



5-FU blocks shuttling of HuR mediated by PKC δ in gastric cancer cells

Wenzhi Wu^{1,2#}, Shengnan Xu^{3#}, Kaiyu Guan³, Weiwei Zhang³, Congde Chen⁴, Xiaoming Chen², Zipu Hong⁶, Dong Hua^{1,5}

¹Department of Oncology, The Second Affiliated Hospital of Soochow University, Suzhou, China; ²Department of Gastroenterology, The First Affiliated Hospital of Wenzhou Medical University, Wenzhou, China; ³Institute of Mental Health, The Affiliated Kangning Hospital of Wenzhou Medical University, Wenzhou, China; ⁴Department of Pediatric Surgery, The Second Affiliated Hospital of Wenzhou Medical University, Wenzhou, China; ⁵Department of oncology, The Affiliated Hospital of Jiangnan University, Wuxi, China; ⁶Department of Burn, The Second Affiliated Hospital of Wenzhou Medical University, Wenzhou, China

Contributions: (I) Conception and design: D Hua, Z Hong, W Wu, S Xu; (II) Administrative support: C Chen, X Chen; (III) Provision of study materials or patients: W Wu, S Xu, K Guan, W Zhang, Z Hong; (IV) Collection and assembly of data: D Hua, Z Hong, W Wu, S Xu; (V) Data analysis and interpretation: D Hua, Z Hong, W Wu; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

[#]These authors contributed equally to this work.

Correspondence to: Dong Hua. Department of Oncology, The Second Affiliated Hospital of Soochow University, Suzhou 215004, China. Email: HUADong@csc.org.cn; Zipu Hong. Department of Burn, The Second Affiliated Hospital of Wenzhou Medical University, Wenzhou 325000, China. Email: hzp120119@163.com.

Background: 5-fluorouracil (5-FU) is a common chemotherapy drug for gastric cancer. Human antigen R (HuR) is an RNA-binding protein that is also known as ELAV like RNA binding protein 1 (ELAVL1) regulates gene expression by binding to target genes 3' UTR region and is highly expressed in tumor tissues. However, the regulatory mechanisms of HuR in 5-FU mediated chemotherapy in stomach cancer are not well understood. In this study, we aimed to investigate 5-FU regulated PKC δ expression and translocation of HuR in SGC cell lines.

Methods: Using Cell viability assay to obtained IC₅₀ doses when SGC cells were treated with 5-FU for 2 and 3 days. Western blot was used to detect the total protein content of HuR and PKC δ after treating with 20 mM 5-FU. Furthermore, after adding 20 mM 5-FU, knocking down PKC δ , HuR and using inhibitor Staurosporine respectively, the protein content of HuR in the cytoplasm were detected by Western blot and Immunofluorescence. MTT assay was used to verify the changes in cell proliferation after knockdown of HuR.

Results: Using Cell viability assay, we obtained IC₅₀ doses of 77 and 20 mM, respectively, when SGC cells were treated with 5-FU for 2 and 3 days. The total protein content of HuR and PKC δ in SGC cells treated with 20 mM 5-FU did not change. However, Western blot and Immunofluorescence detected that the protein content of HuR in the cytoplasm decreased with 20 mM 5-FU treatment. Staurosporine inhibits the nucleation of HuR and is an inhibitor of PKC. Consistently, the expression of HuR in cytoplasm declined while knockdown of PKC δ . And the proliferation of SGC cells decreased after knocking down the HuR.

Conclusions: Our results showed that 5-FU blocked shuttling of HuR from the nucleus to the cytoplasm in SGC cells through PKC δ phosphorylation.

Keywords: 5-fluorouracil (5-FU); human antigen R (HuR); PKC δ ; stomach; cancer

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Introduction

Stomach cancer is the fifth most generally diagnosed cancer, and the third primary cause of cancer-related mortality globally (1,2). The mainstay treatment for stomach cancer is surgical resection. Still, perioperative strategies containing chemotherapy and radiotherapy are regarded as effective approaches to enhance patient survival (3-5). Moreover, combination therapy with 5-fluorouracil (5-FU), platinum, and trastuzumab is recognized as the preferred chemotherapy choice for patients with metastatic stomach cancer (6). Meanwhile, many genes play a crucial role in stomach cancer chemotherapy. For example, compared with standard chemotherapy alone, its combination with trastuzumab can significantly prolong the average survival time of patients with Hairy-related 2 (HER2) positive stomach cancer (7). Also, the combined expression levels of MYC proto-oncogene (MYC), Epidermal growth factor receptor (EGFR), and Fibroblast growth factor receptor 2 (FGFR2) were associated with the poor survival rate of chemotherapy-treated patients (8). Loss of MutL homolog 1 (MLH1) in stomach cancer patients is related to chemoresistance (9).

Human antigen R (HuR) belongs to the Elavl family; it binds to AU-rich RNA elements in 3'-UTR with a high affinity to control the stability and translation of target mRNA (10). HuR is ubiquitously expressed and vital for the modulation of numerous mRNAs involved in cellular proliferation, differentiation, migration inflammation, and tumorigenesis (11,12). Increasing evidence indicated that the upregulated expression level of HuR had been found in a variety of human cancers, including stomach cancer (13). Overexpression of HuR improves the stabilization of mRNA, coding proteins involving TNF α , cyclins, matrix metalloproteinase 9 (MMP-9), and cyclooxygenase 2 (COX-2) (14). Meanwhile, the inhibition of HuR expression can suppress the progression of cancer, such as MS-444, miR-22, and colon carcinoma-1 (OCC-1) (15,16). The cancer cells' characteristics are regulated by HuR in other words (17). In endothelial-specific Elavl1 knock-out mice, oncogene-induced mammary cancer showed decreased revascularization, and Lewis lung carcinoma tumor implant exhibited significantly reduced tumor volume and weight (18). Moreover, Apcmin/p mice, a transgenic animal model of intestinal tumorigenesis, had attenuated polyposis after Intestinal HuR deletion (19). Knocked down of HuR expression caused by small interfering RNA suppressed cell proliferation and migration in the human prostate cancer cell line (20).

HuR is predominantly located in the cell nucleus; its translocation to the cytoplasm plays a vital role in the protective effects on related mRNAs (21-23). The nucleo-cytoplasmic HuR shuttling, in response to various environmental stimuli, is regulated by a major cellular signaling pathway (24). Previous studies have shown that HuR shuttling can be activated by the cell-cycle checkpoint kinase 2 (Chk2), PKC α , and PKC δ through direct phosphorylation (25-27). HuR plays a key role during zebrafish primitive embryonic erythropoiesis, for instance, mediated by PKC (23,28). Since the cytoplasmic abundance of HuR is increased in various types of cancers, it is considered as a likely indicator for the therapeutic effect and survival prognosis of particular cancerous patients (25).

5-FU acts as an anti-metabolite, metabolizing to FdUMP intracellularly, interfering with DNA synthesis and repairment and causing cell damage and death, particularly in tumor cells (29). Therefore it is used to treat various cancers, including gastrointestinal cancer, breast cancer, and pancreatic cancer (2,30). 5-FU improved the HuR cytoplasmic expression in pancreatic ductal adenocarcinoma (PDA) cells and induced the accumulation of HuR in breast cancer cells. However, the underlying molecular mechanism concerning HuR and 5-FU is still unclear in stomach cancer cells. Interestingly, we showed that 5-FU significantly downregulated the protein level of cytoplasmic HuR with 5-FU treatment, without any change in the protein level of total HuR. Also, PKC inhibitor and PKC δ siRNA suppressed nucleo-cytoplasmic HuR shuttling in SGC cells. And decreased expression of HuR resulted in decreased proliferation of SGC cells.

Methods

Cell culture

The SGC cell line (RRID:CVCL_0520) was purchased from Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences, Shanghai, China and maintained in RPMI-1640 medium (Invitrogen, Carlsbad, CA, USA), supplemented with 10% fetal bovine serum, L-glutamine, and penicillin/streptomycin (Invitrogen). Cells were incubated under a humidified atmosphere containing 5% CO₂ at 37 °C.

Cell transfections

The knockdown of PKC δ and HuR were achieved using

small interfering RNAs (siRNA, GenePharma). Transfection of SGC cells was performed using Lipofectamine™ RNAiMAX Transfection Reagent (Invitrogen).

PKC δ : sense (5'-3') GGAGAGUACUUUGCCAUCATT
antisense (5'-3') UGAUGGCAAAGUACUCUCCTT

HuR: sense (5'-3') UACCAGUUUCAUUGGUCAUAA
antisense (5'-3') UUAUGACCAUUGAAACUGGUA.

Drug treatment

5-FU (MedChemExpress, Cat. # HY-9006/CS-0993) and PKC inhibitors, Staurosporine (MedChemExpress, Cat. # HY-15141/CS-2716) were dissolved in Dimethyl sulfoxide (DMSO), then diluted into PBS for the following study. PKC inhibitors, Staurosporine were added to SGC cells 5 hours in addition to 5-FU (20 μ M). Cells were collected at 24, 48, or 72 hours.

Cell viability assay

Cell viability was assessed by MTT cell proliferation and cytotoxicity assay kit (Beyotime Biotechnology, Cat. # C0009). SGC cells were seeded in 96-well plates at a density of 5,000 cells/well. Thiazolyl blue tetrazolium bromide (MTT) solution was prepared following the instructions and incubated with the growing cultures at a final quantity of 10 μ L for 4 hours at 37 °C. Then add 100 microliters of Formazan solution to each well and incubated at 37 °C for 4 hours. Absorbance at 570 nm was measured on a multi-well scanning spectrophotometer (UV-2100; UNICO, Shanghai China), and the results were expressed as a percentage (%) of the control.

Protein extraction and Western blot analysis

Nuclear and cytoplasmic proteins were separated, quantified, and subjected to Western blot analysis as described (28). SGC cells were lysed with RIPA buffer for at least 30 minutes on ice. Protein concentrations were determined by a BCA protein assay kit (Beyotime, Cat. # P0010, China). Total protein was electrophoresed in SDS-PAGE gel, transferred to a nitrocellulose membrane (Pall Corporation), and blocked with 5% skim milk for 1 hour. Then, an incubation overnight at 4 °C was performed with anti-ELAVL1 antibody (Santa Cruz Biotechnology Cat# sc-5261, RRID:AB_627770), anti-PKC δ antibody (Novus Cat# NB100-82142, RRID:AB_1146612), anti-vinculin

antibody (Sigma-Aldrich Cat# V9131, RRID:AB_477629). And anti-vinculin antibody (Sigma-Aldrich Cat# V9131, RRID:AB_477629) was used as internal reference. HRP-conjugated goat anti-mouse (AnaSpec; EGT Group Cat# 28173, RRID:AB_317949) or HRP-conjugated goat anti-rabbit (AnaSpec; EGT Group Cat# 28177, RRID:AB_317953) secondary antibody was added for 1 hour at room temperature, and blots were developed by Super Digital-ECL (Kindle Biosciences).

Immunofluorescence

The SGC cells were fixed with 4% paraformaldehyde, adhered to charged glass microscope slides, permeated with 0.1% Triton X-100 for 10 minutes, and blocked with 1% BSA for 1 hour at room temperature. The cells were incubated with anti-ELAVL1 antibody (Santa Cruz Biotechnology Cat# sc-5261, RRID:AB_627770) at 4 °C overnight. Secondary antibody Goat anti-mouse IgG (H+L) AF488 (SouthernBiotech Cat# 1021-30, RRID:AB_2794251) was added for 1 hour at room temperature. DAPI (Thermo Fisher Scientific Cat# D1306, RRID:AB_2629482) was added for 15 minutes to stain nuclei. Images were obtained by EVOS FL Auto Cell Imaging System (Thermo Fisher Scientific).

Statistical analysis

Statistical analysis was performed by one-way ANOVA followed by Tukey's post-hoc test with Prism (RRID:rid_000081; GraphPad Software, La Jolla, CA, USA). Data are expressed as mean \pm SD. Significant differences were set at $P < 0.05$.

Results

5-FU inhibits the proliferation of human SGC cells

5-FU is a common tumor chemotherapy drug in clinical practice. To evaluate the inhibitory effect of 5-FU on the proliferation of SGC cells, we tested the viability of SGC cells at different concentrations of 5-FU at 2 and 3 days by Cell viability assay. Inconsistent with the expected results, the viability of SGC cells gradually decreased with an increased concentration of 5-FU. The IC₅₀ concentrations of SGC in these two periods were 77 and 20 mM, respectively (Figure 1A,B).

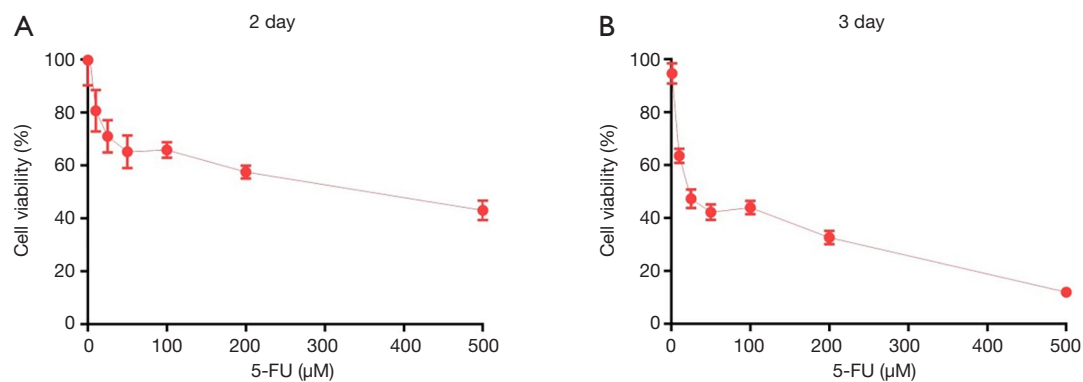


Figure 1 Measurement of IC₅₀ in SGC cells. (A) The viability of SGC cells under a series of 5-FU concentrations treatment at two days, the IC₅₀ concentrations was 77 μM. (B) The viability of SGC cells with different doses of 5-FU treatment at three days, the IC₅₀ concentrations was 20 μM. Data are presented as mean ± SEM. 5-FU, 5-fluorouracil.

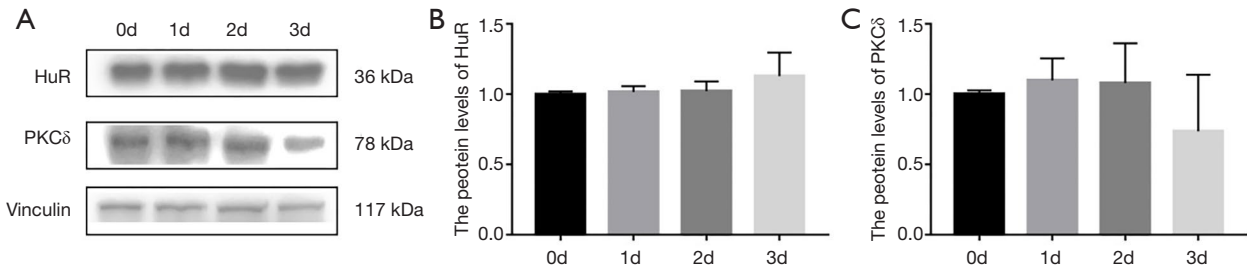


Figure 2 Protein expression levels of HuR and PKCδ after 5-FU treatment. (A) The expression levels of HuR and PKCδ protein. (B) Quantitation of HuR protein level in SGC cells with three days of 5-FU treatment. (C) Quantitation of PKCδ protein level in SGC cells with three days of 5-FU treatment. Statistical analysis was performed with one-way ANOVA. Data are presented as mean ± SEM. HuR, human antigen R; 5-FU, 5-fluorouracil.

5-FU does not affect HuR and PKCδ total protein expression

Previous studies have reported that 5-FU induced accumulation of HuR and PKC can regulate HuR expression by phosphorylating HuR during embryogenesis in zebrafish (2). Indeed, when SGC cells were treated with ~IC₅₀ doses of 5-FU (20 mM), there was no notable change in total protein expression levels of HuR and PKCδ (Figure 2A,B,C).

5-FU blocks shuttling of HuR from nuclear to cytoplasm

Previous studies showed that 5-FU and other chemotherapy agents induced HuR translocation from the nucleus to the cytoplasm in PDA cells, which is a necessary step for HuR function in the stabilization and translational control

of target mRNAs (31). In contrast to the reported results, when we treated SGC cells with IC₅₀ 5-FU (20 mM), western blot results exhibited that the amount of HuR transferred from the nucleus to the cytoplasm gradually declined with one day, two days, and three days treatment (Figure 3A,B). Immunofluorescence of HuR was similar to the results of cytonuclear fraction, and the number of HuR nucleated cells was significantly reduced compared with the control group after treatment with 20 μM 5-FU for three days (Figure 3C,D). These results show that 5-FU can inhibit the nucleation of HuR, which may be caused by different tumor cell properties.

PKCδ inhibition suppressed HuR nucleation

Our previous studies showed that blockade of PKCs suppressed HuR translocation from the nucleus to

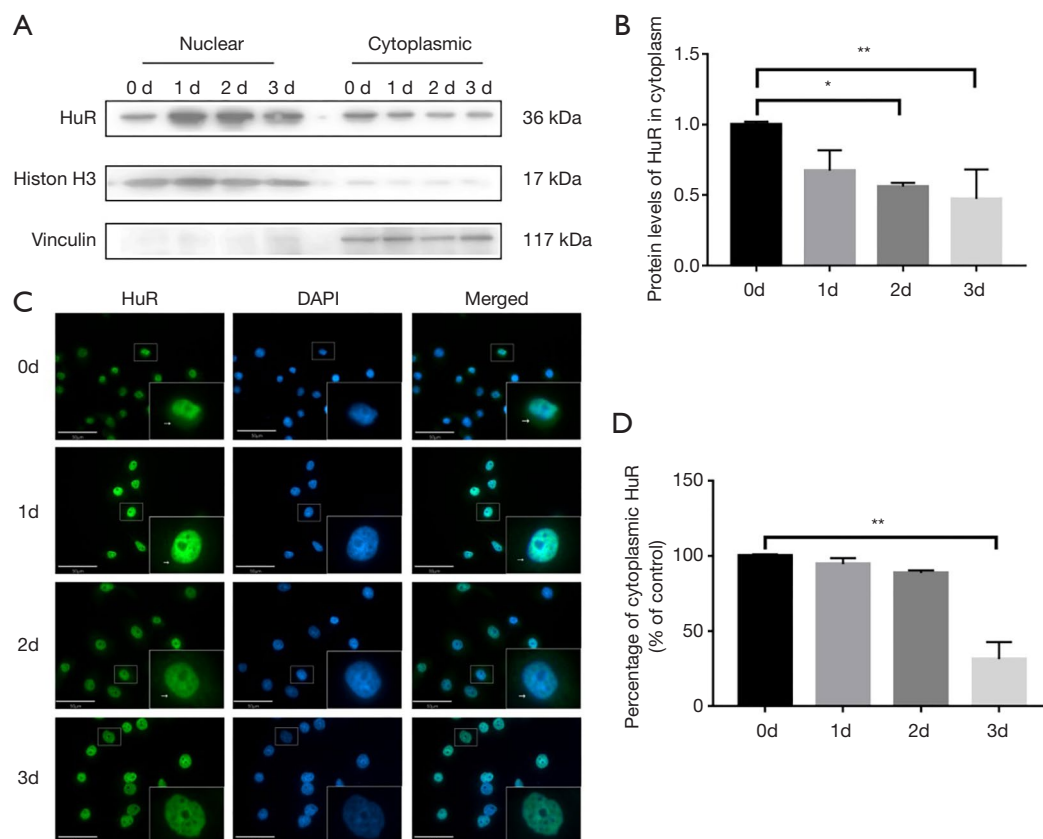


Figure 3 5-FU inhibits HuR shuffling from nuclear to the cytoplasm. (A,B) Western blot analysis of nuclear and cytoplasmic extraction of SGC cells within three days of stimulation with 5-FU of 20 μ M. Vinculin as the cytoplasmic protein control and Histone H3 as the nuclear protein control. (C,D) Immunofluorescence staining of HuR in SGC cells within three days of stimulation with 20 μ M 5-FU. Quantitative analysis of HuR protein in cytoplasmic space. Statistical analysis was performed with one-way ANOVA. Data are presented as mean \pm SEM. *, $P < 0.05$; **, $P < 0.001$. 5-FU, 5-fluorouracil; HuR, human antigen R.

cytoplasm in k562 cells (28). Therefore, we suspected that the PKC δ expression decreased during the 5-FU treatment, which led to the HuR nucleus decreased. The nuclear-cytoplasmic shuttle of HuR is an essential step for function, which is commonly regulated. For example, PKC δ can phosphorylate Ser221 and Ser318 sites of HuR to control its nucleation. To further study the effect of PKC δ on the nucleocytoplasmic shuttle of HuR, we examined the localization of HuR by PKCs inhibitors treatment in SGC cells. Western blot showed that although there is no statistical difference, as the increase of staurosporine concentration, the amount of HuR protein in the cytoplasm gradually decreased (Figure 4A). However, Immunofluorescence showed that HuR nucleated cells were significantly reduced after three days of staurosporine

stimulation at 200 nM (Figure 4B). Also, knockdown of PKC δ induced a significant reduction of HuR shuttling from nuclear to the cytoplasm (Figure 4C,D). In addition, knocking down HuR significantly reduced cell proliferation (Figure 4E,F). These results further showed that the nucleo-cytoplasmic shuttling of HuR regulated by PKC δ is necessary for SGC cells.

Discussion

In this study, 5-FU blocked the translocation of HuR from the nucleus to the cytoplasm in a time-dependent manner. Still, no significant changes in total HuR were observed after 5-FU treatment. Chemical inhibition of PKC δ and PKC δ siRNA further suppressed HuR translocation

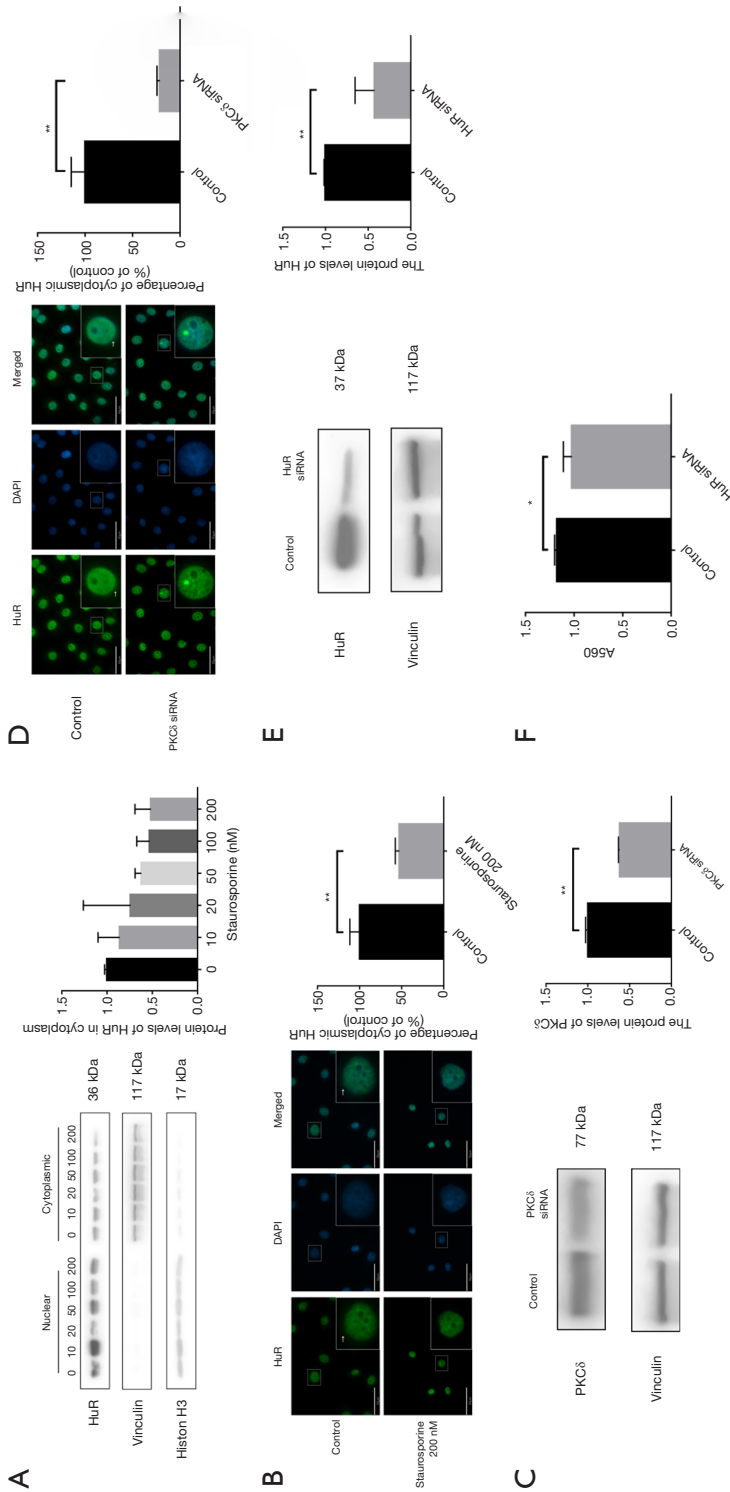


Figure 4 Blockade of PKCδ suppressed HuR shuffling. (A) Detection of HuR in nuclear and cytoplasmic extraction in SGC cells with a series dosages of PKC inhibitor Staurosporine for three days. Vinculin was used as the cytoplasmic protein control and Histone H3 as nuclear protein control. (B) Immunofluorescence staining of HuR in SGC cells after three days of stimulation with Staurosporine of 200 nM. Nucleus was stained with DAPI. Quantitative analysis of HuR protein in the cytoplasm. (C) Protein levels of PKCδ in SGC cells after transfection PKCδ siRNA. (D) Immunofluorescence staining of HuR in SGC cells after the transfection of PKCδ siRNA. Nucleus was stained with DAPI. Quantitative analysis of HuR protein in the cytoplasm was performed using ImageJ. (E) Protein levels of HuR in SGC cells after transfection HuR siRNA. (F) MTT assay in SGC cells after transfection HuR siRNA. Statistical analysis was performed with one-way ANOVA. Data are presented as mean ± SEM. *, P<0.05; **, P<0.001. HuR, human antigen R.

from the nucleus in SGC cells. These results show that the blockade of HuR shuttling was mediated by PKC δ in SGC cells. And decreased expression of HuR resulted in decreased proliferation of SGC cells. Thus, the anti-cancer effect of 5-FU was due to the reduction of PKC mediated cytoplasmic translocation of HuR.

Previous studies have reported that HUR protein levels are increased in a variety of human cancers (32). HuR was expressed at higher levels in gastric adenocarcinoma tissue, which positively associated with enhanced cytoplasmic translocation of HuR (33). Cytoplasmic HuR expression was associated with increased cytoplasmic advanced stage grade, intestinal type, non-resectability, and poor patients' survival (34). High cytoplasmic expression of HuR was an independent prognostic factor of survival for patients with Gallbladder carcinoma and Urothelial cancer (35-37). Recent studies have reported that the expression level of HuR in SGC cells significantly increased compared with the human gastric epithelium cell line (GES-1), the proliferation ability, and the migratory abilities of si-HuR-transfected SGC-7901 cells were significantly decreased (38). Moreover, over-expression of the circ-HuR suppresses HuR expression, resulted in a reduction of the growth and aggressiveness of gastric cancer. Pyrvinium pamoate, a HuR nucleocytoplasmic translocation blocker (39), has been reported to reduce cancer growth for several cancer types *in vivo*. Our results also showed that the cytoplasmic HuR was strongly decreased due to 5-FU, which indicates a negative regulatory role of HuR in gastric cancer. Conversely, in a previous study, HuR expression levels were reported to be significantly enhanced upon 5-FU stimulation in breast cancer cells (2). 5-FU induces the cytoplasmic export of HuR in pancreatic cancer (30). These findings may be related to the effects of 5-FU on cancer appear to be cell type-specific.

Protein kinase C (PKCs) are involved in the regulation of proliferation, cell junctions, apoptosis, and migration (40,41). Although the role of PKCs in cancers remains controversial (42), PKC δ is a widely known important apoptosis regulator, which mainly has the function of promoting apoptosis (26). However, PKC δ also has some anti-apoptotic functions, which have been described in previous studies (43,44). HuR phosphorylation at Ser 318 by PKC δ was found a strong increase in colon carcinomas tissue (45). The knockdown of PKC δ in ovarian cancer suppresses cell proliferation induced by the miR-940 and increased cell apoptosis by miR-940 inhibitor. PKC δ inhibition weakens breast cancer's proliferative capacity

(46,47). 5-FU caused a declining trend of PKC expression in colon cancer cell lines. Our results show that PKC inhibitor Staurosporine and knockdown of PKC δ suppressed HuR translocation from the nucleus to cytoplasm in SGC cells. In a previous study from our group, PKC δ and PKC α -mediated phosphorylation of HuR plays a critical role in cell differentiation and zebrafish embryonic erythropoiesis (28). We suggest that PKC δ -mediated phosphorylation of HuR plays a role in inhibiting cell proliferation, and 5-FU inhibits this process in SGC cells.

The cytoplasmic translocation of HuR was controlled by the phosphorylation of PKC. Staurosporine decreases HuR accumulation in the cytoplasm by blocking PKC δ infractions of human mesangial cells (14). Also, inhibition of PKC α at Ser158 and Ser221 by either knockdown or inhibitors blocked nuclear ELAVL1 export to the cytoplasm mimicking the phenomenon that PKC δ blockade with inhibitors (26). Knockdown or blockade of PKCs showed a degraded tendency to embryonic erythropoiesis by suppressed cytoplasmic translocation of HuR (28). Consistent with these results, administered PKC with Staurosporine and PKC δ siRNA in SGC cells that HUR expression was significantly decreased in the cytoplasm compared with counterpart.

In conclusion, our data propose a new sight of the molecular mechanism of 5-FU for gastric cancer. 5-FU reduces the nucleo-cytoplasmic shuttling of HuR was mediated by PKC δ , leading to a decrease in HuR in the cytoplasm, which may further promote apoptosis and inhibition of cell proliferation to play an anti-cancer effect. Our research advances to understand the molecular mechanism of 5-FU as a chemotherapy drug in gastric cancer. It supplies clues for the development of new strategies for PKC-mediated HuR cytoplasmic shuttle as a potential therapeutic target for the treatment of gastric cancer.

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Footnote

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at <http://dx.doi.org/10.21037/tcr-20-2129>). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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References

1. Bray F, Ferlay J, Soerjomataram I, et al. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin* 2018;68:394-424.
2. Solanki S, Chakinala RC, Haq KF, et al. Inpatient burden of gastric cancer in the United States. *Ann Transl Med* 2019;7:772.
3. Li Z, Gao X, Peng X, et al. Multi-omics characterization of molecular features of gastric cancer correlated with response to neoadjuvant chemotherapy. *Sci Adv* 2020;6:eaay4211.
4. Jabo B, Selleck MJ, Morgan JW, et al. Comparison of perioperative chemotherapy with adjuvant chemoradiotherapy for resectable gastric cancer: findings from a population-based study. *J Gastrointest Oncol* 2018;9:35-45.
5. Shinde A, Novak J, Amini A, et al. The evolving role of radiation therapy for resectable and unresectable gastric cancer. *Transl Gastroenterol Hepatol* 2019;4:64.
6. Takahari D. Second-line chemotherapy for patients with advanced gastric cancer. *Gastric Cancer* 2017;20:395-406.
7. Bang YJ, Van Cutsem E, Feyereislova A, et al. Trastuzumab in combination with chemotherapy versus chemotherapy alone for treatment of HER2-positive advanced gastric or gastro-oesophageal junction cancer (ToGA): a phase 3, open-label, randomised controlled trial. *Lancet* 2010;376:687-97.
8. Kim HK, Choi IJ, Kim CG, et al. Three-gene predictor of clinical outcome for gastric cancer patients treated with chemotherapy. *Pharmacogenomics J* 2012;12:119-27.
9. Hashimoto T, Kurokawa Y, Takahashi T, et al. Predictive value of MLH1 and PD-L1 expression for prognosis and response to preoperative chemotherapy in gastric cancer. *Gastric Cancer* 2019;22:785-92.
10. Brennan CM, Steitz JA. HuR and mRNA stability. *Cell Mol Life Sci* 2001;58:266-77.
11. Cheneval D, Kastelic T, Fuerst P, et al. A review of methods to monitor the modulation of mRNA stability: a novel approach to drug discovery and therapeutic intervention. *J Biomol Screen* 2010;15:609-22.
12. Chatterji P, Rustgi AK. RNA Binding Proteins in Intestinal Epithelial Biology and Colorectal Cancer. *Trends Mol Med* 2018;24:490-506.
13. Yang F, Hu A, Li D, et al. Circ-HuR suppresses HuR expression and gastric cancer progression by inhibiting CNBP transactivation. *Mol Cancer* 2019;18:158.
14. Doller A, Akool el S, Huwiler A, et al. Posttranslational modification of the AU-rich element binding protein HuR by protein kinase Cdelta elicits angiotensin II-induced stabilization and nuclear export of cyclooxygenase 2 mRNA. *Mol Cell Biol* 2008;28:2608-25.
15. Liu Y, Chen X, Cheng R, et al. The Jun/miR-22/HuR regulatory axis contributes to tumorigenesis in colorectal cancer. *Mol Cancer* 2018;17:11.
16. Blanco FF, Preet R, Aguado A, et al. Impact of HuR inhibition by the small molecule MS-444 on colorectal cancer cell tumorigenesis. *Oncotarget* 2016;7:74043-58.
17. Abdelmohsen K, Gorospe M. Posttranscriptional regulation of cancer traits by HuR. *Wiley Interdiscip Rev RNA* 2010;1:214-29.
18. Li X, Lu YC, Dai K, et al. Elavl1a regulates zebrafish erythropoiesis via posttranscriptional control of gata1. *Blood* 2014;123:1384-92.
19. Giammanco A, Blanc V, Montenegro G, et al. Intestinal epithelial HuR modulates distinct pathways of proliferation and apoptosis and attenuates small intestinal and colonic tumor development. *Cancer Res* 2014;74:5322-35.
20. Mitsunari K, Miyata Y, Asai A, et al. Human antigen R is positively associated with malignant aggressiveness via upregulation of cell proliferation, migration, and vascular endothelial growth factors and cyclooxygenase-2 in prostate cancer. *Transl Res* 2016;175:116-28.

21. Kim HH, Yang X, Kuwano Y, et al. Modification at HuR(S242) alters HuR localization and proliferative influence. *Cell Cycle* 2008;7:3371-7.
22. Wang Y, Guo Y, Tang C, et al. Developmental Cytoplasmic-to-Nuclear Translocation of RNA-Binding Protein HuR Is Required for Adult Neurogenesis. *Cell Rep* 2019;29:3101-17 e7.
23. Chang SH, Elemento O, Zhang J, et al. ELAVL1 regulates alternative splicing of eIF4E transporter to promote postnatal angiogenesis. *Proc Natl Acad Sci U S A* 2014;111:18309-14.
24. Doller A, Pfeilschifter J, Eberhardt W. Signalling pathways regulating nucleo-cytoplasmic shuttling of the mRNA-binding protein HuR. *Cell Signal* 2008;20:2165-73.
25. Doller A, Schlepckow K, Schwalbe H, et al. Tandem phosphorylation of serines 221 and 318 by protein kinase Cdelta coordinates mRNA binding and nucleocytoplasmic shuttling of HuR. *Mol Cell Biol* 2010;30:1397-410.
26. Doller A, Huwiler A, Muller R, et al. Protein kinase C alpha-dependent phosphorylation of the mRNA-stabilizing factor HuR: implications for posttranscriptional regulation of cyclooxygenase-2. *Mol Biol Cell* 2007;18:2137-48.
27. Abdelmohsen K, Pullmann R Jr, Lal A, et al. Phosphorylation of HuR by Chk2 regulates SIRT1 expression. *Mol Cell* 2007;25:543-57.
28. Zhou X, Wang S, Zheng M, et al. Phosphorylation of ELAVL1 (Ser219/Ser316) mediated by PKC is required for erythropoiesis. *Biochim Biophys Acta Mol Cell Res* 2019;1866:214-24.
29. Wei Y, Yang P, Cao S, et al. The combination of curcumin and 5-fluorouracil in cancer therapy. *Arch Pharm Res* 2018;41:1-13.
30. McAllister F, Pineda DM, Jimbo M, et al. dCK expression correlates with 5-fluorouracil efficacy and HuR cytoplasmic expression in pancreatic cancer: a dual-institutional follow-up with the RTOG 9704 trial. *Cancer Biol Ther* 2014;15:688-98.
31. Costantino CL, Witkiewicz AK, Kuwano Y, et al. The role of HuR in gemcitabine efficacy in pancreatic cancer: HuR Up-regulates the expression of the gemcitabine metabolizing enzyme deoxycytidine kinase. *Cancer Res* 2009;69:4567-72.
32. Kotta-Loizou I, Giaginis C, Theocharis S. Clinical significance of HuR expression in human malignancy. *Med Oncol* 2014;31:161.
33. Kang MJ, Ryu BK, Lee MG, et al. NF-kappaB activates transcription of the RNA-binding factor HuR, via PI3K-AKT signaling, to promote gastric tumorigenesis. *Gastroenterology* 2008;135:2030-42, 42 e1-3.
34. Rauthe G, Vahrson H, Schlagetter M. Effect of colloidal ¹⁹⁸Au on cell cultures of human tumors (author's transl). *Strahlentherapie* 1980;156:65-8.
35. Sun DP, Lin CY, Tian YF, et al. Clinicopathological significance of HuR expression in gallbladder carcinoma: with special emphasis on the implications of its nuclear and cytoplasmic expression. *Tumour Biol* 2013;34:3059-69.
36. Miyata Y, Watanabe S, Sagara Y, et al. High expression of HuR in cytoplasm, but not nuclei, is associated with malignant aggressiveness and prognosis in bladder cancer. *PLoS One* 2013;8:e59095.
37. Liang PI, Li WM, Wang YH, et al. HuR cytoplasmic expression is associated with increased cyclin A expression and poor outcome with upper urinary tract urothelial carcinoma. *BMC Cancer* 2012;12:611.
38. Matsuki M, Nishida S, Kashiwa Y, et al. Effect of dopamine on ACTH-induced glucocorticoid secretion in rat adrenal suspended cells. *Horm Metab Res* 1985;17:429-31.
39. Guo J, Lv J, Chang S, et al. Inhibiting cytoplasmic accumulation of HuR synergizes genotoxic agents in urothelial carcinoma of the bladder. *Oncotarget* 2016;7:45249-62.
40. Jiang Z, Kong C, Zhang Z, et al. Reduction of protein kinase C alpha (PKC-alpha) promote apoptosis via down-regulation of Dicer in bladder cancer. *J Cell Mol Med* 2015;19:1085-93.
41. Cross T, Griffiths G, Deacon E, et al. PKC-delta is an apoptotic lamin kinase. *Oncogene* 2000;19:2331-7.
42. Antal CE, Hudson AM, Kang E, et al. Cancer-associated protein kinase C mutations reveal kinase's role as tumor suppressor. *Cell* 2015;160:489-502.
43. Ali AS, Ali S, El-Rayes BF, et al. Exploitation of protein kinase C: a useful target for cancer therapy. *Cancer Treat Rev* 2009;35:1-8.
44. Basu A, Pal D. Two faces of protein kinase Cdelta: the contrasting roles of PKCdelta in cell survival and cell death. *ScientificWorldJournal* 2010;10:2272-84.
45. Doller A, Winkler C, Azrilian I, et al. High-constitutive HuR phosphorylation at Ser 318 by PKC{delta} propagates tumor relevant functions in colon carcinoma cells. *Carcinogenesis* 2011;32:676-85.
46. Wang F, Wang Z, Gu X, et al. miR-940 Upregulation Suppresses Cell Proliferation and Induces Apoptosis by

Targeting PKC-delta in Ovarian Cancer OVCAR3 Cells.
Oncol Res 2017;25:107-14.

47. Berardi DE, Flumian C, Rodriguez CE, et al. PKCdelta
Inhibition Impairs Mammary Cancer Proliferative

Capacity But Selects Cancer Stem Cells, Involving
Autophagy. J Cell Biochem 2016;117:730-40.

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