks protocol was used [24]. Briefly, 300 µg of total protein was blotted on a 13% single lane SDS-polyacrylamide gel and transferred to nitrocellulose membrane. By using a 20-lane multiblitter from Bio-Rad (Munich, Germany), the membrane was incubated with different mixtures of up to three antibodies per lane that react with a 75 known phosphorylated cell signaling proteins of distinct molecular masses. After further incubation with a mixture of relevant horseradish peroxidase-conjugated secondary antibodies (Santa Cruz Biotechnology, Heidelberg, Germany), the blots were developed using ECL Plus reagent (Amersham Biosciences, Freiburg, Germany), and signals were quantified using Quantity One software (Bio-Rad). The obtained values were normalized to untreated HepG2 cells and the phosphorylation change in percent was calculated. All values showing an increase or decrease equivalent to 25% or more were considered significant.

Assessment of ethanol-induced apoptosis by DNA fragmentation after GPCR inhibition

Inhibition of G-protein coupled receptors (GPCR) was performed using heparin (0.10 IU/ml) or pertussis toxin (20 pM). All chemicals were purchased from Sigma Aldrich (Seelze, Germany). HepG2 cells were pre-treated for 30 min with GPCR inhibitors followed by a washed step and incubated with ethanol (1 mM) for 24 h. Then, ethanol-induced apoptosis was analyzed by a double-fluorescence staining technique with Hoechst 33342 (excitation 330-380 nm, emission 460 nm; Molecular Probes, MoBiTec, Göttingen, Germany) and propidium iodide (excitation 590 nm and emission 620 nm; Molecular Probes, MoBiTec, Göttingen, Germany) as described previously [25]. Briefly, after 24 hours of ethanol incubation, 20 µg/ml propidium iodide and 100 µg/ml Hoechst 33342 were incubated for 15 min at 37°C in the dark. After staining, the cells were immediately examined using a Leitz DM-IRB fluorescence microscope. The numbers of cells with apoptosis-associated alterations of the nuclei and without membrane barrier dysfunction were determined within a field of view at a magnification of X400. A total of 10 randomly selected fields were counted per well. The numbers of altered cells were averaged an expressed as percentage of total cells.

Quantitative real-time polymerase chain reaction

Quantitative real-time PCR was used to evaluate the effect of ethanol on mRNA expression level of different transcription factors as well as the expression of genes of the Fas receptor signaling pathway, including Fas receptor, Fas ligand, FADD and caspase 8. Total RNA was isolated from HepG2 cells using RNAsy kit (Qiagen) and RNA quality was evaluated using Agilent RNA 6000 Nano Chip Kit and Bioanalyzer 2100 (Agilent, Böbligen, Germany). Real-time RT-PCR was performed using the QuantiTect SYBR green RT-PCR kit (Qiagen, Hilden, Germany). Specific primers for transcription factor (AP1, SRF, Elk1, Stat1 and NFκB) and members of the Fas receptor signaling pathway (Fas receptor, Fas...
Primer design was performed with the Primer Express 2.0 software from ABI Prism, Applied Biosystems (Darmstadt, Germany) and obtained from MWG-Biotech AG (Ebersberg, Germany). Fas receptor forward, 5'-CTT TCC GTG AGC TCG TCT CTG A-3'; Fas receptor reverse, 5'-CTC CCC AGA AGC GTC TTT GA-3'; Fas ligand forward, 5'-CCA GCT TGC CTC CTC GTG AG-3'; Fas ligand reverse, 5'-TCC TGT AGA GCC TGA GGT TTC TTT GA-3'; FAAD forward, 5'-GGT GGA GAA CTG TGC CTC CTC TTG AG-3'; FAAD reverse, 5'-GCA AAA GCA CGG GAG AAA GT-3'; caspase 8 forward, 5'-CCA GCT TGC CTC CTC TTG AG-3'; Fas ligand reverse, 5'-TCC TGT AGA GCC TGA GGT TTC TTT GA-3'; FAAD forward, 5'-GGT GGA GAA CTG TGC CTC CTC TTG AG-3'; caspase 8 reverse, 5'-CCA AAA GCA CGG GAG AAA GT-3'; caspase 8 forward, 5'-CCA GCT TGC CTC CTC TTG AG-3'; and caspase 8 reverse, 5'-TCC TGT AGA GCC TGA GGT TTC TTT GA-3'. Quantitative real-time PCR determination using the Optical System Software (iQ5 version 1.0) provided with the BioRad iQ5 cycler (BioRad, Munich, Germany) was performed.

### Statistical Analysis

Data are expressed as mean values ± standard deviation (SD). Results from HepG2 cells treated with 1mM concentration of ethanol were compared to non-treated HepG2 cells (control cells) using Student's t-test. Statistical significance was assumed at p level <0.05 level. SigmaPlot software version 8.02 (Systat Software, Erkrath, Germany) was used for statistical analysis.

### Results

#### Short exposure to 1 mM ethanol induces phosphorylation of protein kinases

Table 1 shows the effect of 1 mM ethanol on protein kinases phosphorylation in HepG2 cells after 10 min exposure time compared to untreated cells. Ethanol caused a strong inhibition of the GPCR signaling pathway as shown by phosphorylation of GRK2 and PKCα; with values of 109% and 104%, respectively, combined with dephosphorylation of ROKa, PKCd and PKCu; with values of -56%, -44% and -28% of control cells, respectively. These findings resulted in a pro-apoptotic effect that was intensified by a slight inhibition of the JNK and the NFκB signaling pathway as demonstrated by a reduced phosphorylation of MEK4, IKKα, JNK and MEK6; with values of -36%, -33%, -27% and -26%, respectively. The pro-apoptotic effect was also induced by a slight activation of the cell death receptor signaling pathway with increased phosphorylation of DAPK3 and DAPK1 equivalent to 56% and 38%, respectively. On the other hand, an anti-apoptotic effect was also activated, as shown by an increased phosphorylation of members of the ERK and the CDK signaling pathways, including ERK2, RSK2, CDK9, CDK6, CDK7 and RSK1 with values of 50%, 48%, 38%, 35%, 29%, 26%, and 26%, respectively.

### Table 1 Effect of 1 mM ethanol on various signaling transduction pathways leading to apoptosis

| Protein | Abbreviation | Type | Phosphorylation % of control | Ethanol effect |
|---------|--------------|------|------------------------------|---------------|
| G-protein coupled receptor signaling | GRK2 | PSTK | +109 | strong inhibition |
| Protein kinase C alpha | PKCa | PSTK | +104 | pro-apoptotic |
| RhoA kinase alpha | ROKa | PYK | -56 | |
| Protein kinase C delta | PKCd | PSTK | -44 | |
| Protein kinase C mu | PKCu | PSTK | -29 | |
| Mitogen-activated protein kinase signaling | |
| ERK signaling pathway | ERK2 | PSTK | +50 | |
| Extracellular regulated kinase 2 | RSK2 | PSTK | +48 | |
| Ribosomal S6 kinase 2 | RSK1 | PSTK | +26 | |
| JNK signaling pathway | MEK4 | PTYK | -36 | |
| MAP kinase kinase 4 | MEK6 | PTYK | -26 | |
| MAP kinase kinase 6 | |
| NFκB signaling | IKKα | PSTK | -33 | |
| Cyclin-dependent signaling | CDK9 | PSTK | +38 | |
| Cyclin-dependent kinase 9 | |
| Cyclin-dependent kinase 6 | CDK6 | PSTK | +35 | |
| Cyclin-dependent kinase 7 | CDK7 | PSTK | +29 | |
| Cell death receptor signaling | |
| Death associated protein kinase 3 | DAPK3 | PSTK | +50 | |
| Death associated protein kinase 1 | DAPK1 | PSTK | +38 | |

PSTK = protein-serine/threonine kinase, PYK = protein-tyrosine kinase, PTYK = protein-threonine/tyrosine kinase

[apoptotic effect]
Neutralization of membrane receptors blocks ethanol-induced apoptosis

Based on the strong inhibition of the G-protein coupled receptor signaling pathway observed after ethanol exposure, we analyzed further the role of this pathway in ethanol-induced apoptosis. For that purpose, experiments with specific inhibitors with subsequent determination of the apoptotic rate were carried out. As shown in Figure 1, we found that ethanol-induced apoptosis after ethanol exposure was increased by 33% compared to control HepG2 cells (without ethanol exposure). In contrast, specific inhibitors of the G-protein coupled receptor, such as heparin and pertussis toxin, resulted in increased apoptosis with values of 45% and 70%, respectively. These data confirm the role of G-protein coupled receptors as a regulatory mechanism of ethanol-induced apoptosis.

![Figure 1](image1.png)

Figure 1 | Effect of inhibitors of G-protein coupled receptors (GPCR) on ethanol-induced apoptosis in HepG2 cells. Results are expressed as percent of control HepG2 cells (without ethanol exposure). Results are the mean of six different experiments (n=6). Error bars represents standard deviations. Significance is shown as the difference between ethanol-treated and control cells * p < 0.05.

Ethanol-induced protein kinases phosphorylation activates the expression of transcription factors

Figure 2 shows the effect of a millimolar ethanol concentration on mRNA levels of AP1, Elk1, Stat1, SRF and NFκB after 2, 4 and 6 hours of ethanol exposure using relative quantitative real time PCR. The transcription factor AP1 did not reveal any significant difference after 2, 4 and 6 hr ethanol exposure compared to controls. The mRNA expression level of Elk1 and Stat1 showed time dependent increases in mRNA expression levels of approximately 1-fold after 2 h of ethanol exposure, and 2-fold after 6 hr exposure. SRF was significantly increased after 2 hr by 7-fold with subsequent decrease to about only 1-fold after 4 hr and 6 hr. In contrast, NFκB was significantly increased after 2 hr of ethanol exposure with a maximal effect seen after 6 hr, representing increases of 1- and 12-fold, respectively. These data confirm the effect of ethanol on intracellular signal transduction resulting in increased expression of transcription factors.

![Figure 2](image2.png)

Figure 2 | Time course changes in mRNA levels of transcription factors in HepG2 cells exposed to 1mM ethanol concentration expressed as a relative fold change compared to control HepG2 cells (not exposed to ethanol). White, gray and black bars represent 2, 4 and 6 hr exposure times, respectively. Results are the mean of six different experiments (n=6). Error bars represents standard deviations. Significance is shown as the difference between ethanol-treated and control cells at 2 h (* p < 0.05), 4 h († p < 0.05), and 6 h (# p < 0.05).

Ethanol regulates the expression of genes of the Fas signaling cascade

To further investigate the role of ethanol-induced apoptosis on gene expression, the mRNA level of Fas receptor, Fas ligand, FADD, and caspase 8, were examined using relative quantitative real time PCR. As shown in Figure 3,
HepG2 cells exposed to ethanol for 6 h showed significantly increased mRNA expression levels for Fas receptor, Fas ligand, FADD and caspase 8; equivalent to 1.3, 1.8, 3.7 and 7.3-fold, respectively, when compared to cells non treated with ethanol. Fas receptor expression was similar after 6 and 24 hr, while Fas ligand expression significantly increased by 2.7-fold after 24 hr of ethanol exposure. In contrast, mRNA expression of FADD and caspase 8 were significantly decreased by 2.2 and 5.6, respectively, after 24 hr exposure of ethanol.

Figure 3 mRNA expression levels of members of the Fas signaling cascade in HepG2 cells exposed to 1mM ethanol concentration expressed as a relative fold change compared to control HepG2 cells (not exposed to ethanol). White and black bars represent 6 and 24 hr exposure times, respectively. Results are the mean of six different experiments (n=6). Error bars represents standard deviations. Significance is shown as the difference between ethanol-treated and control cells at 6 h (*p < 0.05) and 24 h (†p < 0.05).

4. Discussion
The working hypothesis of the present study was that apoptosis induced by low concentration of ethanol (namely 1 mM) in HepG2 cells is regulated through the interaction of both pro-apoptotic and anti-apoptotic signaling pathways. The pro-apoptotic effect induced by ethanol was demonstrated by a strong inhibition of GPCR signaling pathway in association with a slight inhibition of JNK and NFκB signaling in association with a slight activation of the cell death receptor signaling. In addition, ethanol also phosphorylated protein kinases belonging to signal transduction pathways with an anti-apoptotic effect such as ERK and CDK. These data, corroborate that ethanol-induced phosphorylation of protein kinases modulates both anti-apoptotic and pro-apoptotic mechanisms, and suggests that ethanol at low concentration shifts the balance into the apoptotic direction after its initial effect on protein kinases phosphorylation.

The most strikingly and novel finding obtained hereby was the strong inhibition of the GPCR signaling pathway induced mainly by increased phosphorylation of GRK2. This finding suggests an important regulatory role of ethanol-induced apoptosis through early phosphorylation of GRK2. Our data correlates with the reported involvement of this protein kinase in the regulation of signal transduction initiated through the GPCR [26, 27]. Furthermore, the regulatory mechanism of ethanol-induced apoptosis through the GPCR signaling pathway was also confirmed by the increased apoptotic rate observed after neutralization of GPCR. These findings provide a novel insight into the molecular mechanism of action exerted by the exposure of 1 mM ethanol concentration in human hepatocellular HepG2 cells.

We found that ethanol-induced phosphorylation of protein kinases lead to an increased expression of transcription factors (AP1, Elk1, Stat1, SRF and NFκB). This finding correlates with the reported activation of transcriptions factors through the different signaling pathways [28-32]. In addition, the observed increased expression of transcription factors subsequently resulted in an induction of gene expression, which seems to be also regulated through the interaction of various intracellular signaling pathways. It has been reported that MAPK signal cascade induces Fas ligand expression as well as Fas ligand promoter activation in T lymphocytes [33]. Our data confirmed the increased expression of genes associated with the Fas-signaling cascade (including Fas receptor, Fas ligand, FADD and caspase 8).

The NFκB signaling pathway is known to be one of the key regulators of anti-apoptotic processes observed in human hepatocytes derived cell lines [34, 35]. Studies performed in T cells have demonstrated
that the NFκB signaling pathway mediates cell growth and protection against apoptotic cell death [36]. The kinase IKKα plays a key step in the activation of NFκB [37]. Our data demonstrated that ethanol inhibits the anti-apoptotic effect of the NFκB signaling pathway by reducing the phosphorylation of IKKα. Moreover, the increased expression of mRNA levels of NFκB after 6 hr of ethanol exposure, suggests a regulatory role of this pathway in ethanol-induced apoptosis.

Taking these data together, we propose a regulatory mechanism of ethanol-induced apoptosis in HepG2 cells (as described in Figure 4 and 5). This regulatory mechanism results from the balanced interaction between pro-apoptotic and anti-apoptotic effects of various intracellular signaling pathways. Specifically, ethanol at low concentration induced a biphasic effect in HepG2 cells, characterized by a rapid transient effect on protein kinases, after 10 min exposure, followed by increased expression of transcription factors up to 6 hr (early events phase) and a second persistent activation of gene expression that began after 6 hr and persisted for more than 24 hr (late events phase). The early events phase was characterized by significant change in the phosphorylation state of protein kinases with subsequent induction of transcription factors. Such changes were associated with either inhibition or activation of various intracellular signaling pathways leading to a pro-apoptotic effect. These findings suggest that low concentration of ethanol shifts the balance into the apoptotic direction after its initial effect on protein kinases. The second phase or late events, between 6 and 24 hr is characterized by expression of Fas-associated genes (including Fas receptor, Fas ligand, FADD and caspase 8).

**Figure 4** Regulatory mechanism of low concentration of ethanol resulting from the balanced interaction between pro-apoptotic and anti-apoptotic effects of various intracellular signaling pathways.

![Diagram](https://example.com/diagram.png)
In this study, treatment with 1mM ethanol concentration induced up-regulation of caspase 8, which is the key molecule of ethanol-induced apoptosis through Fas receptor activation. These data corroborate the increased activity of caspase 8 previously reported after 12 hr of ethanol exposure [12]. This finding suggests that low concentration of ethanol plays a very important role in the modulation of ethanol-induced apoptosis through simultaneous induction of anti- and pro-apoptotic events. Thus, apoptosis induced by a low concentration of ethanol seems to be regulated by protective mechanisms resulting from activation of anti-apoptotic pathways. Moreover, the interaction between both pro- and anti-apoptotic processes, induced by short-term exposure of ethanol at low concentration, does represent an important new mechanism to consider in the pathophysiology of ethanol-induced effects on HepG2 cells.

In conclusion, the data obtained in the present study support our working hypothesis and confirm a balance between pro- and anti-apoptotic effects. Ultimately, the fine balance between these two processes would lead to a controlled regulation of ethanol-induced apoptosis in HepG2 cells exposed to low concentrations of ethanol. The findings of the present study provide new evidence that ethanol at low concentration is not simply an activator of Fas receptor-induced apoptosis. This model represents the basis for further studies addressing the pathophysiological mechanisms of exposure to low concentration of ethanol in HepG2 cells, and provides information about potential targets for the treatment strategies of human hepatocellular carcinoma.

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Conflicts of interest
The authors have declared that no conflict of interest exists.

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