ORIGINAL RESEARCH

Genetic Variant of PP2A Subunit Gene Confers an Increased Risk of Primary Liver Cancer in Chinese

Youxin Wang[1],*
Qiuyue Huang[1],*
Xinglei Huang[1]
Huiliu Zhao[2]
Bin Guan[1]
Kechen Ban[3]
Xuefeng Zhu[1]
Zhixing Ma[1]
Yanmei Tang[1]
Zhaohui Su[1]
Qingqing Nong[1]

[1]Department of Environmental Health, School of Public Health, Guangxi Medical University, Nanning, 530021, People's Republic of China; [2]Department of Clinical Laboratory, The Affiliated Tumor Hospital of Guangxi Medical University, Nanning, 530021, People's Republic of China; [3]Department of Pathology, The University of Texas M. D. Anderson Cancer Center, Houston, TX, 77030, USA

*These authors contributed equally to this work

Correspondence: Qingqing Nong
Department of Environmental Health, School of Public Health, Guangxi Medical University, Nanning, 530021, People's Republic of China
Tel +86-771-5358146
Fax +86-771-5350823
Email nnqq@gxmu.edu.cn

Background: Protein phosphatase 2A (PP2A, a serine/threonine phosphatase) is frequently inactivated in many types of cancer, including primary liver cancer (PLC). Genetic variations in PP2A subunits have been reported to be associated with the risk of many types of cancer but rarely in PLC. This study aims to assess the association between functional polymorphisms of PP2A subunit genes and the risk of PLC in Chinese.

Methods: In a case-control study with a total of 541 PLC patients and 547 controls in Guangxi province of Southern China, we genotyped six putatively functional polymorphisms (rs10421191G>A, rs11453459del>insG, rs1560092T>G, rs7840855C>T, rs1255722G>A and rs10151527A>C) of three PP2A subunit genes (PPP2R1A, PPP2R2A and PPP2R5E) using matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrometry platform.

Results: The rs11453459insG variant genotypes (ins/ins+del/ins) of PPP2R1A were found to be significantly associated with an increased risk of PLC compared with the del/del genotype (adjusted OR = 1.290, 95% CI = 1.009–1.650), and the number of insert G allele worked in a dose-dependent manner (P_{trend}= 0.007). The stratified analysis showed that the effects of rs11453459insG variant genotypes were more evident in the subgroup who drink pond-ditch water (adjusted OR = 3.051, 95% CI = 1.264–7.364) than those never drink (P = 0.041). The carriers of rs11453459 del/ins genotype had a significantly lower level of PPP2R1A mRNA expression in liver cancer tissues than those of the del/del genotype (P = 0.021). Furthermore, we used microcystin-LR, a carcinogen presents in the pond-ditch water, to treat human peripheral blood mononuclear cells and found that the cells from carriers of rs11453459insG variant genotypes induced more DNA oxidative damages than those from the del/del genotype carriers (P < 0.001).

Conclusion: These findings suggest that the PPP2R1A rs11453459del>insG polymorphism is associated with an increased risk of PLC, especially for persons with a history of drinking pond-ditch water. This insertion/deletion polymorphism may be a susceptible biomarker for PLC in Chinese.

Keywords: PP2A, primary liver cancer, insertion/deletion polymorphism, pond-ditch water

Introduction

Primary liver cancer (PLC), 80% of which is hepatocellular carcinoma (HCC), ranks the sixth most common cancer and the third leading cause of cancer-related deaths in the world, accounting for 830,180 deaths in 2020.[1,2] More than half of the global new PLC cases and deaths occurred in China in 2015.[3] Particularly, Guangxi province in Southern China has the highest incidence of PLC, and accounts for 30.7% of PLC deaths in China.[4] Chronic hepatitis B virus (HBV) infection, aflatoxin B1 exposure, and consumption of microcystins (MCs)-contaminated drinking water have been reported to contribute to the unusually high incidence

of PLC in Guangxi province.[5,6] Multiple studies have revealed the contributions of genetic variants to the effects of chronic HBV infection and aflatoxin B1 exposure on PLC risk.[7,8] However, little is known about the roles of genetic variants in the effects of MCs causing PLC.

It has been reported that the underlying carcinogenic mechanism of MCs was through inhibition of protein phosphatase 2A (PP2A).[9,10] PP2A is a serine/threonine phosphatase that involves the regulation of many cellular processes, including metabolism, cell cycle, DNA replication, growth, and apoptosis.[11,12] The diminished activity of PP2A contributes to the malignant transformation and tumor development.[13] As a confirmed tumor suppressor, PP2A participates in AKT, WNT and c-MYC signaling pathways.[14] The PP2A dimeric core enzyme is composed of a catalytic C subunit (PPP2CA or PPP2CB) and a scaffold A subunit (PPP2R1A or PPP2R1B). The core enzyme interacts with a variable regulatory B subunit to form the holoenzyme PP2A.[15] Previous studies have shown that genetic variants in PP2A subunit genes are associated with risks of several cancers,[16–22] such as lung cancer,[20] breast cancer[21] and soft tissue sarcoma.[22] The mutation W257G of PPP2R1A enhances cancer cell migration through the SRC-JNK-c-Jun pathway and the mutation of B56γ-PP2A partially lost tumor-suppressive function.[23,24] Lin et al reported that a common insertion/deletion polymorphism (−241 ins/delG rs11453459) in the promoter of PPP2R1A affected PPP2R1A transcription by disturbing the binding ability of transcription factor NF-κB.[16] However, the association between genetic variations in PP2A and the risk of PLC has been rarely substantiated.

In the present study, we performed a case-control study to test the hypothesis that the genetic variants in PP2A subunit genes (PPP2R1A, PPP2R2A and PPP2R5E) may contribute to the susceptibility of PLC in Guangxi province of Southern China.

Materials and Methods
Study Subjects
In this study, a total of 541 pathologically proved PLC patients (49.8±10.3 years old, 465 males and 76 females) were recruited between 2016 and 2018 in the Affiliated Tumor Hospital of Guangxi Medical University. During the same period, 547 cancer-free individuals (49.1±9.7 years old, 461 males and 86 females) were enrolled as a control group. Also, 30 PLC tumor tissues were collected from surgical patients at the same time. All patients

were confirmed by at least two pathologists. Patients and normal controls affected by other types of cancer were excluded from this study. After signing a written informed consent, all individuals were interviewed according to a structured questionnaire in order to collect personal information, including alcohol use, HBV infection, drinking pond-ditch water, and other factors including family history of cancer. The participants who had smoked less than 100 cigarettes in their lifetime were defined as non-smokers, otherwise as smokers.[25] Similarly, the participants who had consumed alcohol at least once a week for more than one year were defined as drinkers of alcohol and the remaining as nondrinkers of alcohol.[26] A pond/ditch was defined as a water storage place, about 5–6 m in width, 100 m in length, and 1–2 m deep, usually located nearby residential areas.[5] Those subjects who had used ponds/ditches as the source of drinking water for 20 years or more were defined as drinkers of pond-ditch water. Each participant was asked to donate 5 mL of blood for examination. This study was approved by the Medical Ethics Committee of Guangxi Medical University (GXMU-20160303-9). All subjects gave their written informed consent, and the study complied with the Declaration of Helsinki.

SNP Selection
SNPs in the promoter, exon, and 3′-untranslated regions (3′-UTR) were defined to be potentially functional. By searching the dbSNP database (http://www.ncbi.nlm.nih.gov/), we found four potentially functional SNPs in the three subunits of PP2A with the Minor Allele Frequency (MAF) more than 5% in the Chinese Han population. They were rs11453459del>insG in the promoter of PPP2R1A, rs10421191G>A in the 3′-UTR of PPP2R1A, rs7840855C>T in the promoter of PPP2R2A, and rs1255722G>A in the promoter of PPP2R5E. In addition, we also selected two intron SNPs, rs10151527A>C of PPP2R5E and rs1560092T>G of PPP2R1A, since they had been reported to be associated with cancer risk previously.[22] Taken together, we selected six SNPs of PP2A subunit genes (10421191G>A, rs1560092T>G and rs11453459del>insG of PPP2R1A; rs7840855C>T of PPP2R2A; rs1255722G>A and rs10151527A>C of PPP2R5E) in this study.

Genotype Analysis
The Sequenom MassArray platform was utilized for genotyping (Sequenom, San Diego, CA). The design of

primers for PCR and extension was accomplished with the software MassARRAY Assay Design 3.1 (Sequenom). The reaction system for a multiplex PCR included 1 μL (20 ng/μL) of DNA sample, 1.850 μL of H_2O, 0.625 μL of PCR buffer, 0.325 μL of $MgCl_2$ (25 mM), 0.100 μL of dNTP mix (25 mM) and 1.000 μL of primer mix (500 nM), 0.100 μL of HotStar Taq (5 units/μL). PCR reaction conditions were as follows: 5 min at 94°C, 45 cycles of DNA amplification (20 s at 94°C, 30 s at 56°C and 1 min at 72°C), 3 min at 72°C and cooling at 4°C. The operations of shrimp alkaline phosphatase and iPLEX Gold extension reactions were strictly in compliance with the manufacturer's instructions. During the experiment, the samples were placed on a SpectroCHIP for detection under the Sequenom matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) mass spectrometer. Next, we used Mass Array Typer 4.0 to identify the genotypes according to the calculated mass of the products after the extension reaction.

PPP2R1A mRNA Expression Analysis

We only examined the mRNA levels of PPP2R1A by quantitative real-time polymerase chain reaction (qRT-PCR), because there were no significant associations of PLC risk with SNPs of other PP2A subunit genes. Thirty PLC tumor tissues were used to detect the mRNA levels of PPP2R1A. Genotypes of these samples were all confirmed by sequencing. Total RNA was extracted using TriPure Reagent (Roche Applied Science, Vilvoorde, Belgium) and reverse transcribed to complementary DNA using cDNA synthesis kit ThermoScript™ RT-PCR System (Invitrogen, California, USA). Relative mRNA expression level of PPP2R1A and an internal reference gene GAPDH were detected on the ABI Prism 7500 sequence detection system (Applied Biosystems, California, USA) according to the $2^{-\Delta\Delta Ct}$ method. The primers used for PPP2R1A were 5′-AAC TTC GAC AGT ACT TCC GG AA-3′ (forward) and 5′-ATG ATC TCA CTC TTG ACG TTG T-3′ (reverse); and for GAPDH 5′-CAT GAG AAG TAT GAC AAC AGC CT-3′ and 5′-AGT CCT TCC ACG ATA CCA AAG T-3′. We predicted the binding of transcription factor to PPP2R1A promoter region that the SNPs reside in using UCSC (https://genome.ucsc.edu/) and JASPAR (http://jaspar.genereg.net/) databases.

Detection of DNA Oxidative Damage

To further explore the contents of DNA damage in individuals with different PPP2R1A genotypes, we determined the basal and MC-LR-induced DNA oxidative damage by 8-hydroxy-2′-deoxyguanosine (8-OHdG) analysis. MC-LR exposure sensitivities were evaluated in 50 additional control subjects. Each control subject donated 6 mL of human peripheral blood and then human peripheral blood mononuclear cells (PBMCs) were obtained by density gradient centrifugation using lymphocyte separation medium (Tianjin Haoyang Biotechnology Co., Ltd., Tianjin, China). PBMCs were cultured at a density of 1×10^6 cells/mL in RPMI 1640 medium supplemented with 20% fetal calf serum, 2 mM L-glutamine, 100 U/mL penicillin and 100 ug/mL streptomycin, and incubated at 37°C, 5% CO_2 for 24 h. Then the PBMCs of each subject were treated with 3 μg/mL MC-LR (treated group) or medium alone (untreated group) for 24 h. The content of 8-OHdG was detected by an ELISA kit (Nanjing Jiancheng Biology Engineering Institute, Jiangsu, China).[27] The exact 8-OHdG value of each subject was expressed as the 8-OHdG value of MC-LR-treated group minus the 8-OHdG value of untreated group.[28]

Water Sample Collection and Microcystin-LR Analysis

In August 2017, eighty-nine samples of different kinds of drinking water (29 samples of pond/ditch water, 30 samples of well water, 30 samples of tap water) were collected in Nanning district of Guangxi province. Briefly, 500 mL of the water samples were directly collected from 0.5 m below the surface of the water, and sub-packed in plastic tubes. The tubes were subsequently kept frozen at −20°C until analysis. The levels of microcystin-LR (MC-LR) in water were determined with direct competitive enzyme-linked immunosorbent assay kits (Beacon Analytical Systems Inc., Maine, USA), by strictly following the kit instructions.

Statistical Analysis

Chi-square (χ^2) test was used to analyze the differences of selected parameters (age, gender, ethnicity, smoking status, alcohol drinking, HBV infection, family history of cancer, and drinking pond-ditch water) between cases and controls. The Hardy-Weinberg equilibrium (HWE) was tested by a goodness-of-fit chi-square test to compare the expected genotype frequencies with observed genotype frequencies in controls. Odds ratios (ORs) and 95% confidence intervals (CI) were calculated to measure the potential association between genotypes and PLC risk.

An unconditional logistic regression model was applied to analyze crude and adjusted OR and its 95% CI without and with adjustment for age, gender, ethnicity, smoking status, alcohol drinking. The logistic model was also used for the trend test. Student's t-test and one-way analysis of variance (ANOVA) test were used to examine the difference in the levels of microcystin-LR, 8-OHdG or PPP2R1A mRNA expressions between different subgroups. All the tests were analyzed by the SPSS 22.0 software. Furthermore, the statistical power was calculated by using the PASS Software (https://www.ncss.com). The heterogeneity analysis was calculated by Stata 16.0 software. A two-sided P value <0.05 was considered statistically significant.

Results

Distribution of PP2A Subunit Genes Genotypes and Their Associations with Risk of PLC

In order to determine whether the six candidate SNPs of PP2A genes were associated with PLC, we examined the genotype distribution of these SNPs in both cases and controls. The demographics of PLC cases and controls are shown in Table 1. There were no significant differences in the distributions of gender, age, ethnicity, smoking status and alcohol drinking between cases and controls, respectively ($P > 0.05$ for all). However, the proportions of HBV infection, family history of cancer and drinking pond-ditch water were significantly different between cases and controls ($P < 0.001$ for all).

As shown in Table 2, the genotype frequencies of six SNPs among controls are all in agreement with the Hardy-Weinberg equilibrium ($P > 0.05$ for all). The chi-square test showed that the genotype distribution of rs11453459del>insG was significantly different between the cases and controls ($P = 0.011$). However, for other five SNPs, no significant association with PLC risk was found ($P > 0.05$ for all). Under the co-dominant genetic model, the logistic regression analysis showed that the rs11453459 ins/ins genotype conferred a 2.364-fold risk of PLC compared to the common del/del genotype (adjusted OR = 2.364, 95% CI = 1.302–4.295, $P = 0.005$), but the del/ins genotype failed (adjusted OR = 1.184, 95% CI = 0.916–1.53, $P = 0.196$). According to the smallest Akaike's information criterion, the effect of rs11453459del>insG best fitted the dominant model. The rs11453459insG variants (ins/ins+del/

Table 1 The Selected Characteristics Between PLC Cases and Controls Used for Association Study

Variables	Cases (n = 541) N (%)	Controls (n = 547) N (%)	P^a
Gender			
Male	465 (86.0)	461 (84.3)	0.438
Female	76 (14.0)	86 (15.7)	
Age (years)			
<50	240 (44.4)	247 (45.2)	0.793
≥50	301 (55.6)	300 (54.8)	
Ethnicity			
Han	321 (59.3)	330 (60.3)	0.738
Zhuang	220 (40.7)	217 (39.7)	
Smoking status			
Ever	217 (40.1)	213 (38.9)	0.693
Never	324 (59.9)	334 (61.1)	
Alcohol drinking			
Ever	200 (37.0)	189 (34.6)	0.406
Never	341 (63.0)	358 (65.4)	
HBV infection			
HBsAg (+)	425 (78.6)	63 (11.5)	**<0.001**
HBsAg (-)	116 (21.4)	484 (88.5)	
Family history of cancer			
Yes	153 (28.3)	37 (6.8)	**<0.001**
No	388 (71.7)	510 (93.2)	
Drinking pond-ditch water			
Ever	131 (24.2)	74 (13.5)	**<0.001**
Never	410 (75.8)	473 (86.5)	

Note: $^a P$ values for a two-sided χ^2 test.
Abbreviation: PLC, primary liver cancer.

ins) increased the PLC risk by 29% when compared to the del/del genotype (adjusted OR = 1.290, 95% CI = 1.009–1.650, $P = 0.043$). Meanwhile, the trend analysis indicated that the detrimental effect of PPP2R1A rs11453459del>insG exhibited a G allele dose-dependent manner to increase PLC risk (P_{trend}= 0.007).

Stratification Analysis of Association Between PPP2R1A rs11453459del>insG Genotypes and PLC Risk

We performed stratified analysis to evaluate the effects of other factors on associations between PPP2R1A

Table 2 Frequency Distribution of Genotypes in PP2A SNPs and Results of Logistic Regression Analysis for Their Associations with PLC Risk

SNPs	Cases (n =541) N (%)	Controls[a] (n = 547) N (%)	P[b]	Crude OR (95% CI)	Adjusted OR (95% CI)[c]
rs11453459del>insG					
del/del	319 (59.0)	355 (64.9)	0.011	1.000 (ref.)	1.000 (ref.)
del/ins	186 (34.4)	175 (32.0)		1.183 (0.916–1.528)	1.184 (0.916–1.531)
ins/ins	36 (6.6)	17 (3.1)		2.357 (1.298–4.278)	2.364 (1.302–4.295)
Trend test P value				0.008	0.007
del/ins+ins/ins	222 (41.0)	192 (35.1)		1.287 (1.007–1.644)	1.290 (1.009–1.650)
rs10421191G>A					
GG	326 (60.2)	326 (59.6)	0.885	1.000 (ref.)	1.000 (ref.)
GA	187 (34.6)	189 (34.6)		0.989 (0.768–1.275)	0.993 (0.770–1.281)
AA	28 (5.2)	32 (5.8)		0.875 (0.515–1.486)	0.878 (0.516–1.493)
Trend test P value				0.713	0.734
GA+AA	215 (39.8)	221 (40.4)		0.973 (0.763–1.240)	0.976 (0.766–1.245)
rs1560092T>G					
TT	217 (40.1)	221 (40.4)	0.430	1.000 (ref.)	1.000 (ref.)
TG	261 (48.2)	249 (45.5)		1.068 (0.827–1.378)	1.062 (0.822–1.372)
GG	63 (11.7)	77 (14.1)		0.833 (0.569–1.221)	0.835 (0.569–1.224)
Trend test P value				0.601	0.595
TG+GG	217 (59.9)	221 (59.6)		1.012 (0.794–1.290)	1.008 (0.751–1.285)
rs7840855C>T					
CC	352 (65.1)	346 (63.3)	0.510	1.000 (ref.)	1.000 (ref.)
CT	160 (29.6)	177 (32.3)		0.889 (0.685–1.153)	0.891 (0.686–1.158)
TT	29 (5.4)	24 (4.4)		1.188 (0.678–2.081)	1.191 (0.679–2.090)
Trend test P value				0.812	0.832
CT+TT	189 (34.9)	201 (36.7)		0.924 (0.721–1.184)	0.928 (0.723–1.190)
rs1255722G>A					
GG	191 (35.3)	207 (37.8)	0.582	1.000 (ref.)	1.000 (ref.)
GA	261 (48.2)	247 (45.2)		1.145 (0.881–1.489)	1.141 (0.877–1.485)
AA	89 (16.5)	93 (17.0)		1.037 (0.730–1.473)	1.033 (0.727–1.469)
Trend test P value				0.641	0.659
GA+AA	350 (64.7)	340 (62.2)		1.116 (0.876–1.428)	1.112 (0.868–1.424)
rs10151527A>C					
AA	167 (30.9)	156 (28.5)	0.198	1.000 (ref.)	1.000 (ref.)
AC	252 (46.6)	284 (51.9)		0.829 (0.629–1.093)	0.825 (0.626–1.089)
CC	122 (22.5)	107 (19.6)		1.065 (0.759–1.495)	1.063 (0.756–1.493)
Trend test P value				0.881	0.892
AC+CC	374 (69.1)	391 (71.5)		0.894 (0.689–1.159)	0.890 (0.686–1.156)

Notes: [a]The observed genotype frequencies of the six SNPs among the control subjects were all in agreement with the Hardy-Weinberg equilibrium ($P > 0.05$ for all); [b]χ^2 test for differences in distribution of genotype frequencies between cases and controls; [c]Adjusted in an unconditional logistic regression model that included age, gender, ethnicity, smoking status and alcohol drinking.
Abbreviations: SNPs, single nucleotide polymorphisms; PLC, primary liver cancer.

genotypes and PLC risk. Only the result of rs11453459del>insG was presented because the other SNPs had no significant findings. As shown in Table 3, the increased PLC risk caused by rs11453459insG variant genotypes (ins/ins+del/ins) is more pronounced among individuals who have had a history of drinking pond-ditch water (adjusted OR = 3.051, 95% CI = 1.264–7.364) than those who have not ($P_{heterogeneity}$= 0.041). In contrast, there were no significant differences in stratum-ORs among gender, age, ethnicity groups, smoking status, alcohol drinking, HBV infection and family history of cancer ($P > 0.05$ for all).

Table 3 Stratification Analysis of the PPP2R1A rs11453459del>insG Genotypes by Selected Variables in PLC Cases and Controls

Variables	Cases (n = 541)		Controls (n = 547)		Adjusted OR (95% CI)[a]	P_heterogeneity[b]
	del/del N (%)	del/ins+ins/ins N (%)	del/del N (%)	del/ins+ins/ins N (%)	del/ins+ins/ins vs del/del	
Gender						
Male	273 (50.5)	192 (35.5)	304 (55.6)	157 (28.7)	1.354 (0.937–1.957)	0.550
Female	46 (8.5)	30 (5.5)	51 (9.3)	35 (6.4)	1.003 (0.403–2.497)	
Age (years)						
<50	142 (26.3)	98 (18.1)	174 (31.8)	73 (13.3)	1.568 (0.915–2.687)	0.323
≥50	177 (32.7)	124 (22.9)	181 (33.1)	119 (21.8)	1.100 (0.700–1.729)	
Ethnicity						
Han	185 (34.2)	136 (25.1)	211 (38.6)	119 (21.8)	1.251 (0.803–1.949)	0.686
Zhuang	134 (24.8)	86 (15.9)	144 (26.3)	73 (13.3)	1.446 (0.840–2.489)	
Smoking status						
Ever	125 (23.1)	92 (17.0)	138 (25.2)	75(13.7)	1.268 (0.739–2.176)	0.698
Never	194 (35.9)	130 (24.0)	217 (39.7)	117(21.4)	1.458 (0.926–2.293)	
Alcohol drinking						
Ever	123 (22.8)	77 (14.2)	120 (21.9)	69 (12.6)	1.153 (0.652–2.042)	0.488
Never	196 (36.2)	145 (26.8)	235 (43.0)	123 (22.5)	1.488 (0.959–2.309)	
Drinking pond-ditch water						
Ever	77 (14.2)	54 (10.0)	54 (9.9)	20 (3.7)	3.051 (1.264–7.364)	0.041
Never	242 (44.7)	168 (31.1)	301 (55.0)	172 (31.4)	1.125 (0.772–1.640)	
HBV infection						
HBsAg (+)	251 (46.4)	174 (32.1)	44 (8.0)	19 (3.5)	1.552 (0.860–2.799)	0.425
HBsAg (-)	68 (12.6)	48 (8.9)	311 (56.9)	173 (31.6)	1.157 (0.755–1.773)	
Family history of cancer						
Yes	82 (15.2)	71 (13.1)	22 (4.0)	15 (2.7)	1.330 (0.467–3.786)	0.915
No	237 (43.8)	151 (27.9)	333 (60.9)	177 (32.4)	1.252 (0.870–1.801)	

Notes: [a]ORs were adjusted for age, gender, ethnicity, smoking status, alcohol drinking, HBV infection, family history of cancer and drinking pond-ditch water; [b]P_heterogeneity values were calculated to compare the difference of ORs within each stratum.
**Abbreviation**: PLC, primary liver cancer.

Association of the PPP2R1A rs11453459del>insG Genotypes and PPP2R1A Gene Expression Level

Next, the mRNA levels of PPP2R1A were determined in tumor tissues of 30 PLC patients by qRT-PCR. As shown in Figure 1, the mRNA levels of PPP2R1A are much lower in cases with rs11453459 del/ins genotypes than those with del/del genotype ($P = 0.021$), suggesting that the rs11453459del>insG participated in the regulation of PPP2R1A expression.

Effects of the PPP2R1A rs11453459del>insG Polymorphism on the MC-LR-Induced DNA Oxidative Damage in PBMCs

We then investigated whether PPP2R1A rs11453459del>insG genotypes may affect the responses of PBMCs against MC-LR-induced DNA oxidative damage. After genotyping, we collected PBMCs from 50 control subjects and then treated with MC-LR for one day. As shown in Figure 2, the average level of 8-OHdG in the ins/ins subgroup (N = 4) was
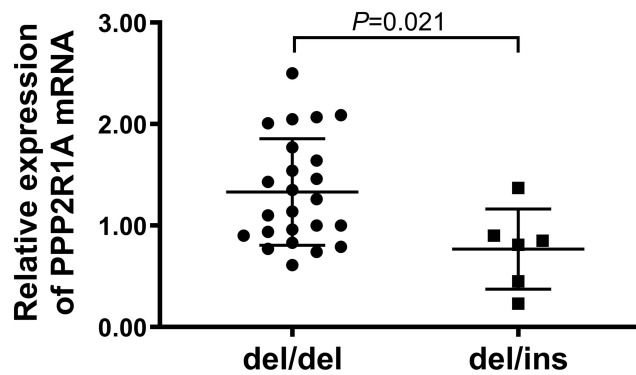
Figure 1 Relative mRNA expression levels of PPP2R1A in liver cancer tissues with different genotypes of rs11453459del>insG. The differences in the expression levels were analyzed by Student's *t*-test. The mRNA expression levels of PPP2R1A in 6 cases of liver cancer tissues harboring del/ins genotypes were significantly lower than those in 24 cases harboring del/del genotype.

significantly higher than that of del/ins subgroup (N = 14, P = 0.006) and that of del/del subgroup (N = 32, P < 0.001). When combined ins/ins and del/ins genotypes, the average level of 8-OHdG in PBMCs of del/del carriers was much lower than that of the other genotypes (P < 0.001).

Moreover, we determined the levels of MC-LR in different drinking water types collected in Guangxi province. As shown in Figure 3, the average MC-LR concentration in the pond-ditch water was 0.485±0.064 μg/L, much higher than that observed in well water (0.244±0.058 μg/L) and in tap water (0.215±0.018 μg/L). Note that the MC-LR levels in all three water types were lower than the limit of 1.0 μg/L in water supply proposed by the World Health Organization.[29]

Bioinformatics Analysis

Because the site of rs11453459 is located in the promoter of PPP2R1A, we further performed bioinformatics analysis to predict whether the rs11453459del>insG polymorphism might affect the binding ability of potent transcription factors (TF) to PPP2R1A promoter. By the use of UCSC and JASPAR databases, we found that the insertion of allele G at the site of −241 in the promoter of PPP2R1A may lose binding site of transcription factor Dp-1 (TFDP1).

Discussion

In the present case-control study (541 PLC cases and 547 controls) conducted in Guangxi province, we analyzed the associations between six common PP2A SNPs (10421191G>A, rs1560092T>G, rs11453459del>insG, rs7840855C>T, rs1255722G>A and rs10151527A>C) and PLC risk. We found that the PPP2R1A rs11453459insG variant genotypes (ins/ins+del/ins) contributed to an increased PLC risk in a G allele dose-dependent manner. The increased risk on PLC was more significant in subjects who had a history of drinking pond-ditch water. PLC patients carrying the rs11453459 del/ins genotype showed a lower PPP2R1A expression than those carrying the del/del genotype. In contrast, we did not observe significant associations between any other SNPs in PP2A and PLC risk. To the best of our knowledge, this is the first study to show the association between genetic variants of PP2A and the risk of PLC, especially in individuals with a history of drinking pond-ditch water.
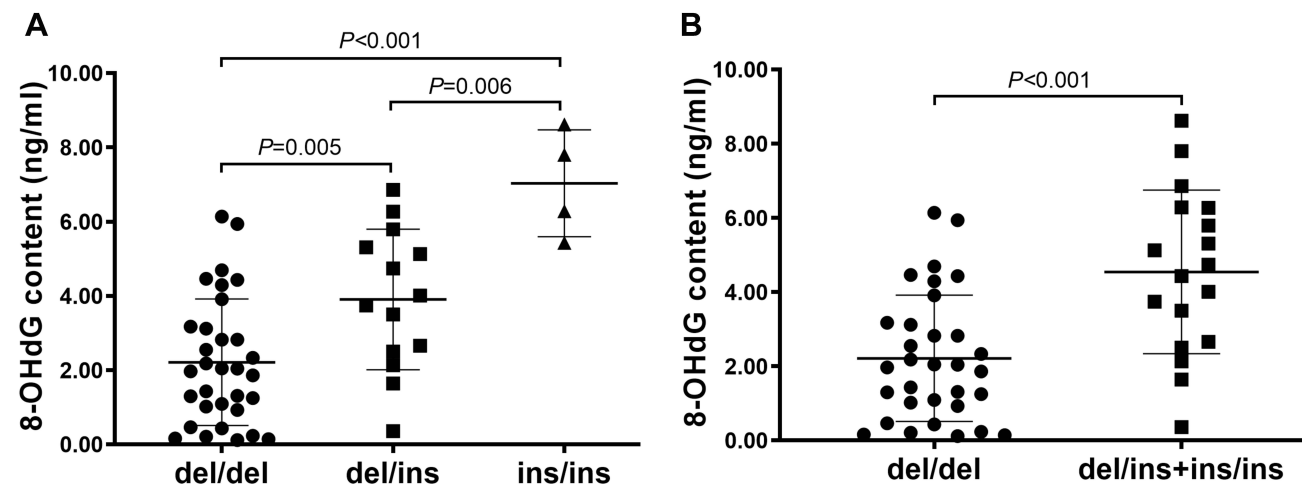
Figure 2 The PPP2R1A rs11453459del>insG polymorphism potentiated MC-LR-induced DNA oxidative damage in PBMCs. (**A**) Comparison of 8-OHdG contents by the three rs11453459del>insG genotype subgroups. (**B**) Comparison of 8-OHdG content between the del/del genotype and the other genotypes carriers (del/ins and ins/ins). The levels of 8-OHdG in subjects with different genotypes of rs11453459del>insG were analyzed with Student's *t*-test and one-way ANOVA test.
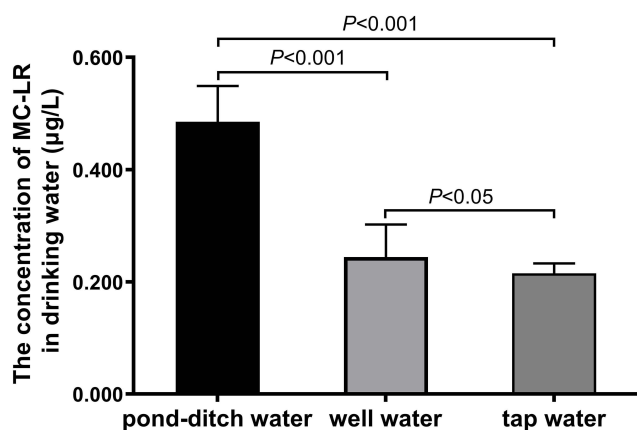
Figure 3 Microcystin-LR levels in different drinking water types collected in Guangxi province. The differences in the microcystin-LR levels were analyzed by one-way ANOVA test.

PPP2R1A serves as a scaffolding molecule to coordinate the assembly of the catalytic subunit and the regulatory B subunit to constitute PP2A. As a negative regulator on controlling cell growth and division, PPP2R1A loss can transform immortalized cells and spur tumorigenicity.[21] Several studies have reported associations between the polymorphisms of PPP2R1A and cancer risks. Dupont et al found that the SNPs in PPP2R1A were significantly associated with an increased risk of breast cancer in the Nashville population.[21] Yang et al reported that the rs11453459insG of PPP2R1A contributed to an increased risk of lung cancer in a Chinese population.[20] Kuhn et al revealed that the SNP in PPP2R1A was associated with uterine serous carcinoma.[30] These evidences highlighted the application of PPP2R1A SNPs as a possible biomarker for predicting cancer risk.

Among these reported SNPs, the PPP2R1A rs11453459del>insG is probably the most investigated. The rs11453459 site is in the promoter of PPP2R1A. It has been reported that the rs11453459del>insG was a functional genetic variant to affect the binding of NF-κB to PPP2R1A promoter and to influence PPP2R1A transcription in human liver cells.[19] Consistently in the present study, we found that the rs11453459del>insG polymorphism conferred an increased risk of PLC in a Chinese population, possibly through decreased expression of PPP2R1A mRNA in PLC tissues. The bioinformatics analysis further showed that the insertion of allele G at the site of −241 in PPP2R1A may interfere the binding of TFDP1, a transcription factor positively associated with liver cancer.[31] Therefore, it is biologically conceivable that the genetic variant of PPP2R1A is a possible biomarker for cancer susceptibility.

Particularly, we found that the effects of rs11453459insG variant genotypes on increasing PLC risk were more pronounced in individuals with ever drinking pond-ditch water. Long-term consumption of pond-ditch water is a potential risk factor for liver cancer,[5,32] and the high incidence of PLC in Guangxi province may be partially attributed to the high contamination of pond-ditch water. It had been reported that pond-ditch water was often contaminated by high concentration of blue-green algal toxin microcystin.[5,32–34] Among the identified over 100 structural analogues of MCs, MC-LR is one of the most abundant and toxic congeners.[35] MC-LR was classified by the International Agency for Research on Cancer (IARC) as a possible human carcinogen (Group 2B).[36] In our present study, much higher levels of MC-LR were observed in pond-ditch water than in tap water.

Interestingly, a previous study revealed that MCs in the pond-ditch water inhibited PP2A activity.[37] We used MC-LR to treat PBMCs and found that the cells from carriers of rs11453459insG variant genotypes were more susceptible to DNA oxidative damage than those from the del/del genotype carriers. The gene–environment interaction between the PPP2R1A rs11453459insG variants and consumption of pond/ditch water on cancer risk is biologically relevant, because the PPP2R1A rs11453459insG variants may decrease PPP2R1A transcription and consequently increase susceptibility to DNA oxidative damage from MC-LR. Further studies are warranted to test the function of the rs11453459insG variants on modulating the protumorigenic effect of MC-LR or pond/ditch water.

There are some limitations in our study. First, the clinical sample size is relatively small. Second, the present study was a hospital-based case-control study, and restricted to Chinese Han and Zhuang population, suggesting the possible presence of selection bias. Third, a considerable proportion of patients lacked clinical information such as tumor staging and differentiation status, which limited our analysis on the association between PPP2R1A variants and clinical features. Nevertheless, the genotype frequencies among controls fitted the Hardy-Weinberg disequilibrium law. It suggests the randomness of subject selection. Furthermore, the study powers were acceptable. We achieved a 90% study power (two-sided test, $\alpha = 0.05$) to detect an OR of 2.364 for the rs11453459 ins/ins and an OR of 1.290 for the rs11453459insG variant genotypes (ins/ins +del/ins), suggesting that our findings were reliable.

Conclusions

In summary, we identified that the PPP2R1A rs11453459del>insG polymorphism was associated with the risk of PLC in a Guangxi population of China. The detrimental effects of the rs11453459insG variants on increasing PLC risk were more pronounced in subjects who had a history of drinking pond-ditch water. Notably, this insertion/deletion polymorphism was functional in PLC as it was associated with the decreased expression of PPP2R1A mRNA. These data suggest that the genetic variant in PPP2R1A (rs11453459del>insG) may act as a potential susceptible biomarker for PLC in Chinese.

Institutional Review Board Statement

This study was approved by the Medical Ethics Committee of Guangxi Medical University (GXMU-20160303-9). Written informed consent was obtained from all included subjects.

Informed Consent Statement

Informed consent was obtained from all subjects involved in the study.

Author Contributions

All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; agreed to submit to the current journal; gave final approval of the version to be published; and agree to be accountable for all aspects of the work.

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Disclosure

The authors declare no conflict of interest.

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