Association between **TP53** gene Arg72Pro polymorphism and Wilms’ tumor risk in a Chinese population

Wen Fu,1,2,* Zhen-Jian Zhuo,3,* Wei Jia,1,2 Jinhong Zhu,4 Shi-Bo Zhu,1,2 Ze-Feng Lin,1,2 Feng-Hua Wang,1,2 Huimin Xia,1,2 Jing He,1,2 Guo-Chang Liu1,2

1Department of Pediatric Urology, Guangzhou Women and Children’s Medical Center, Guangzhou Medical University, Guangzhou, Guangdong; 2Department of Pediatric Surgery, Guangzhou Institute of Pediatrics, Guangzhou Women and Children’s Medical Center, Guangzhou Medical University, Guangzhou, Guangdong; 3Department of Medicine, School of Chinese Medicine, The Chinese University of Hong Kong, Hong Kong, 4Molecular Epidemiology Laboratory, Department of Laboratory Medicine, Harbin Medical University Cancer Hospital, Harbin, Heilongjiang, People’s Republic of China

*These authors contributed equally to this work

Correspondence: Guo-Chang Liu
Department of Pediatric Urology, Guangzhou Women and Children’s Medical Center, Guangzhou Medical University, 9 Jinsui Road, Guangzhou 510623, Guangdong, People’s Republic of China
Email starbless2003@126.com

Jing He
Department of Pediatric Surgery, Guangzhou Institute of Pediatrics, Guangzhou Women and Children’s Medical Center, Guangzhou Medical University, 9 Jinsui Road, Guangzhou 510623, Guangdong, People’s Republic of China
Email heijing198374@gmail.com

Abstract: Wilms’ tumor is one of the most prevalent pediatric malignancies, ranking fourth in childhood cancer worldwide. **TP53** is a critical tumor suppressor gene, which encodes a 53 kDa protein, p53. The p53 functions to protect against cancer by regulating cell cycle and apoptosis and maintaining DNA integrity. **TP53** gene is highly polymorphic. Several **TP53** gene polymorphisms have been considered to be associated with cancer risk. Of them, a nonsynonymous polymorphism, Arg72Pro (rs1042522 C>G), has been most extensively studied for the association with cancer risk; however, few studies have investigated its effect on Wilms’ tumor. Because of the central role of p53 in cell cycle control, the **TP53** gene Arg72Pro polymorphism is also a good potential candidate predisposition locus for this pediatric cancer. We genotyped this polymorphism in 145 patients and 531 cancer-free controls recruited from Chinese children by Taqman methodology. Overall, our result suggested a lack of association between the **TP53** gene Arg72Pro polymorphism and Wilms’ tumor. In the stratified analysis, we found that carriers of CG/GG genotypes had a significantly increased Wilms’ tumor risk in children not older than 18 months (adjusted odds ratio = 2.04, 95% confidence interval = 1.003–4.13, P = 0.049) compared with CC genotype carriers. Our study indicated that the **TP53** gene Arg72Pro polymorphism may have a weak, age-related effect on Wilms’ tumor risk in Chinese children. These findings need further validations in other populations with larger sample size.

**Keywords:** **TP53**, polymorphism, Wilms’ tumor, susceptibility

Introduction

Wilms’ tumor, also known as nephroblastoma, is the fourth most frequently diagnosed childhood cancer worldwide. It constitutes ~6% of all cancers detected among children younger than 15 years. The incidence rate of Wilms’ tumor varies geographically. The occurrence rate of Wilms’ tumor is ~1/8,000 children, with 400–650 new cases annually, in the USA1,2 and ~1/10,000 children in Western populations.3 Wilms’ tumor is also one of the most common renal tumors in children in China but less prevalent than that in Western countries, with an incidence rate of ~3.3 per million.4 Up to 2% of Wilms’ tumor cases are familial, with a known causal genetic lesion.5 No more than 5% of cases can be attributed to known causes, and the genetic basis underlying most Wilms’ tumors remains largely unknown.6,7

The human **TP53** gene, encoding a 53 kDa protein (p53), is located on chromosome 17p13.1. **TP53** gene is one of the most commonly mutated genes in human cancers.8 The **TP53** gene is a tumor suppressor gene, playing an important role in the maintenance of DNA integrity, cell cycle control, and apoptosis.9–11 According to dbSNP database (http://www.ncbi.nlm.nih.gov/projects/SNP), at least 1,462 single-nucleotide
polymorphisms (SNPs) have been identified in the TP53 gene so far, supporting its highly polymorphic nature. Among them, the most commonly reported SNP is a nonsynonymous Arg72Pro (rs1042522 C>G) polymorphism. This polymorphism is located in the exon 4 of the TP53 gene and can lead to the replacement of arginine (Arg) with proline (Pro) at codon 72. This polymorphism is able to alter the primary structures as well as biochemical functions of the p53 protein. It has been reported that the Arg72 form of p53 is more effective in inducing apoptosis and protecting cells from malignant transformation than the Pro72 variant. Numerous studies have been carried out to investigate the association between this polymorphism and cancer susceptibility. However, very few studies have investigated its association with Wilms’ tumor susceptibility and clinical outcomes until now. In light of the important tumor-suppressing role of TP53 gene, we investigated the association between the TP53 gene Arg72Pro polymorphism and Wilms’ tumor susceptibility in the current hospital-based case–control study with 145 cases and 531 cancer-free controls.

Subjects and methods

Subjects

This study consisted of 145 newly diagnosed and histopathologically confirmed Wilms’ tumor patients from the Department of Pediatric Urology, Guangzhou Women and Children’s Medical Center. Patients were enrolled between March 2001 and June 2016. Controls were chosen from children who visited for routine physical examination, as we described previously. A total of 531 ethnicity-, age-, and gender-matched cancer-free controls were included in the study. Written informed consent was obtained from all subjects or their legal guardians in accordance with the principles of the Declaration of Helsinki. This study was approved by the Institutional Review Board of Guangzhou Women and Children’s Medical Center.

DNA extraction and genotyping of TP53 gene Arg72Pro polymorphism

Total genomic DNA was mainly isolated from the peripheral blood leukocytes using the TIANamp Blood DNA Kit (TianGen Biotech Co., Ltd., Beijing, China) as described previously. The TP53 gene Arg72Pro polymorphism was genotyped using the TaqMan real-time PCR method. Genotyping was performed blindly to the status of the case or control. Moreover, 10% of the samples were randomly selected to perform repeated assays, and the results were 100% concordant.

Statistical analysis

χ² test was adopted to compare demographic characteristics and the frequency distributions of genotypes between cases and controls. Deviation from Hardy–Weinberg equilibrium (HWE) was tested using the χ² goodness-of-fit test for the control subjects. Odds ratios (ORs) and 95% confidence intervals (CIs) were calculated to assess the association between the TP53 gene Arg72Pro polymorphism and Wilms’ tumor susceptibility. Adjusted ORs and corresponding 95% CIs adjusting for age and gender were calculated by unconditional multiple logistic regression model. Stratified analyses were performed by age, gender, and clinical stages. All the statistical tests were two sided, and P-values <0.05 were considered as statistically significant. Statistical analysis was performed using the SAS software (Version 9.4; SAS Institute, Cary, NC, USA).

Results

Subject characteristics

Epidemiological characteristics of Wilms’ tumor patients and cancer-free controls are listed in Table S1. The mean age was 26.17 months (±21.48, range =1–132 months) for Wilms’ tumor patients and 29.73 months (±24.86, range =0.07–156 months) for controls. No significant differences were observed between the patients and controls in the distribution of age (P=0.725) and gender (P=0.956). According to the classification of childhood renal tumors defined by NWTS-5 criteria, 49% (32.76%), 49 (33.79%), 50 (34.48%), and 33 (22.76%) patients had clinical stages I, II, III, and IV Wilms’ tumor, respectively. Unfortunately, tumor specimens were not available for nine (6.21%) patients.

Association between TP53 gene Arg72Pro polymorphism and Wilms’ tumor risk

The genotype frequencies of the TP53 gene Arg72Pro polymorphism in the Wilms’ tumor patients and controls are listed in Table 1. Genotype analysis for the TP53 gene Arg72Pro polymorphism revealed that there was no significant deviation from the HWE in the control group (P=0.440). We did not observe any significant association between the TP53 gene Arg72Pro polymorphism and Wilms’ tumor susceptibility (CG vs CC: adjusted OR =1.48, 95% CI =0.95–2.32, P=0.086; GG vs CC: adjusted OR =1.02, 95% CI =0.58–1.78, P=0.948; CG/GG vs CC: adjusted OR =1.33, 95% CI =0.87–2.04, P=0.191; and GG vs CC/CG: adjusted OR =0.78, 95% CI =0.49–1.25, P=0.307).

We further explored the association between the TP53 gene Arg72Pro polymorphism and Wilms’ tumor susceptibility
Table 1 Genotype distributions of TP53 gene rs1042522 C>G polymorphism and Wilms’ tumor risk

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Cases (N=144), n (%)</th>
<th>Controls (N=530), n (%)</th>
<th>p-valuea</th>
<th>Crude OR (95% CI)</th>
<th>P-value</th>
<th>Adjusted OR (95% CI)b</th>
<th>P-valueb</th>
</tr>
</thead>
<tbody>
<tr>
<td>rs1042522 (HWE = 0.440)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CC</td>
<td>34 (23.61)</td>
<td>155 (29.25)</td>
<td>1.00</td>
<td>1.00</td>
<td></td>
<td>1.00</td>
<td></td>
</tr>
<tr>
<td>CG</td>
<td>83 (57.64)</td>
<td>255 (48.11)</td>
<td>1.48 (0.95–2.32)</td>
<td>0.083</td>
<td>1.48 (0.95–2.32)</td>
<td>0.086</td>
<td></td>
</tr>
<tr>
<td>GG</td>
<td>27 (18.75)</td>
<td>120 (22.64)</td>
<td>1.03 (0.59–1.79)</td>
<td>0.929</td>
<td>1.02 (0.58–1.78)</td>
<td>0.948</td>
<td></td>
</tr>
<tr>
<td>Additive</td>
<td></td>
<td></td>
<td>0.127</td>
<td>1.04 (0.80–1.35)</td>
<td>0.792</td>
<td>1.03 (0.79–1.34)</td>
<td>0.814</td>
</tr>
<tr>
<td>Dominant</td>
<td>110 (76.39)</td>
<td>375 (70.75)</td>
<td>1.34 (0.87–2.05)</td>
<td>0.183</td>
<td>1.33 (0.87–2.04)</td>
<td>0.191</td>
<td></td>
</tr>
<tr>
<td>Recessive</td>
<td>117 (81.25)</td>
<td>410 (77.36)</td>
<td>0.310</td>
<td>0.79 (0.50–1.26)</td>
<td>0.317</td>
<td>0.78 (0.49–1.25)</td>
<td>0.307</td>
</tr>
</tbody>
</table>

Notes: a χ² test for genotype distributions between Wilms’ tumor patients and controls. b Adjusted for age and gender.

Abbreviations: CI, confidence interval; OR, odds ratio; HWE, Hardy–Weinberg equilibrium.

Discussion

In the current hospital-based case–control study with 145 cases and 531 unrelated cancer-free controls, we did not observe any significant association between the TP53 gene Arg72Pro polymorphism and Wilms’ tumor susceptibility among Southern Chinese children. Interestingly, we found that the carriers of CG/GG genotypes, if 18 months old or younger, had a significantly increased Wilms’ tumor risk compared with counterparts carrying wild-type CC genotypes.

The TP53 gene (gene ID: 7157) is also known as P53 gene. It encodes a p53 tumor suppressor protein that contains DNA binding, transcriptional activation, and oligomerization domains.32,33 The p53 protein can regulate the expression of target genes in response to diverse cellular stresses, thereby inducing DNA repair, cell cycle arrest, metabolism changes, apoptosis, and cellular senescence accordingly.9–11,34 The p53 protein has been recognized as the guardian of human cells against cancer.9,35 However, this gene is highly mutated, and mutations in the TP53 gene have been implicated in the various types of cancer in human beings.36 Apart from the predisposition to mutations, TP53 is also highly polymorphic. A great number of SNPs has been identified in the TP53 gene. Several TP53 gene SNPs have been shown to confer cancer susceptibility, such as the rs78378222 A>C polymorphism discovered by whole-genome sequencing in 2011. We recently performed a meta-analysis to investigate the association between TP53 gene rs78378222 A>C and overall cancer risk.37 Evidence from 34 studies including 36,599 cases and 91,272 controls confirmed that this polymorphism was significantly associated with an increased overall cancer risk.37 The TP53 gene Arg72Pro polymorphism is the most extensively investigated variant among the known TP53 gene susceptibility loci, which was first reported by Matlashewski et al.38 Intriguingly, the two alleles of this nonsynonymous polymorphism differ in the capacity of inducing target gene

Table 2 Stratification analysis for the association between TP53 gene rs1042522 C>G polymorphism and Wilms’ tumor risk

<table>
<thead>
<tr>
<th>Variables</th>
<th>rs1042522 (cases/controls)</th>
<th>Crude OR (95% CI)</th>
<th>P-value</th>
<th>Adjusted OR* (95% CI)</th>
<th>P-value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, months</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤18</td>
<td>11/68</td>
<td>0.99 (0.57–1.71)</td>
<td>0.973</td>
<td>1.00 (0.58–1.73)</td>
<td>0.993</td>
</tr>
<tr>
<td>&gt;18</td>
<td>23/87</td>
<td>1.13 (0.61–2.08)</td>
<td>0.702</td>
<td>1.12 (0.61–2.07)</td>
<td>0.711</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Females</td>
<td>18/71</td>
<td>1.57 (0.86–2.87)</td>
<td>0.143</td>
<td>1.56 (0.85–2.86)</td>
<td>0.150</td>
</tr>
<tr>
<td>Males</td>
<td>16/84</td>
<td>1.74 (0.85–3.55)</td>
<td>0.130</td>
<td>1.77 (0.86–3.65)</td>
<td>0.119</td>
</tr>
<tr>
<td>Clinical stage</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>I + II</td>
<td>10/155</td>
<td>1.15 (0.68–1.93)</td>
<td>0.609</td>
<td>1.14 (0.68–1.92)</td>
<td>0.627</td>
</tr>
<tr>
<td>III + IV</td>
<td>22/155</td>
<td>1.74 (0.85–3.55)</td>
<td>0.130</td>
<td>1.77 (0.86–3.65)</td>
<td>0.119</td>
</tr>
</tbody>
</table>

Notes: * Adjusted for age and gender. The results are presented in bold if the 95% CI excluded 1 or P<0.05.

Abbreviations: CI, confidence interval; OR, odds ratio.
transcription, consequentially modifying cancer susceptibility differentially.\textsuperscript{13,39,40}

As so far, numerous studies have been conducted to investigate the association between this polymorphism and cancer susceptibility.\textsuperscript{16–20} Klug et al\textsuperscript{16} carried out a meta-analysis by including a total of 49 studies with 7,946 cases and 7,888 controls. They found that the \textit{TP53} gene Arg72Pro polymorphism was not associated with cervical cancer risk when the analyses were restricted to methodologically sound studies. Dahabreh et al\textsuperscript{19} did not find any significant association between \textit{TP53} gene Arg72Pro polymorphism and colorectal cancer with evidence from 23 studies including 6,514 cases and 9,334 controls. Xu et al\textsuperscript{18} combined 15 publications and found that the \textit{TP53} gene Arg72Pro polymorphism was associated with a significantly increased bladder cancer risk among Asians but not among Caucasians. Dahabreh et al\textsuperscript{19} carried out a meta-analysis involving breast cancer (68 studies), lung cancer (42), colorectal cancer (26), ovarian cancer (16), and endometrial cancer (8). They found that the association with the \textit{TP53} gene Arg72Pro polymorphism only existed among studies using tumor tissue as the source of genotyping material for cases (22 studies) but not among studies using other sources of genotyping material (eg, blood). However, it should be noted that the use of tumor tissue as the source of genotyping material for cases can lead to bias in the association studies. In the study by Tian et al\textsuperscript{20} with 14 investigations, including 2,506 cases and 4,386 controls, no association was observed between the \textit{TP53} gene Arg72Pro polymorphism and leukemia risk.

However, there is lack of evidence of the association between the \textit{TP53} gene Arg72Pro polymorphism and Wilms’ tumor risk to date. Only one case–control study has been performed to examine the association of this polymorphism with Wilms’ tumor risk,\textsuperscript{22} using controls from other studies. In the study with 46 cases and 300 controls, Andrade et al\textsuperscript{22} found that carriers of the rs1042522 C allele may associate with an increased Wilms’ tumor risk. In our present study, we failed to find any significant association between the \textit{TP53} gene Arg72Pro polymorphism and Wilms’ tumor risk, which was similar to most previous investigations in other types of cancer.\textsuperscript{16,17,20} Failure to find any significant association between the \textit{TP53} gene Arg72Pro polymorphism and Wilms’ tumor risk may be ascribed to the weak effect of low-penetrant SNPs, ethnicity difference, as well as the limited sample size in the current study. We found that the \textit{TP53} gene Arg72Pro polymorphism may contribute to a weak effect for Wilms’ tumor risk in Chinese children aged 18 months and younger, which may be ascribed to the fact that young children may be more genetic susceptible to Wilms’ tumor risk; besides, it may be a chance finding for the sample size, which is relatively small.

Although this is the first investigation to assess the association between \textit{TP53} gene Arg72Pro polymorphism and Wilms’ tumor risk in Chinese children, our results should be interpreted with caution for several limitations should be addressed. First, due to the low occurrence rate of Wilms’ tumor, only 145 cases were included in this study, which may have limited statistical power. Second, because of the nature of retrospective study design, we were only able to collect age, gender, ethnicity, and geographical factors for cases and controls. Other factors, such as dietary intake and parental environment exposures, were not available. Third, we just performed a case–control designed study to investigate the association between \textit{TP53} Arg72Pro polymorphism and Wilms’ tumor risk and we did not explore the potential mechanisms in cell lines for this polymorphism, which need to be investigated in future. Finally, we included only the most frequently investigated Arg72Pro polymorphism, and other \textit{TP53} gene polymorphisms (such as rs7837822 A>C) were not examined in this study.

**Conclusion**

This study indicated that the \textit{TP53} gene Arg72Pro polymorphism may contribute to a weak effect for Wilms’ tumor risk in Chinese children aged 18 months and younger. However, further prospective studies with larger sample size, different ethnicities, and more polymorphisms are required to confirm our findings.

**Acknowledgments**

This study was supported by the grant of State Clinical Key Specialty Construction Project (Pediatric Surgery) 2013 (no GJLCZD1301) and the grant of Clinical Medicine Research and Transformation Center of Brain Injury in Premature Infant in Guangzhou (no 520101-2150092). We thank Hongjiao Chen, Yanlu Tong, and Hezhen Wang for their assistance in DNA extraction and Zhang Zhao and Jin-Hua Hu for their assistance in sample collection and medical histories information collection.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**


Supplementary material

Table S1  Frequency distribution of selected variables in Wilms’ tumor patients and controls

<table>
<thead>
<tr>
<th>Variables</th>
<th>Cases (n=145)</th>
<th>Controls (n=531)</th>
<th>P-valuea</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N</td>
<td>%</td>
<td>N</td>
</tr>
<tr>
<td>Age range, months (mean ± SD)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤18</td>
<td>66</td>
<td>45.52</td>
<td>66</td>
</tr>
<tr>
<td>&gt;18</td>
<td>79</td>
<td>54.48</td>
<td>79</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Females</td>
<td>64</td>
<td>44.14</td>
<td>233</td>
</tr>
<tr>
<td>Males</td>
<td>81</td>
<td>55.86</td>
<td>298</td>
</tr>
<tr>
<td>Clinical stage</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>I</td>
<td>4</td>
<td>2.76</td>
<td></td>
</tr>
<tr>
<td>II</td>
<td>49</td>
<td>33.79</td>
<td></td>
</tr>
<tr>
<td>III</td>
<td>50</td>
<td>34.48</td>
<td></td>
</tr>
<tr>
<td>IV</td>
<td>33</td>
<td>22.76</td>
<td></td>
</tr>
<tr>
<td>NA</td>
<td>9</td>
<td>6.21</td>
<td></td>
</tr>
</tbody>
</table>

Note: *Two-sided χ2 test for distributions between Wilms’ tumor patients and controls.

Abbreviations: NA, not available; SD, standard deviation.