Review Article

Association between \textit{NFKB1} −94ins/del ATTG Promoter Polymorphism and Cancer Susceptibility: An Updated Meta-Analysis

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Nuclear factor-κB (NF-κB) was initially identified in 1986 as a transcription factor which binds to a 10 bp DNA element in kappa immunoglobulin light-chain enhancer in B cells [3]. The NF-κB family consists of p50 (NF-κB1), p52 (NF-κB2), p65 (RelA), c-Rel (Rel), and RelB. The major form of NF-κB is a heterodimer of the p50 and p65/RelA subunits which are encoded by the \textit{NFKB1} and \textit{NFKB2} genes, respectively [4]. The human \textit{NFKB1} gene is mapped to chromosome 4q24 and encodes a 50 kDa DNA-binding protein (p50) that can act as a master regulator of inflammation and cancer development [5–7].

A common insertion/deletion polymorphism (−94ins/del ATTG, rs28362491) in the promoter region of the \textit{NFKB1} gene elicits a regulatory effect on the \textit{NFKB1} gene [8]. A previous meta-analysis concluded that the deletion allele serves as a risk or protective allele for cancer susceptibility in Caucasian or Asian populations, respectively; however, it revealed no association between the polymorphism and cancer risk [9]. An increasing number of studies have assessed the association between the \textit{NFKB1} promoter −94ins/del ATTG polymorphism and cancer risk [10–12]. However, these studies obtained conflicting results. Therefore, we collected all available data to perform an updated meta-analysis.

1. Introduction

Cancer is a major public health problem worldwide; it is the primary and secondary causes of death in economically developed and developing countries, respectively [1]. The global concern on cancer continues to intensify as a result of the aging and expanding world population and the increasing adoption of cancer-causing habits. The mechanism of carcinogenesis remains largely unknown although genetic susceptibility is a known possible explanation for the interindividual variation in cancer risk [2].

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that generates a precise estimation to comprehensively and objectively investigate the association between the \textit{NFKB1} promoter $-94\text{ins/del ATTG}$ polymorphism and cancer risk.

2. Materials and Methods

2.1. Search Strategy and Identification of Relevant Studies. A comprehensive literature search for relevant articles published (last search updated in September 15, 2013) in PubMed (http://www.ncbi.nlm.nih.gov/pubmed/) was performed with the following key words: (“genetic polymorphism,” “polymorphism,” “SNP,” “single nucleotide polymorphism,” “gene mutation,” or “genetic variant”), (“neoplasm,” “cancer,” “tumor,” “carcinoma,” or “carcinogenesis”), and (“NFKB1,” “NF-$\kappa$B1,” “nuclear factor kappa B1,” “NF kappa B1,” or “nuclear factor $\kappa$B1”). The search was limited to human studies in English. All eligible studies were retrieved. The reviews and references of eligible studies were hand-searched for additional relevant publications. The most recent or complete study was selected when more than one publications contain overlapping data. A flow diagram of the study selection process is presented in Figure 1.

2.2. Inclusion Criteria. Case-control studies that evaluated the association of the \textit{NFKB1} promoter $-94\text{ins/del ATTG}$ polymorphism with cancer risk and described in detail the genotype distributions of the polymorphism in cases and controls were included in this meta-analysis.

2.3. Exclusion Criteria. Studies that were not for cancer research, were only case population, and were duplication of previous publication were excluded in this meta-analysis.
2.4. Data Extraction. Information was carefully extracted from eligible studies independently by two investigators (Xiao Yang and Pengchao Li) according to the inclusion criteria listed above, and the result was reviewed by a third investigator (Jun Tao). The following data were collected from each study: surname of first author, year of publication, ethnicity, genotyping method, source of controls, frequencies of the genotypes in cases and controls, cancer type, and ethnicity. Studies that investigated more than one type of cancer were regarded as individual datasets only in subgroup analysis. For each study, ethnicity was categorised as "Asian" or "Caucasian." Studies that investigated more than one type of cancer were regarded as individual datasets only in subgroup analyses according to cancer type. No minimum number of patients was required for this meta-analysis. Articles that reported different ethnic groups or countries or locations were considered different study samples for each category cited above.

2.5. Statistical Analysis. The strength of association between the NFKB1 promoter −94ins/del ATTG polymorphism and cancer risk was estimated through pooled odds ratio (OR) with its corresponding 95% CI. Pooled ORs were calculated for insertion allele versus deletion allele, ins/ins versus del/del, ins/del versus del/del, ins/ins + ins/del versus del/del, and ins/ins versus ins/del + del/del. Subgroup stratification analyses by ethnicity and cancer type were conducted to identify the association of the −94ins/del ATTG polymorphism with cancer susceptibility.

The between-study heterogeneity of the studies included in this meta-analysis was evaluated using the Q and I^2 statistic tests, where I^2 > 50% indicated heterogeneity [13]. The random-effects model was selected when I^2 was significant (>50%); otherwise, the fixed-effects model was selected. The allele frequencies of the NFKB1 promoter −94ins/del ATTG polymorphism from the respective study were determined by allele counting. In addition, a chi-square test was used to determine whether or not the observed frequencies of genotypes conformed to HWE. Pooled OR in the current meta-analysis was performed by weighting individual ORs by the inverse of their variance. The significance of the pooled OR was determined by the Z-test. In addition to the between-study heterogeneity assessment, we performed funnel plots and Egger’s test to evaluate the publication bias in our meta-analysis [14, 15]. All statistical analyses were performed by STATA 10.0 software (StataCorp, College Station, TX, USA).

3. Results

3.1. Eligible Studies and Meta-Analysis Databases. A total of 21 case-control studies involving 6127 cases and 9239 controls were analysed. The characteristics of all studies are presented in Table 1. The allele and genotype frequencies of the NFKB1 promoter −94ins/del ATTG polymorphism were extracted.
from all eligible studies. In total, this meta-analysis included 3 bladder cancer studies, 4 colorectal cancer studies, 2 ovarian cancer studies, 2 oral cancer studies, 2 prostate cancer studies, and 8 studies with the “other cancers.” Of the 21 studies, 14 were conducted among Asians and 7 were conducted among Caucasians. All cases were clinically pathologically confirmed.

The results of HWE test for the genotype distribution in the control population are shown in Table 1. Six of the eligible studies were not in HWE [10, 11, 18, 21, 31].

3.2. Quantitative Synthesis. The pooled ORs of the included case-control studies revealed a statistically significant association between the NFKB1 promoter –94ins/del ATTG polymorphism and cancer risk across the four genetic models ins/ins versus del/del, OR=1.47, 95%, CI=1.11—1.93; dominant model, OR=1.26, 95% CI=1.03—1.53; recessive model, OR=1.26, 95% CI=1.05—1.51; and ins allele versus del allele, OR=1.19, 95%, CI=1.05—1.35 (Table 2, Figure 2). Stratified analyses also revealed a significant association between the polymorphism and ovarian, oral, and prostate cancers in the various models. Ethnic subgroup analyses revealed significant increases in cancer risk in the four models among Asians but not among Caucasians. The results became prominent when the six studies that deviated from HWE were excluded (see Supplementary Table 1 and Supplementary Figure 1 in Supplementary Material available online at http://dx.doi.org/10.1155/2014/612972).

3.3. Evaluation of Publication Bias. Publication bias was evaluated by Begg’s funnel plot and Egger’s test, and the visual asymmetry was determined in the funnel plot analysis (Figure 3). We further evaluated the publication bias in the subgroups. The results of Egger’s tests for all genetic models are shown in Supplementary Table 2 (ins allele versus del allele, P = 0.004).

4. Discussion

NF-κB serves important functions in pathogenetic regulation and influences cancer development and aggressiveness by enhancing tumor angiogenesis, antiapoptosis, and proliferation and by repressing immune response [7, 33, 34]. Several investigators reported the constitutive activation of NF-κB in various malignancies [35, 36], including nonsmall cell lung carcinoma and colon, prostate, breast, bone, and brain cancers. p50 overexpression is frequently observed in various tumor tissues; hence, p50 is potentially involved in tumourigenesis. A polymorphism in the promoter region of NFKB1 encoding the p50 subunit of NF-κB modulates gene activity. This polymorphism has been recently reported to influence cancer risk.

A meta-analysis of all eligible studies in 2010 suggested that the deletion allele serves as a protective or risk allele for cancer susceptibility among Asians or Caucasians, respectively [9]. However, no significant association was detected for the overall population [9]. After the reported study, numerous studies further assessed the relationship between the NFKB1 promoter –94ins/del ATTG polymorphism and cancer among Asians and Caucasians [10, 12, 32]. However, the association remains inconclusive because of the inconsistent results from the published studies. Li et al. [12] found an association between del/del genotype and bladder cancer risk but none between the polymorphism and hepatocellular carcinoma susceptibility [30].

In this study, we analysed 21 eligible case-control studies with 6127 cases and 9239 controls. The results of this meta-analysis revealed a significant association between insertion allele careers and enhanced cancer risk. The probable mechanism behind the observed association may be linked to the enhanced expression and activity of p50 (NF-κB1). The insertion allele is reportedly associated with the increased promoter activity and enhanced NFKB1 mRNA expression [8, 12, 17]. This association might influence cancer development.

The major effect of p50 (NF-κB1) is mediated by its function as a component of the transcription factor NF-κB, which is among the major signalling pathways involved in the cellular response to environmental stress [7]. p50 serves an important function in inhibiting cell apoptosis by modulating the expression levels of several survival genes, such as Bcl-2 homologue A1 [37], PAI-2 [38], and IAP gene family [39]. Certain antiapoptosis proteins, such as Bcl-xl and Fas-associated death domain-like IL-1-converting enzyme inhibitor protein, are upregulated through the NFκB signalling pathway [40—42]. In addition, accumulated evidence illustrated that the p50 (NF-κB1) signalling pathways participate in cellular proliferation by increasing IL-5 [43], promoting MAPK phosphorylation [7, 44], and modulating cyclin D1 expression [45]. Therefore, the observed association between the –94ins/del ATTG polymorphism and cancer risk can be accounted for by the insertion allele that can inhibit apoptosis and promote cellular proliferation by upregulating the expression of p50 (NFKB1) [8, 12, 17], which was implicated in the abovementioned mechanism.

In the stratified analyses, the increased cancer risk remained in subgroups of Asians but not in those of Caucasians. The ethnic differences in the allele frequencies may be caused by natural selection or balance to other related genetic variants. Possible differences in genetic backgrounds and gene environment may also interact with the etiology. The increased cancer risk also remained in the subgroups of ovarian, oral, and prostate cancers. This result suggested that the NFKB1 gene might function as a prominent factor in these cancers. Therefore, further investigations are warranted to validate ethnic difference and cancer specificity in the effect of this functional polymorphism on cancer susceptibility.

This study has several limitations. First, significant between-study heterogeneity was detected in some comparisons and may be distorting the meta-analysis. Second, the genotype distribution among controls did not completely agree with HWE. However, the association between the insertion allele and cancer risk in the overall population and in the Asian population became pronounced when the six studies that deviated from HWE were excluded. Third, the studies included in the analysis used different genotyping methods with different quality control issues that may have also influenced the results. Fourth, publication bias was observed
Table 2: Meta-analysis of the NFKB1 −94ins/del ATTG promoter polymorphism and cancer risk.

| Variables                  | n$^a$ | Cases/Controls | ins/ins versus del/del OR (95% CI) | $I^2$ (%) | ins/del versus del/del OR (95% CI) | $I^2$ (%) | ins/ins + ins/del versus del/del (dominant) OR (95% CI) | $I^2$ (%) | ins/ins versus ins/del + del/del (recessive) OR (95% CI) | $I^2$ (%) | ins allele versus del allele OR (95% CI) | $I^2$ (%) |
|---------------------------|------|----------------|-------------------------------|---------|-----------------------------------|---------|-------------------------------------------------|---------|-------------------------------------------------|---------|-----------------------------------|---------|
|                           |      |                |                               |         |                                   |         |                                                 |         |                                                 |         |                                   |         |
| Total                     | 21   | 6127/9239      | 1.47 (1.11–1.93)$^b$          | 84.8    | 1.15 (0.97–1.37)$^b$              | 67.7    | 1.26 (1.03–1.53)$^b$                          | 77.5    | 1.26 (1.05–1.51)$^b$                          | 82.3    | 1.19 (1.05–1.35)$^b$                    | 84.0    |
| Cancer types              |      |                |                               |         |                                   |         |                                                 |         |                                                 |         |                                   |         |
| Bladder cancer            | 3    | 1058/1175      | 1.07 (0.45–2.53)$^b$          | 90.1    | 1.00 (0.46–2.18)$^b$              | 88.8    | 1.04 (0.46–2.33)$^b$                          | 90.7    | 1.04 (0.73–1.48)$^b$                          | 72.7    | 1.03 (0.70–1.51)$^b$                    | 89.0    |
| Colorectal cancer         | 4    | 1275/1890      | 0.84 (0.47–1.50)$^b$          | 82.9    | 0.93 (0.77–1.13)                 | 0       | 0.88 (0.73–1.06)                              | 0       | 0.89 (0.51–1.55)$^b$                          | 88.9    | 0.90 (0.72–1.12)$^b$                    | 76.2    |
| Ovarian cancer            | 2    | 366/444        | 2.57 (1.66–3.98)              | 0       | 1.88 (1.23–2.89)                 | 0       | 2.17 (1.45–3.25)                              | 0       | 1.59 (1.19–2.11)                            | 0       | 1.55 (1.27–1.90)                        | 0       |
| Oral cancer               | 2    | 674/721        | 2.10 (1.54–2.87)              | 33.0    | 1.42 (1.10–1.83)                 | 0       | 1.59 (1.25–2.03)                              | 3.9     | 1.67 (1.29–2.17)                            | 0       | 1.43 (1.23–1.66)                        | 6.9     |
| Prostate cancer           | 2    | 451/477        | 1.59 (1.09–2.33)              | 23.0    | 1.28 (0.89–1.84)                 | 28.6    | 1.40 (1.00–1.98)                              | 35.8    | 1.33 (1.01–1.74)                            | 0       | 1.26 (1.05–1.52)                        | 0       |
| Other cancers             | 8    | 2303/4532      | 1.72 (1.13–2.61)              | 80.9    | 1.16 (0.88–1.53)                 | 61.3    | 1.34 (0.99–1.83)                              | 72.4    | 1.46 (1.12–1.90)$^b$                          | 78.0    | 1.29 (1.07–1.57)$^b$                    | 79.9    |
| Ethnicities               |      |                |                               |         |                                   |         |                                                 |         |                                                 |         |                                   |         |
| Asian                     | 14   | 4143/5169      | 1.83 (1.30–2.57)$^b$          | 84.8    | 1.23 (0.97–1.58)$^b$              | 75.9    | 1.42 (1.08–1.86)$^b$                          | 82.5    | 1.50 (1.26–1.78)$^b$                          | 66.8    | 1.32 (1.14–1.54)$^b$                    | 82.2    |
| Caucasian                 | 7    | 1984/4070      | 0.90 (0.64–1.27)$^b$          | 71.2    | 1.00 (0.85–1.18)                 | 18.5    | 0.95 (0.81–1.10)                              | 24.0    | 0.90 (0.66–1.23)$^b$                          | 83.7    | 0.95 (0.82–1.12)$^b$                    | 70.9    |

$^a$Number of comparisons.

$^b$Random effects estimate.
in our study, which may affect the validity of conclusion. In the stratified analysis, we found that the publication bias was significant among the Asian groups and other cancer groups but not significant among the Caucasian, bladder, and colorectal cancer groups. The sample sizes of the included studies were diverse, and most of them were insufficiently large. These conditions might partly interpret the publication bias. Finally, only three controls were population based; thus, they may not represent the general population. Therefore, the results of this study should be interpreted with caution.

In conclusion, the NFKB1 promoter −94ins/del ATTG polymorphism is associated with cancer risk. Well-designed studies with representative sample sizes are necessary to validate these findings.

### Conflict of Interests

The authors have declared that no conflict of interests exist.

### Authors’ Contribution

Xiao Yang, Pengchao Li, and Jun Tao contributed equally to this work.

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