Research Paper

MZF-1/Elk-1/PKCα is Associated with Poor Prognosis in Patients with Hepatocellular Carcinoma

Je-Chiuan Ye1, Li-Sung Hsu2,3, Jen-Hsiang Tsai4, Hsin-ling Yang5, Meen-Woon Hsiao6, Jin-Ming Hwang6, Chia-Jen Lee7, Jer-Yuh Liu8,9

1. Bachelor Program of Senior Services, Southern Taiwan University of Science and Technology, Tainan, Taiwan;
2. Institute of Biochemistry and Biotechnology, Medical College, Chung Shan Medical University, Taichung, Taiwan;
3. Clinical Laboratory, Chung Shan Medical University Hospital, Taichung, Taiwan;
4. Department Physical Therapy, School of Medical and Health Sciences, Fooyin University, Kaohsiung, Taiwan;
5. Institute of Nutrition, College of Biopharmaceutical and Food Sciences, China Medical University, Taichung, Taiwan;
6. Department of Medical Applied Chemistry, College of Health Care and Management, Chung Shan Medical University, Taichung, Taiwan;
7. Department of Medical Research, Tungs’ Taichung MetroHarbor Hospital, Taichung, Taiwan;
8. Center for Molecular Medicine, China Medical University Hospital, Taichung, Taiwan;
9. Graduate Institute of Biomedical Sciences, China Medical University, Taichung, Taiwan.

Corresponding authors: Jer-Yuh Liu PhD., Graduate Institute of Cancer Biology, College of Medical, China Medical University, No 6, Hsueh-Shih Road, Taichung 404, Taiwan. Tel: +886-4-22052121 ext. 7932; Fax: +886-4-22333496; E-mail: jyl@mail.cmu.edu.tw Chia-Jen Lee PhD., Department of Medical Research, Tungs’ Taichung MetroHarbor Hospital, No.699, Sec. 8, Taiwan Blvd., Taichung City 435, Taiwan. Tel: +886-4-26581919 ext. 79004; E-mail: chiajenlee54@gmail.com

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Abstract

Background: Protein kinase C alpha (PKCα) is a key signaling molecule in human cancer development. As a therapeutic strategy, targeting PKCα is difficult because the molecule is ubiquitously expressed in non-malignant cells. PKCα is regulated by the cooperative interaction of the transcription factors myeloid zinc finger 1 (MZF-1) and Ets-like protein 1 (Elk-1) in human cancer cells.

Methods: By conducting tissue array analysis, herein, we determined the protein expression of MZF-1/Elk-1/PKCα in various cancers.

Results: The data show that the expression of MZF-1/Elk-1 is correlated with that of PKCα in hepatocellular carcinoma (HCC), but not in bladder and lung cancers. In addition, the PKCα down-regulation by shRNA Elk-1 was only observed in the HCC SK-Hep-1 cells. Blocking the interaction between MZF-1 and Elk-1 through the transfection of their binding domain MZF-160–72 decreased PKCα expression. This step ultimately depressed the epithelial-mesenchymal transition potential of the HCC cells.

Conclusion: These findings could be used to develop an alternative therapeutic strategy against patients with the PKCα-derived HCC.

Key words: MZF-1; Elk-1; PKCα; HCC.

Introduction

Protein kinase C alpha (PKCα) is a member of the protein kinase C family, consisting of at least ten isoforms (α, βI, βII, γ, δ, ε, η, θ, and ζ and ι), that regulates multiple biological processes, including cell proliferation, apoptosis, differentiation, migration, and adhesion [1]. The enhanced expression of PKCα has been reported in the tumor tissues of various cancer types [2–8]. Such expression has been demonstrated to promote tumor growth and metastasis [9]. Thus, developing therapeutic agents that target PKCα has become the focus of many research laboratories [10]. However, the off-target effects of targeting PKCα and the limited understanding of the signaling mechanisms upstream
of PKCα have hampered this effort.

Several mechanisms that contribute to PKCα overexpression have been investigated. These mechanisms include I) the shift in signaling from the epidermal growth factor receptor to the platelet-derived growth factor receptor during the progression from non-stem cells to cancer stem cells (CSCs) [11], II) the epithelial–mesenchymal transition (EMT), which results in the dominance of pro-invasive pathways downstream G-protein receptors [12], and III) the overexpression or activation of ErbB2 [13]. We recently discovered that the transcription factors Ets-like protein-1 (Elk-1) and myeloid zinc finger-1 (MZF-1) regulate PKCα expression in human hepatocellular carcinoma (HCC) cells [14–17]. These transcription factors are also direct upstream inducers of the expression of the PKCα protein and mRNA; the former molecules bind to the PKCα promoters. The Elk-1 transcription factor activates the c-fos promoter by associating with serum response factors; moreover, Elk-1 is a target of both extracellular-signal-regulated kinase and c-Jun N-terminal kinase cascades [18]. The transcription factor also controls the expression of genes involved in cell-cycle progression, differentiation, and apoptosis in response to extracellular signals [19–21].

MZF-1 belongs to the Kruppel family of zinc finger proteins and is preferentially expressed in myeloid progenitor cells [22–25]. MZF-1 plays an important role in cell growth, differentiation, and apoptosis.

We previously found that knocking down Elk-1 and/or MZF-1 results in significantly decreased PKCα mRNA and protein expression [14]. This finding suggests that the transcription factors regulate PKCα cooperatively. In the present study, we investigated the correlation between MZF-1/Elk-1 and PKCα expression in human HCC and explored a peptide-based strategy that inhibits EMT in malignant cells.

**Materials and Methods**

**Immunohistochemical (IHC) staining**

Array slides (HCC BS03014, lung cancer LC10011, and bladder cancer BL482) were purchased from US Biomax, Inc. (Rockville, MD, USA). The BS03014 slides included 60 cases of carcinoma, but 1 case was lost during evaluation. The LC10011 slides included 40 cases of non-small cell carcinomas, but 5 cases were lost during evaluation. The BL482 slides included 48 cases of transitional cell carcinomas. Detailed information on this array can be viewed at http://www.biomax.us/tissue-arrays/.

The slides were deparaffinized in xylene and rehydrated in an alcohol series. The sections were then incubated with 3% H2O2 for 5 min. After washing with PBS, the sections were boiled in EDTA solution (1 mM EDTA, 0.1% NP-40; pH 8.0) for 5 min (for PKCα) or in citric acid solution (10 mM citric acid monohydrate; pH 6.0) for 15 min (for Elk-1 and MZF-1 detection) in a microwave oven. After cooling for 1 h, the sections were washed thrice in PBS for 5 min. Then, the sections were incubated in PBS with 5% normal bovine serum for 25 min. The sections were washed with PBS and incubated with antibodies against PKCα (10 ng/ml PBS plus 0.2% BSA) (Sigma-Aldrich, St Louis, MO), Elk-1 (1:400) (Santa Cruz, CA), and MZF-1 (1:400) (Santa Cruz) at 4 °C overnight. After washing thrice with PBS for 5 min, the sections were incubated with biotinylate-labeled goat anti-rabbit IgG or rabbit anti-mouse IgG (Sigma-Aldrich) at room temperature for 1 h. Then, the sections were washed with PBS and incubated with peroxidase-conjugated ABC reagent (Avidin/Biotin kit, Vector Laboratories, Inc., Burlingame, CA, USA) at room temperature for 30 min. The sections were visualized by adding 3,3′-diaminobenzidine substrate (Sigma-Aldrich). The reaction was terminated by rinsing the sections with distilled water. The sections were counterstained with Gill’s hematoxylin V (Mute Pure Chemicals Ltd., Tokyo, Japan) and dehydrated in an alcohol series. Afterwards, the sections were cleared with xylene before mounting with Malinol (Muto Pure Chemicals Ltd., Tokyo, Japan) and examined under a BX40 system microscope (Olympus, Tokyo, Japan) with a CCD DRII camera (Olympus). Resulting images were analyzed using Image-Pro® Plus software (Media Cybernetics, Silver Spring, MD, USA).

Plasmid construction

Plasmids containing different fragments of MZF-1-c-Myc (encoding amino acids 60–72) were amplified from pcDNA-MZF-1-c-Myc by PCR (25). Then, the PCR products were isolated and cloned into a pcDNA™ 3.1/myc-His vector (reverse-transcription PCR and cloned into the pcDNA™ 3.1/myc-His vector [Invitrogen]).

**Cell lines**

Cancer cells from various human organs were obtained directly from the ATCC (Manassas, VA, USA). The liver cancer cells include HCC HA22T (BCRC no.60168), Hep3B (BCRC no.60434), HepG2 (BCRC no.RM60025), SK-Hep-1 (ATCC no. HTB-52), and Huh-7 (ATCC no. JCRB-0403) cells. Meanwhile, the lung cancer cells were A549 (ATCC no. CCL185),
H322 (ATCC no. CRL-5806), H1299 (ATCC no. CRL-5803), and H928. All the cells were cultured in media specific to each cell line and supplemented with 10% fetal bovine serum, 100 units/mL penicillin G, and 100 µg/mL streptomycin (Gibico, Grand Island, NY, USA) in a humidified atmosphere containing 5% CO₂ at 37°C.

**Transfection and stable clone establishment**

Lipofectin was used for transfection. Cells were cultured in 60 mm dishes containing minimum essential medium (MEM) supplemented with 10% fetal calf serum (FCS-MEM) at 37 °C for 24 h before rinsing with serum-free MEM. Then, the sample was transferred to 1 mL serum-free MEM containing 15 µg/mL Lipofectamine 2000 transfection reagent (Invitrogen) and various doses of the indicated plasmid. After incubating for a minimum of 6 h, 1 mL MEM supplemented with 20% FCS was added to the medium. After incubating for another 18 h, the medium was replaced with fresh FCS-DMEM. Then, the cells were incubated for at least 48 h before they were lysed for subsequent assays.

Stable clones were established by seeding low-passage cells at a density of 3 × 10⁵ cells in 60mm tissue culture dishes. The cells were transfected with the MZF-160–72 plasmid (5 µg/6 mL) using tissue culture dishes. The cells were transfected with medium was replaced with fresh FCS-DMEM. Then, medium. After incubating for another 18 h, the cells were incubated for at least 48 h before they were lysed for subsequent assays.

Individual clones were then transferred to 96-well plates and grown until confluence. After being transferred to flasks, the cells were cultured until confluence, harvested, and frozen in liquid nitrogen for further experimentation.

**Immunoblotting analysis**

Cancer cell lysates were lysed in radioimmunoprecipitation assay (RIPA) buffer (150 mM NaCl, 5 mM EDTA, 50 mM HEPES [pH 7.5], 0.5% [w/v] sodium deoxycholate, 1% Nonidet P-40 [NP-40], 10 mM 2-mercaptoethanol) containing 2 mM phenylmethylsulfonyl fluoride (PMSF), 50 mg/mL aprotinin A, 25 mg/mL leupeptin, and 25 mg/mL pepstatin. The cell lysates were then centrifuged at 12000 × g for 5 min and maintained on ice. The cell lysates were resolved by 10% sodium dodecyl sulfate–polyacrylamide gel electrophoresis, after which the proteins were transferred to polyvinylidene fluoride membranes (Immobilon-P; Millipore, Bedford, MA, USA). The membranes were then incubated in blocking buffer (5% [w/v] non-fat dry milk, 0.1% [v/v] Tween 20) in Tris-buffered saline (TBST) at room temperature for 30 min. Next, the membranes were probed with the following specific antibodies: anti-PKCα (BD Biosciences, San Jose, CA, USA); anti-E-cadherin (CDH1); anti-VIM and anti-SNAI2l (Cell Signaling, Beverly, MA, USA); anti-p38 MAPK (p38), anti-phospho-p38 MAPK (P-p38), and anti-urokinase-type plasminogen activator (uPA) (GeneTex, Inc., Irvine, CA, USA); anti-Elk-1, anti-MZF-1; and anti-β-actin polyclonal antibodies (Santa Cruz). Then, the blots were incubated in blocking buffer at 4 °C overnight, after which they were incubated with horseradish peroxidasedelabeled anti-mouse or anti-rabbit secondary antibodies (Promega) at room temperature for 2 h. After three washes with TBST buffer, antibody-reactive proteins were detected using a chemiluminescent substrate (Pierce, Rockford, IL, USA).

**Antisense knockout assays**

A shRNA Elk-1-expressing plasmid vector was constructed using the pcDNA-HU6 vector (a gift from Dr. J. Tsai Chang, Institute of Toxicology, College of Medicine, Chung Shan Medical University, Taichung, Taiwan) as the vector backbone. The shRNA Elk-1 duplex sequence, which was obtained from human Elk-1 genes (GenBank, NCBI), was designed using the BLOCK-iT™ RNAi Designer software available at http://www.invitrogen.com. The sequence corresponded to the coding regions relative to the first nucleotide of the start codon. The sequences designed to produce hairpin RNAs identical to the oligonucleotide shRNA duplex sequences were as follows: sense, 5′-GATCCGCAAGAACAAGACCAA CATTTCAAGAGAT–3′ and antisense, 5′-AGCTTA AAAAGCAAGAACAAGACCAA CATTTCAAGAGAT–3′. To generate the shRNA duplex, 40 µM each of sense and antisense oligonucleotides were annealed in the thermocycler using the following profile: 37°C for 30 min and 65°C for 15 min. The resulting shRNA duplex was then cloned into the pcDNA-HU6 vector in frame with the BamHI and HindIII sites. The insert was screened by PCR, with the HU6 primer and confirmed by sequencing with the HU6 primer. After transfection for 6 h, the cells were washed thrice in serum-free MEM and allowed to recover for 24 h in fresh medium. Stable clones were selected with geneticin (G418; 600 µg/ml) at 37°C for 4 weeks.

**Statistical analysis**

Data were expressed as mean ± standard deviation and were analyzed by ANOVA. Pearson’s chi-squared test [26] and Student’s t-test were used in
two-group comparisons. Here, $p < 0.05$ was considered as statistically significant.

**Results**

**PKCα expression is correlated with MZF-1/Elk-1 expression in HCC**

To determine the clinical relevance of the correlation between PKCα and Elk-1 and/or MZF-1, we examined the expression of these proteins in tissue microarrays of liver, lung, and bladder cancers by IHC staining. We observed a positive correlation between moderate-to-strong PKCα staining and either Elk-1 and/or MZF-1 staining in HCC (Figure 1) but not in lung or bladder cancers (Figures 2 and 3, respectively). Moreover, the moderate-to-strong staining of PKCα/Elk-1/MZF-1 was most common in the grades 2 and 3 HCCs (Figure 1). We also validated these results in the HCC cells (Figure 4), and found that high PRKCA expression levels were significantly correlated with those of Elk-1 and MZF-1 in poorly differentiated HCC (r = 0.86, $p < 0.05$ and r = 0.92, $p < 0.05$, respectively). Reducing Elk-1 expression by siRNA gene silencing decreased the PKCα protein expression in the poorly differentiated HCC SK-Hep-1 cells (Figure 5). Collectively, these results suggest that PKCα, along with Elk-1 and MZF-1, function as important mediators of tumor progression in HCC.

**Figure 1.** Correlations between the expression of PKCα and Elk-1/MZF-1 in human HCC. IHC analyses and correlations of PKCα and Elk-1/MZF-1 expression in human HCC. The up-panel shows the representative staining results for the samples scored by visual assessment as “weak,” “moderate,” or “strong” on the basis of staining intensity. The label (i.e. F4) at the bottom-left corner of each sample is the serial number of the patients indicated by US Biomax, Inc. (Rockville, MD, USA), and can be viewed at http://www.biomax.us/tissue-arrays/. The numbers of each group classified on the basis of PKCα, Elk-1, or MZF-1 staining intensity or grade are depicted in the down-panel. A positive rating was given to the moderate or strong expression of the genes of interest; otherwise, a negative rating was given. The clinical characteristic grades of I, II, and III were obtained from US Biomax, Inc. *$p < 0.05$, Pearson’s chi-squared test.
Inhibition of MZF-1 and Elk-1 heterodimer formation attenuates the malignant phenotypes of HCC cells by reducing PKCα expression

In our previous work, we reported that the expression of MZF-1ΔDBD (contains MZF-160–72 but lacks the DNA-binding domain) or Elk-1ΔDBD (contains Elk-1145–157 but lacks the DNA-binding domain) decreased PKCα expression, cell migration and invasion, and cell proliferation in HCC cells [27]. Compared with the tumors that developed in the mice injected with empty-vector-treated cancer cells, the tumors in the mice with MZF-1ΔDBD-treated cancer cells were smaller throughout almost the entire experimental time interval. Furthermore, the MB-231 and Hs578T breast cancer cells stably expressing MZF-160–72(MB-231-M(v3), MB-231-M(v4), Hs578T-M(s2), and Hs578T-M(s3)) were more rounded than the elongated parental and vector control cells [28]. These observations suggest that MZF-160–72 hindered the endogenous Elk-1 and MZF-1 interaction, subsequently moderated the transcription factors’ binding to the PRKCA promoter, and ultimately inhibited PKCα and EMTcore genes.

Next, we examined the effects of MZF-160–72 on HCC cells. The changes in the protein expressions of these EMTcore genes were also similar to those in the SK-Hep-1 HCC cells stably expressing MZF-160–72(Figure 6). The changes in the HCC SK-Hep-1 cells were consistent with those observed in the cells with PKCα knockdown by transfection with shRNA [28]. The expressions of PKCα, phosphorylated-mitogen-activated protein kinase (MAPK) p38, and urokinase-type plasminogen activator (uPA) were reduced. The expression of the EMTcore genes was also altered.
Figure 3. Correlations between the expression of PKCα and Elk-1/MZF-1 in human bladder cancer. IHC analyses and correlations of PKCα and Elk-1/MZF-1 expression in human bladder cancer. The representative staining results, labels, and clinical characteristic grades for the samples were described as in Figure 1. *p<0.05, Pearson's chi-squared test.

Figure 4. PKCα expression correlated with MZF-1/Elk-1 expression in HCC cell lines. Immunoblotting analysis of the protein levels of PKCα, Elk-1, and MZF-1 in HCC and lung cancer cells.
Discussion

We previously demonstrated that MZF-1 and Elk-1 interact cooperatively to regulate PKCα expression [14, 26, 29], and a decrease in either endogenous MZF-1 or Elk-1 level affects PKCα expression, cell migration activity, and tumorigenesis [14–17]. We further identified specific domains within MZF-1 and Elk-1, which are responsible for the disruption of their interactions. Such disruption leads to decreased DNA binding activity, followed by reduced PKCα expression, and the eventual attenuation of cell migration and EMT potential. Present findings indicate a positive correlation between moderate-to-strong PKCα expression and either Elk-1 and/or MZF-1 staining in HCC. The moderate-to-strong staining of PKCα/Elk-1/MZF-1 was most commonly observed in the grades 2 and 3 HCCs. Thus, introducing peptides that saturate the binding surfaces of the molecules of interest is a potential strategy for developing alternative anti-cancer therapies.

PKCα is an important signaling molecule in the progression of many carcinomas and plays a key role in EMT [11, 30–32]. The fluctuating intensities of stress factors (e.g., hypoxia, inflammation, and the either cooperative or hostile interactions of tumor intercellular networks), which are known to induce EMT, all increase the adaptation potential of cancer cells; such adaptation mechanisms include bypassing cellular senescence and the subsequent development of CSCs [33]. CSCs appear to be responsible for driving tumor growth, recurrence, and metastasis [34, 35]. These stem cells also possess mesenchymal phenotypes associated with highly aggressive cancer traits [36, 37]. Our data support previous findings, which indicate that EMT-inducing agents enriched CSCs and increased PKCα expression. The increase in PKCα expression correlated with increased Elk-1 and MZF-1 expression in the clinical tissue array in the present study. This observation demonstrated that the induction of PKCα overexpression by Elk-1 and MZF-1 expression may have induced the development of high-grade malignancy. However, further studies on the mechanism underlying the increase in Elk-1 and MZF-1 expression are still underway.

Figure 6. EMT reduction by the disruption of MZF-1/Elk-1 heterodimer formation in HCC cell lines. Changes in protein levels in the parental and MZF-160–72-transfected stable HCC SK-Hep-1 cells as detected by immunoblotting analysis. SK-Hep-1-V indicates the control vector-transfected stable cells; SK-Hep-1-M indicates the MZF-160–72 vector-transfected stable cells. The number in the () indicates the designation of the stable-clone cell. β-Actin was used as internal control.
In HCC research, where PKCa expression is higher in human HCC [5], PKCa overexpression is known to increase the molecular potential for activating the mitogen-activated protein kinase (MAPK) p38 signaling pathway [29]. PKCa overexpression is also known to be associated with cell malignancy. Thus, PKCa inhibition decreases the malignancy of HCC cells and suppresses metastasis [38]. Herein, we showed that inhibiting MZF-1/Elk-1 heterodimer formation decreased PKCa expression and MAPK p38 activation. Therefore, considering the high correlation between PKCa and Elk-1/MZF-1 expression in HCC patients, inhibiting the Elk-1/MZF-1 interaction represents a novel and feasible strategy to specifically inhibit PKCa expression, and eventually, tumor development. This approach may also increase the effectiveness of HCC treatment.

Conclusions

PKCa inhibitors are currently being investigated in human clinical trials, both alone and in combination with other modalities [10, 39]. However, formulating a viable treatment strategy, which specifically targets PKCa in cancer cells, is challenging due to the ubiquity of PKCa and potential off-target effects [10, 40]. Peptide-based therapy is currently both under clinical and preclinical stages of development, and this approach is available in the market for treating human diseases [41]. An example of such therapy is TAT-beclin1 [42] derived from beclin1. The peptide has been shown to induce autophagy, decrease the replication of several pathogens (including HIV-1), and increase the survivability of mice from viruses such as chikungunya and the West Nile virus. Another example is the PKCa antagonistic peptide [9], which is derived from a highly variable region V5 of the enzyme. The peptide inhibits intravasation, cell migration, and metastasis; protects against liver damage; and normalizes blood cell count in animal tests. The direct targeting of the Elk-1/MZF-1 interaction by PKCa treatment methods could allow the targeting of PKCa, Elk-1, and MZF-1-expressing malignant cells without any adverse effects on non-cancerous cells that show non-prominent Elk-1 and MZF-1 expression. Moreover, the discovery of the interactive sequences of Elk-1 and MZF-1 could later lead to a new strategy of treating PKCa-derived HCC involving the identification of small molecules that increase drug delivery potential and inhibitory effectiveness in metastasis.

Abbreviations

PKCa: protein kinase C alpha; CSCs: cancer stem cells; EMT: epithelial-mesenchymal transition; Elk-1: Ets-like protein-1; MZF-1: myeloid zinc finger-1; HCC: hepatocellular carcinoma; IHC: immunohistochemical; MAPK: mitogen-activated protein kinase; uPA: urokinase-type plasminogen activator.

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Competing Interests

The authors have declared that no competing interest exists.

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