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Thyroid Cancer: The Quest for Genetic Susceptibility Involving DNA Repair Genes

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Abstract: The incidence of thyroid cancer (TC), particularly well-differentiated forms (DTC), has been rising and remains the highest among endocrine malignancies. Although ionizing radiation (IR) is well established on DTC aetiology, other environmental and genetic factors may also be involved. DNA repair single nucleotide polymorphisms (SNPs) could be among the former, helping in explaining the high incidence. To further clarify the role of DNA repair SNPs in DTC susceptibility, we analyzed 36 SNPs in 27 DNA repair genes in a population of 106 DTCs and corresponding controls with the aim of interpreting joint data from previously studied isolated SNPs in DNA repair genes. Significant associations with DTC susceptibility were observed for XRCC3 rs861539, XPC rs2228001, CCNH rs2230641, MSH6 rs1042821 and ERCC5 rs2227869 and for a haplotype block on chromosome 5q. From 595 SNP-SNP combinations tested and 114 showing relevance, 15 significant SNP combinations (p < 0.01) were detected on paired SNP analysis, most of which involving CCNH rs2230641 and mismatch repair variants. Overall, a gene-dosage effect between the number of risk genotypes and DTC predisposition was observed. In spite of the volume of data presented, new studies are sought to provide an interpretability of the role of SNPs in DNA repair genes and their combinations in DTC susceptibility.

Keywords: Thyroid cancer; DNA repair; genetic susceptibility; genetic markers; SNPs

1. Introduction

Thyroid cancer (TC) is the most common endocrine malignancy and its increasing incidence raises concern. It is two to four times more frequent in women than in men and one of the most common malignancies in adolescent and young adults, ages 15–39 years, the median age at diagnosis being lower than that for most other types of cancer [1,2]. Papillary (PTC) and follicular (FTC) thyroid cancer,
representing 85–90% and 5–10% of cases, respectively, are the most common histological varieties
and are often collectively referred to as well-differentiated thyroid carcinoma (DTC). In contrast to
anaplastic thyroid cancer (ATC), DTC prognosis is generally good, with high long-term survival and
low disease-specific mortality [3,4].

DTC aetiology is multifactorial, resulting from the interplay between genetic and environmental
factors: exposure to ionizing radiation (IR), particularly during childhood, remains the best-established
modifiable risk factor, despite others – such as dietary habits (e.g., iodine intake), obesity and xenobiotic
exposure – have also been proposed [2,4,5]. The importance of hereditary factors on DTC susceptibility
is evidenced from familial studies demonstrating high disease risk among first-degree relatives and
placing DTC as one of the cancers with higher heritability [6]. So far, the most robust evidence –
provided by several genome wide association studies (GWASs), with independent replication across
different populations – establishes markers at 9q22.33 (FOXE1), 14q13.3 (NKX2-1), 2q35 (DIRC3), 8p12
(NRG1) and 1q42.2 (PCNL2) as the strongest genetic susceptibility markers for DTC (reviewed in [6,7]).
Further candidate markers such as single nucleotide polymorphisms (SNPs) within genes involved in
cell cycle control and apoptosis, DNA repair, intracellular signalling and transcriptional regulation
have been proposed (reviewed in [8–10]) but many of these findings have not been properly replicated.
Overall, currently proposed DTC risk markers are still largely insufficient to explain the high heritability
of DTC [6]. It is possible that other, yet unidentified, genetic variants have a relevant impact on DTC
susceptibility and thus explain part of the missing heritability of the disease. Their identification is
therefore highly desirable.

DNA repair safeguards genomic integrity upon exposure to genotoxic agents, its absence or
impairment leading to cancer-driving mutations in oncogenes or tumour suppressor genes (reviewed
in [11,12]). A great number of DNA repair SNPs has been associated with cancer susceptibility
(reviewed in [12,13]), strongly suggesting that such variants may, if functionally significant, modulate
the individual sensitivity to genotoxic agents and, hence, contribute to cancer predisposition.

Considering the important role that IR and, possibly, other DNA damaging agents play in DTC
aetiology, DNA repair SNPs could, through interference with DNA repair capacity, contribute to
DTC susceptibility. Indeed, prior studies by our team do suggest that SNPs across different DNA
repair pathways – e.g., RAD51 and XRCC3 (HR pathway), CCNH (NER pathway) and MSH6 (MMR
pathway) – may be implicated in TC (or, more specifically, DTC) predisposition [14–18]. Such studies
add on to prior and subsequent work by other teams [8,12,19–25] that propose additional markers
and reinforce the notion that DNA repair SNPs may contribute to DTC risk. However, besides being
scarce, these studies provide only limited information on the impact of the studied SNP in specific
subpopulations, e.g., male versus female patients or early-onset versus late-onset DTC. Considering
the specificities of DTC regarding gender distribution and median age at diagnosis [1,2] such detailed
analysis could prove useful. Although gene-gene interactions could be of utmost importance in the real
context, possibly decisive, they have only seldom evaluated and, when considered [19,20,22,24,26,27],
analyses were usually limited to the combined effect of SNPs in the same gene or in genes of the
same pathway. DNA repair proteins functionally interact with each other, both within the same DNA
repair pathway and across different pathways, establishing ground for additive or even multiplicative
effects of different SNPs (irrespective of their pathway) on DNA repair activity and, hence, cancer
risk. This has been previously demonstrated for other types of cancer such as breast cancer [28–30]
and, most likely, also applies to DTC. Such hypothesis has not, to the best of our knowledge, been
investigated, justifying the usefulness of assessing the effect of combined genotypes on DTC risk.

In the present work we grouped and analysed all studies performed by our group on a Caucasian
Portuguese population [14–18]. Since the actual biological situation reflects the concerted action of
various alleles in the repair of DNA lesions that may be carcinogenic, all the data was re-analysed
in order to identify intra and inter-pathway genotype combinations and thus further characterize
the potential contribution of those DNA repair SNPs to DTC susceptibility. Such screening efforts
may allow the identification of candidate SNPs for future use as susceptibility biomarkers, hence, the development of tailored DTC prevention policies and perhaps implementation of guidelines.

2. Material and Methods

2.1. Study Subjects

Overall, 335 Caucasian Portuguese subjects were enrolled in this hospital-based case-control study: 106 histologically confirmed DTC patients were recruited in the Service of Nuclear Medicine of the Portuguese Oncology Institute, Lisbon, Portugal where they were treated according to the hospital current practice and 229 unrelated age (±2 years) and gender-matched controls (two for each DTC case, in each of the previously published studies) were recruited at the Department of Clinical Pathology of the São Francisco Xavier Hospital, West Lisbon Hospital Centre, Portugal where they were seeking healthcare for non-neoplastic pathology. None of the study participants had personal history of prior malignancy nor familial history of thyroid disease.

In order to verify eligibility criteria and to account for potential confounding factors, information on demographic characteristics (e.g., gender, age, occupation), family history of cancer, lifestyle habits (e.g., smoking, alcohol drinking) and IR exposure was collected from each study participant, on recruitment, through a pre-designed questionnaire performed by trained interviewers. Prior exposure to relevant levels of ionizing radiation (i.e., other than that from natural and standard diagnostic sources) was denied by all subjects included in the study. Former smokers were considered as non-smokers if they gave up smoking 2 years before DTC diagnosis or 2 years before their inclusion as controls. The response rate was >95% for both cases and controls.

All studies were previously approved by the local ethics boards of the involved institutions and conducted in compliance with the Helsinki Declaration. On recruitment, prior to blood withdrawal, all eligible subjects were informed about the objectives of the study. Those agreeing to participate gave their written informed consent and were enrolled in the study. The anonymity of all participants was guaranteed.

2.2. SNP Selection

The selection of SNPs for genotyping was performed according to criteria that were predefined individually for each original study [14–18]. Briefly, eligible SNPs were required to exhibit a minor allele frequency (MAF) greater than 0.05 in Caucasian populations, the remaining criteria (e.g., being located in a coding or splice region, altering the amino acid sequence, being a tagging SNP, having been previously referred to in MEDLINE) varying according to the individual study, as indicated in the original studies of individual alleles.

Overall, a total of 36 DNA repair SNPs across all DNA repair pathways were selected for genotyping and analysed. Details on the genomic location, base and amino acid exchange and MAF of selected SNPs are presented on Table 1.
Table 1. Selected SNPs and detailed information on the corresponding base and amino acid exchanges, minor allele frequency (MAF) and AB assay used for genotyping.

| Gene   | Location | db SNP Cluster ID (rs no.) | Base Change | Aminoacid Change | MAF (%) | AB Assay ID |
|--------|----------|-----------------------------|-------------|-----------------|---------|-------------|
| **Base Excision Repair (BER)** |
| XRCC1  | 19q13.31 | rs1799782                    | C → T       | Arg194Trp       | 13.1    | -e          |
| XRCC1  | 19q13.31 | rs25487                      | G → A       | Arg399Gln       | 26.6    | -e          |
| OGG1   | 3p25.3   | rs1052133                    | C → G       | Ser326Cys       | 29.9    | -e          |
| APEX1  | 1q11.2   | rs1130409                    | T → G       | Asp148Glu       | 44.0    | C__8921503_10 |
| MUTYH  | 1p34.1   | rs3219489                    | G → C       | Gln335His       | 31.9    | C_27504565_10 |
| PARP1  | 1q42.12  | rs1136410                    | T → C       | Val762Ala       | 24.4    | C_1515368_1_ |
| **Nucleotide Excision Repair (NER)** |
| CCNH   | 5q14.3   | rs2230641                    | T → C       | Val270Ala       | 13.8    | C_11685807_10 |
| CDK7   | 5q13.2   | rs2972388                    | A → G       | Asn33Asn        | 40.5    | C_1191757_10 |
| ERCC5  | 1q33.1   | rs2227869                    | G → C       | Cys529Ser       | 4.9     | C_15956775_10 |
| ERCC1  | 19q13.32 | rs3212986                    | G → T       | Asp1104His      | 37.7    | C_1891743_10 |
| RAD23B | 9q31.2   | rs1805329                    | C → T       | Ala249Val       | 16.7    | C_11493966_10 |
| ERCC6  | 10q11.23 | rs2228529                    | A → G       | Gln413Arg       | 15.6    | C_16171343_10 |
| ERCC4  | 1p13.12  | rs4253211                    | G → C       | Arg1230Pro      | 6.4     | C_25762749_10 |
| XPC    | 3p25.1   | rs2228001                    | C → T       | Ala499Val       | 24.8    | -e          |
| **Mismatch Repair (MMR)** |
| MLH1   | 3p22.2   | rs1799777                    | A → G       | Ile219Val       | 13.0    | C_1219076_20 |
| MSH3   | 5q14.1   | rs26279                      | A → G       | Thr1045Ala      | 28.0    | C_800002_1_  |
| MLH3   | 5q14.1   | rs184967                     | G → A       | Arg949Gln       | 9.8     | C_907914_10  |
| MSH4   | 1p31.1   | rs5745549                    | G → A       | Ser914Asn       | 6.4     | C_11848003_10 |
| MSH6   | 1p31.1   | rs5743235                    | G → A       | Ala97Thr        | 21.3    | C_3286081_10 |
| PMS1   | 2q32.2   | rs5742933                    | G → C       | –            | 21.9    | C_29329633_10 |
| MLH3   | 14q24.3  | rs175080                     | G → A       | Pro844Leu       | 36.4    | C_1082805_10 |
| MSH6   | 2p16.3   | rs1042821                    | C → T       | Gly39Glu        | 20.1    | C_8760558_10 |

* MAFF (%) for MMR genes: 6.4, 8.9, 9.8, 13.0, 13.4, 21.9, 28.0, 36.4, 41.0, 49.9, 50.1, 51.5, 52.9, 61.3, 64.0, 66.5, 68.9, 74.4, 75.6, 76.9, 78.1, 80.4, 81.6, 82.8, 84.1, 85.3, 86.5, 87.8, 89.0, 90.2, 91.4, 92.6, 93.8, 95.0, 96.2, 97.4, 98.6, 99.8, 100.0.

\( ^{a} \) MAF for BER genes: 13.1, 26.6, 29.9, 44.0, 31.9, 24.4.

\( ^{b} \) MAF for NER genes: 13.8, 40.5, 4.9, 37.7, 16.7, 15.6, 6.4, 24.8.

\( ^{c} \) MAF for MMR genes: 13.0, 28.0, 9.8, 6.4, 21.3, 21.9, 36.4, 20.1.
| Gene   | Location | db SNP Cluster ID (rs no.) | Base Change | Aminoacid Change     | MAF (%) a | AB Assay ID        |
|--------|----------|---------------------------|-------------|---------------------|-----------|--------------------|
| RAD51  | 15q15.1  | rs1801321                 | G → T       | – c                 | 25.7      | C__7482700_10     |
| NBN    | 8q21.3   | rs1805794                 | C → G       | Glu185Gln           | 35.7      | C__26470398_30    |
| XRCC2  | 7q36.1   | rs3218536                 | G → A       | Arg188His           | 5.3       | – e                |
| XRCC3  | 14q32.33 | rs861539                  | C → T       | Thr241Met           | 21.7      | – e                |

**Homologous Recombination (HR)**

**Non-homologous End Joining (NHEJ)**

| Gene   | Location | db SNP Cluster ID (rs no.) | Base Change | Aminoacid Change     | MAF (%) a | AB Assay ID        |
|--------|----------|---------------------------|-------------|---------------------|-----------|--------------------|
| XRCC4  | 5q14.2   | rs1805377                 | G → A       | – d                 | 37.5      | C__11685997_10    |
| LIG4   | 13q33.3  | rs1805388                 | C → T       | Thr9Ile             | 14.6      | C__11427969_20    |
| XRCC4  | 5q14.2   | rs28360135                | T → C       | Ile134Thr           | 1.4       | C__25618660_10    |
|        | 2q35     | rs1051685                 | A → G       | – b                 | 17.2      | C__8838368_1      |
| XRCC5  | 2q35     | rs1051677                 | T → C       | – b                 | 15.6      | C__8838367_1      |
|        | 2q35     | rs6941                    | C → A       | – b                 | 15.7      | C__8838374_10     |
|        | 2q35     | rs2440                    | C → T       | – b                 | 42.0      | C__3231046_10     |

a Minor Allele Frequency, according to [http://www.ncbi.nlm.nih.gov/projects/SNP/](http://www.ncbi.nlm.nih.gov/projects/SNP/). b SNP located on 3’ UTR. c SNP located on 5’ UTR. d SNP located on intron. e not applicable (genotyping performed by PCR-RFLP). SNPs, single nucleotide polymorphisms.
2.3. Practical Methodologies—Brief Description

All DNA samples were obtained after collection of peripheral venous blood samples from each participant. The DNA extraction was performed as described previously [14–18] using a commercial available kit (QIAamp® DNA mini kit; Qiagen GmbH, Hilden, Germany), according to the manufacturer’s recommendations. All samples were stored at −20 °C until further analysis.

Genotyping was carried out through either real-time polymerase chain reaction (PCR) or conventional PCR-restriction fragment length polymorphism (RFLP) techniques, as described in previous studies [14–18]. For real-time PCR—the option for the vast majority of SNPs considered in this study – genotyping was performed on an ABI 7300 Real-Time PCR system thermal cycler (Applied Biosystems; Thermo Fisher Scientific, Inc., Waltham, MA, USA), using the commercially available TaqMan® SNP Genotyping Assays (Applied Biosystems) identified in Table 1. Conventional techniques of polymerase chain reaction (PCR) and restriction fragment length polymorphism (RFLP) were employed to genotype XRCC1 rs1799782, XRCC1 rs25487 and OGG1 rs1052133 (BER pathway); XPC rs2228000 and XPC rs2228001 (NER pathway); and XRCC3 rs861539 and XRCC2 rs3218536 (HR pathway). Primer design methods and sequences, PCR conditions, PCR product sizes, restriction analysis conditions and expected digestion pattern for each genotype have been described in full detail elsewhere [14,16,17] and will therefore not be reproduced here. Irrespective of the genotyping method, all inconclusive samples were reanalysed. Also, for quality control, at least 10–15% of genotype determinations were run in duplicates through independent experiments, with 100% concordance between experiments.

2.4. Statistical Analysis

Prior to analysis, genotype distributions for each studied SNP were checked for deviation from Hardy–Weinberg equilibrium (HWE) using SNPstat platform [31], in both case and control populations. Variable transformation was applied to categorize the only continuous variable (age of diagnosis) and the Chi-square test was then used to evaluate differences in genotype frequency, smoking status, age class and gender distributions between DTC patients and controls. Whenever the construction of 2 × 2 contingency tables was possible, the two-sided Fisher’s exact test was employed instead of the Chi-square test.

Logistic regression was used to estimate the risk of DTC associated with each genotype: risk estimates were calculated under the codominant, dominant and recessive models and expressed as crude and adjusted odds ratios (OR) and corresponding 95% confidence intervals (CI). Whenever adjustment was performed, terms for gender (male/female), age class (<30, 30–49, 50–69 and ≥70 years) and smoking habits (smokers/non-smokers) were included in the model, the most common homozygous genotype, female gender, lower age group and non-smoking status being considered the reference classes for such calculations. As data on prior IR exposure was not suitable for rigorous quantitative transformation, it was not possible to include such term in the adjustment model. Risk estimates were calculated in the whole population and after stratification according to histological type of tumour (papillary or follicular TC), gender (male and female) and age (<50 and ≥50 years).

Finally, the joint effect of multiple SNPs on DTC risk was estimated from application of logistic regression analysis (1) to relevant haplotypes, (2) to individual genetic risk scores calculated from genotype variables significant on single SNP analysis and (3) to all possible 2 × 2 combinations of the DNA repair SNPs included in this study. For the purpose of risk score calculations, genotypes presenting significant results on single SNP analysis were attributed a +1 score, the risk score for each participant corresponding to the sum of such scores. Samples with one or more missing genotypes were excluded from these calculations to avoid bias due to missing data. For paired SNP analysis, the combination of the most common homozygous genotypes of each individual SNP in the control group was taken as the reference category in OR calculations. Also, paired genotypes with frequency <5% in the study population were pooled together.
This is not a conclusive final study but an exploratory one that should be regarded as ‘proof of concept’. As such, the Bonferroni adjustment was deemed as not necessary as it is too conservative. Also, the complement of the false negative rate \( \beta \) to compute the power of a test \((1 - \beta)\) was not taken into account at this stage since further studies with more patients and controls should be undertaken to change over this preliminary study into a confirmatory positive one. All statistical analyses were performed with SPSS 22.0 (IBM SPSS Statistics for Windows, version 22.0, IBM Corp, Armonk, NY, USA) except for assessment of HWE deviation, MAF calculations, haplotype estimation and linkage disequilibrium (LD) analysis which were carried out using SNPstats [31]. Results were considered significant when the corresponding two-tailed \( p \)-values were <0.05 except for paired SNP analysis where, because of the high number of SNP-SNP combinations being tested, a more stringent significance level \((p < 0.01)\) was employed. The study was approved by the Ethical Committee of Nova Medical School, Faculty of Medical Sciences with the number 05/2008 dated of January 9th, 2008. The approval was also obtained by the ethical committee of Portuguese Oncology Institute (IPO), the hospital responsible for blood samples collection with the reference GIC/357 dated of July 14th 2004.

3. Results

3.1. General Analysis

The general characteristics of the 106 DTC patients and their 229 age- and gender-matched controls included in this study are depicted in Table 2. The overall mean age of the study population was 51 years (52.1 in the patient group and 51.0 in the control group). As expected from the worldwide gender distribution for DTC [1,2], female patients greatly outnumbered male patients in the case group. Twelve (11.3%) DTC patients were categorized as smokers. Age distribution, gender and smoking habits were not significantly different between case and control populations. Concerning histological classification of tumours, 78 (73.6%) patients were diagnosed as papillary TC while 28 (26.4%) presented follicular tumours, in line with DTC histotype distributions commonly reported in the literature [4]. Three additional cases of poorly differentiated TC were also present in some of our original studies but, since this study concerns only with DTC, such cases (and the corresponding controls) were excluded from this analysis. Prior IR exposure (except for diagnostic X-rays) was denied by all cases.

| Characteristics   | Controls n (%) | Cases n (%) | \( p \)-Value c |
|-------------------|----------------|-------------|-----------------|
| **Gender**        |                |             |                 |
| Male              | 43 (18.8)      | 16 (15.1)   | 0.445           |
| Female            | 186 (81.2)     | 90 (84.9)   |                 |
| **Age a, b**      |                |             |                 |
| <30               | 14 (6.1)       | 4 (3.8)     |                 |
| 30–49             | 85 (37.1)      | 38 (35.8)   | 0.817           |
| 50–69             | 100 (43.7)     | 49 (46.2)   |                 |
| ≥70               | 30 (13.1)      | 15 (14.2)   |                 |
| **Smoking habits**|                |             |                 |
| Non-smokers       | 184 (80.3)     | 94 (88.7)   | 0.084           |
| Smokers           | 43 (18.8)      | 12 (11.3)   |                 |
| Missing           | 2 (0.9)        | 0 (0.0)     |                 |

a Age of diagnosis, for cases. b Age at the time of diagnosis of the matched case, for controls. c \( p \)-value for cases versus control group determined by two-sided Fisher’s exact test (gender, smoking habits) or \( \chi^2 \) test (age).

Abbreviations: DTC, well-differentiated thyroid cancer.

3.1. All DTC Cases

Allelic and genotypic frequencies as well as crude/adjusted ORs were calculated for all 36 DNA repair SNPs analysed in our study. Significant findings are reported in Table 3. The allelic and genotypic frequencies observed in the control group were in agreement with those expected for Caucasian populations. Also, for the majority of SNPs, genotype distributions were in Hardy-Weinberg
equilibrium (HWE, p ≥ 0.05), in both case and control populations. Significant deviations from HWE were observed for OGG1 rs1052133, MUTYH rs3219489 and CDK7 rs2972388 in the control group and for XRCC1 rs1799782, XPC rs2282000 and MSH3 rs184967 in the DTC group. Further, strong linkage disequilibrium was observed between XRCC5 rs1051677 and rs6941, but not between any other pair of SNPs. XRCC5 rs6941 was thus excluded from further analysis, the conclusions taken for XRCC5 rs1051677 being valid for XRCC5 rs6941, since they behave as tag SNPs.

Table 3. Genotype distribution in case and control populations and associated DTC risk (crude and adjusted ORs). Only SNPs presenting significant findings are shown.

| Genotype | MAF  | Genotype Frequency | p-value a | OR (95% CI) | Adjusted OR (95% CI) b |
|----------|------|--------------------|-----------|-------------|------------------------|
|          | Controls | Cases | Controls n (%) | Cases n (%) |                         |
|          | CCNH rs2230641 | | 212 (100) | 210 (100) | 1 (Reference) | 1 (Reference) |
|          | Val/Val | | 148 (69.8) | 60 (56.6) | 1.000 | 0.94 (0.60–1.48) |
|          | Val/Ala | | 56 (26.4) | 43 (40.6) | 1.000 | 0.97 (0.60–1.57) |
|          | Ala/Ala | C: 0.17 | 8 (3.8) | 3 (2.8) | 0.775 | 0.74 (0.19–2.86) |
|          | Dominant model | | 64 (30.2) | 46 (43.4) | 1.000 | 1.00 (0.60–1.61) |
|          | Recessive model | | 8 (3.8) | 3 (2.8) | 0.775 | 0.74 (0.19–2.86) |
|          | ERCC5 rs2227869 | | 212 (100) | 210 (100) | 1 (Reference) | 1 (Reference) |
|          | Cys/Cys | | 184 (86.8) | 99 (93.4) | 1 (Reference) | 1 (Reference) |
|          | Cys/Ser | C: 0.07 | 27 (12.7) | 6 (5.7) | 0.135 | 0.21 (0.12–0.37) |
|          | Ser/Ser | C: 0.04 | 1 (0.5) | 1 (0.9) | 0.86 | 0.86 | 0.47 (0.20–1.10) |
|          | Dominant model | | 28 (13.2) | 7 (6.6) | 0.088 | 0.47 (0.20–1.10) |
|          | Recessive model | | 1 (0.5) | 1 (0.9) | 1.000 | 1.00 (0.47–2.29) |
|          | XPC rs2238001 | | 212 (100) | 210 (100) | 1 (Reference) | 1 (Reference) |
|          | Lys/Lys | | 82 (38.7) | 39 (36.8) | 1 (Reference) | 1 (Reference) |
|          | Lys/Gln | | 108 (50.9) | 47 (44.3) | 0.103 | 0.95 (0.57–1.60) |
|          | Gln/Gln | C: 0.36 | 22 (10.4) | 20 (18.9) | 1.000 | 1.00 (0.54–1.89) |
|          | Dominant model | | 130 (61.5) | 67 (63.2) | 1.000 | 1.00 (0.54–1.89) |
|          | Recessive model | | 22 (10.4) | 20 (18.9) | 0.052 | 2.01 (1.04–3.87) |
|          | MSH6 rs1042821 | | 210 (100) | 210 (100) | 1 (Reference) | 1 (Reference) |
|          | Gly/Gly | | 127 (60.5) | 68 (64.2) | 1 (Reference) | 1 (Reference) |
|          | Gly/Glu | T: 0.21 | 78 (37.1) | 30 (28.3) | 0.042 | 0.28 (0.14–0.55) |
|          | Glu/Glu | T: 0.22 | 5 (2.4) | 8 (7.5) | 0.042 | 0.29 (0.14–0.61) |
|          | Dominant model | | 83 (39.5) | 38 (35.8) | 0.343 | 0.96 (0.53–1.79) |
|          | Recessive model | | 5 (2.4) | 8 (7.5) | 0.037 | 3.35 (1.07–10.50) |
|          | XRCC3 rs861539 | | 209 (100) | 208 (100) | 1 (Reference) | 1 (Reference) |
|          | Thr/Thr | | 70 (33.5) | 36 (34.0) | 1 (Reference) | 1 (Reference) |
|          | Thr/Met | T: 0.40 | 112 (53.6) | 44 (41.5) | 0.021 | 0.63 (0.29–1.38) |
|          | Met/Met | T: 0.45 | 27 (12.8) | 26 (24.5) | 1.000 | 1.00 (0.60–1.61) |
|          | Dominant model | | 139 (66.5) | 70 (66.0) | 1.000 | 1.00 (0.60–1.61) |
|          | Recessive model | | 27 (12.9) | 26 (24.5) | 0.011 | 2.19 (1.20–3.99) |

a p-value for cases versus control group determined by two-sided Fisher’s exact test (whenever 2 × 2 contingency tables are possible) or χ² test (remaining cases). b ORs were adjusted for gender (male and female), age (<30, 30–49, 50-69, ≥ 70 years) and smoking status (non-smoker and smoker). c p < 0.05. Abbreviations: DTC, well-differentiated thyroid cancer; MAF, minor allele frequency; OR, odds ratio; CI, confidence interval.

As expected, both the comparison of genotype frequency distributions between case and control populations and the logistic regression analysis (Table 3) yielded results similar to those previously reported [14–18]: significant differences on the distribution of genotypic frequencies between cases and controls were observed for CCNH rs2230641 (p = 0.037 on the codominant model and p = 0.024 on the dominant model), for MSH6 rs1042821 (p = 0.042, on the codominant model and p = 0.037 on the recessive model) and for XRCC3 rs861539 (p = 0.021 on the codominant model and p = 0.011 on the recessive model). On logistic regression analysis, after adjustment for age, gender and smoking status, DTC risk was significantly increased in CCNH rs2230641 heterozygotes (adjusted OR = 1.89, 95% CI: 1.14–3.14, p = 0.014) and also in variant allele carriers, according the dominant model (adjusted OR = 1.79, 95% CI: 1.09–2.93, p = 0.021), in MSH6 rs1042821 variant allele homozygotes (adjusted OR = 3.42, 95% CI: 1.04–11.24, p = 0.042 on the codominant model; adjusted OR = 3.84, 95% CI: 1.18–12.44, p = 0.025 on the recessive model), in XRCC3 rs861539 variant allele homozygotes (adjusted OR = 2.20, 95% CI: 1.20–4.03, p = 0.011 on the recessive model) and in XPC rs2282001 variant allele homozygotes (adjusted OR = 2.00, 95% CI: 0.20–4.03, p = 0.049). The association between XPC rs2282001 and DTC risk is a new finding emerging from this reanalysis, since the recessive model of inheritance had not been applied in the original study [17].

Abbreviations: DTC, well-differentiated thyroid cancer; MAF, minor allele frequency; OR, odds ratio; CI, confidence interval.
No additional significant differences in genotype frequency distributions nor associations with DTC risk were found, irrespective of the model assumed.

### 3.3. Stratified Analysis

Stratified analysis according to histological tumour type, gender and age may be important to identify any subgroup-specific risk association but was only partially performed in prior studies in this population. On stratification according to histological criteria (Table 4), this study confirmed prior observations [14,17,18] of increased papillary TC risk in XPC rs2228001 and XRCC3 rs861539 variant allele homozygotes (XPC rs2228001: adjusted OR = 2.31, 95% CI: 1.07–4.98, \( p = 0.033 \); XRCC3 rs861539: adjusted OR = 2.10, 95% CI: 1.07–4.11; \( p = 0.031 \), both on the recessive model), decreased papillary TC risk in ERCC5 rs2227869 heterozygotes (adjusted OR = 0.23, 95% CI: 0.07–0.81, \( p = 0.022 \), on the codominant model) or variant allele carriers (adjusted OR = 0.22, 95% CI: 0.06–0.77, \( p = 0.018 \), on the dominant model) and increased follicular TC risk in MLH3 rs175080 variant allele carriers (crude OR = 3.95, 95% CI: 1.05–14.81, \( p = 0.042 \)) and MSH6 rs1042821 variant allele homozygotes (adjusted OR = 20.98, 95% CI: 1.08–406.53, \( p = 0.044 \), on the codominant model; adjusted OR = 23.70, 95% CI: 1.25–449.32, \( p = 0.035 \), on the recessive model). Interestingly, three other significant associations were observed in this reanalysis that were not present or had not been detected in the original studies, while two previously observed associations were lost in this reanalysis: a previously undetected decreased papillary TC risk was observed in MUTYH rs3219489 heterozygotes (crude OR = 0.56, 95% CI: 0.32–1.00, \( p = 0.048 \)) and variant allele carriers (crude OR = 0.57, 95% CI: 0.33–0.99, \( p = 0.048 \)) as well as in NBN rs1805794 variant allele homozygotes (adjusted OR = 0.28, 95% CI: 0.08–0.97, \( p = 0.045 \), on the recessive model) while the presence of the variant allele of XRCC2 rs3218536 exhibited a protective effect for follicular TC (crude OR = 0.21, 95% CI: 0.04–1.00, \( p = 0.049 \), either for heterozygotes in the codominant model and for variant allele carriers in the dominant model). In contrast, the associations of XRCC5 rs2440 and CCNH rs2230641 genotypes with papillary and follicular TC risk, respectively, reported in our original studies [15,17], were no longer observed.

On gender stratification (Table 4), when considering female patients only, a significantly increased DTC risk was evident for CCNH rs2230641 heterozygotes (adjusted OR = 1.97, 95% CI: 1.13–3.43, \( p = 0.017 \)) and variant allele carriers (adjusted OR = 1.90, 95% CI: 1.11–3.24, \( p = 0.020 \)), for XPC rs2228001 variant allele carriers (adjusted OR = 2.00, 95% CI: 1.01–3.96, \( p = 0.048 \), on the recessive model), for MSH6 rs1042821 variant allele homozygotes (adjusted OR = 4.78, 95% CI: 1.17–19.56, \( p = 0.030 \), on the codominant model; adjusted OR = 5.42, 95% CI: 1.34–21.92, \( p = 0.018 \), on the recessive model) and for XRCC3 rs861539 variant allele homozygotes (adjusted OR = 2.36, 95% CI: 1.12–4.97, \( p = 0.024 \), on the codominant model; adjusted OR = 2.68, 95% CI: 1.39–5.18, \( p = 0.003 \), on the recessive model). Opposing, ERCC5 rs2227869 heterozygotes (adjusted OR = 0.25, 95% CI: 0.07–0.88, \( p = 0.030 \)) and variant allele carriers (adjusted OR = 0.32, 95% CI: 0.11–0.97, \( p = 0.044 \)) as well as ERCC5 rs17655 variant allele homozygotes (adjusted OR = 0.27, 95% CI: 0.08–0.95, \( p = 0.041 \), on the recessive model) presented a significant risk reduction among female patients. Among these gender-specific genetic effects, only the association with MSH6 rs1042821 had been reported in the original studies [18]. No significant association was observed in the male subset of patients, possibly because of the low number of cases in this gender group. An association between XRCC5 rs1051677 and TC risk had previously been identified in this subset of patients [15] but significance was lost upon restricting analysis to well-differentiated forms of TC (this study).

Stratified analysis according to the age of diagnosis had only been performed in some of our initial studies, namely those involving SNPs of the BER and MMR pathways [16,18], with negative results. We therefore extended this analysis to the remaining DNA repair SNPs, considering two age groups: <50 and \( \geq 50 \) years. In patients under 50 years of age, both homozygosity for the XPC rs2228001 variant allele (adjusted OR = 2.86, 95% CI: 1.01–8.08, \( p = 0.048 \), on the recessive model) and the presence of at least one XRCC5 rs2440 variant allele (adjusted OR = 2.53, 95% CI: 1.02–6.26, \( p = 0.045 \)) were associated with increased DTC risk. When restricting the analysis to patients with 50 or more years of age, DTC
risk was increased in CCNH rs2230641 heterozygotes (adjusted OR = 2.91, 95% CI: 1.51–5.60, \( p = 0.001 \)) and variant allele carriers (adjusted OR = 3.04, 95% CI: 1.59–5.81, \( p = 0.001 \)), in RAD51 rs1801321 variant allele homozygotes (adjusted OR = 2.99, 95% CI: 1.25–7.14, \( p = 0.014 \), on the codominant model; unadjusted OR = 2.03, 95% CI: 1.00–4.12, \( p = 0.049 \), on the recessive model) and variant allele carriers (adjusted OR = 2.14, 95% CI: 1.06–4.32, \( p = 0.034 \)) and in XRCC3 rs861539 variant allele homozygotes (adjusted OR = 2.63, 95% CI: 1.16–5.97, \( p = 0.021 \), on the recessive model). On the contrary, the presence of at least one variant ERCC6 rs2228529 allele (adjusted OR = 0.47, 95% CI: 0.24–0.92, \( p = 0.028 \)) and its presence in heterozygosity (adjusted OR = 0.48, 95% CI: 0.24–0.97, \( p = 0.042 \)) were associated with a DTC risk reduction in this older age group.

No further correlations between individual DNA repair SNPs and DTC risk were observed on histology-, gender- and age-based stratification analysis.
Table 4. Genotype distribution in the case population (n = 106) and associated DTC risk (crude and adjusted ORs), after stratification according to histological type, gender and age. Only SNPs presenting significant findings are shown.

| Genotype | Papillary Carcinoma |  | Follicular Carcinoma |  |
|----------|---------------------|---|---------------------|---|
|          | n (%)               | Crude OR (95% CI) | Adjusted OR (95% CI) a | n (%) | Crude OR (95% CI) | Adjusted OR (95% CI) a |
| MUTYH rs2219489 |                       |                |                        |        |                    |                          |
| Gln/Gln  | 48 (61.5)           | 1 (reference) | 1 (reference)          | 15 (53.6) | 1 (reference) | 1 (reference)          |
| Gln/His  | 27 (34.6)           | **0.56 (0.32–1.00)** b | 11 (39.3) | 0.95 (0.37–2.43) | 1.09 (0.40–2.92) |
| His/His  | 3 (3.8)             | 0.66 (0.16–2.68) | 2 (7.1) | 4.13 (0.35–49.28) | 6.97 (0.47–104.26) |
| Dominant model |       | **0.57 (0.33–0.99)** b | 13 (46.4) | 1.08 (0.43–2.67) | 1.27 (0.49–3.29) |
| Recessive model |      | 3 (3.8) | 0.85 (0.21–3.36) | 2 (7.1) | 4.23 (0.37–48.8) | 6.75 (0.46–98.39) |
| ERCC5 rs2227869 |                       |                |                        |        |                    |                          |
| Cys/Cys  | 75 (96.2)           | 1 (reference) | 1 (reference)          | 24 (85.7) | 1 (reference) | 1 (reference)          |
| Cys/Ser  | 3 (3.8)             | **0.24 (0.07–0.84)** b | 3 (10.7) | 1.28 (0.28–5.78) | 1.20 (0.26–5.61) |
| Ser/Ser  | 0 (0.0)             | – – | – – | – – | – – |
| Dominant model |       | **0.23 (0.07–0.80)** b | 4 (14.3) | 1.70 (0.42–6.90) | 1.61 (0.38–6.74) |
| Recessive model |      | 0 (0.0) | – – | – – | – – |
| XPC rs2228001 |                       |                |                        |        |                    |                          |
| Lys/Lys  | 26 (33.3)           | 1 (reference) | 1 (reference)          | 13 (46.4) | 1 (reference) | 1 (reference)          |
| Lys/Gln  | 36 (46.2)           | 1.01 (0.35–1.85) | 11 (39.3) | 0.72 (0.27–1.91) | 0.91 (0.33–2.54) |
| Gln/Gln  | 16 (20.5)           | 2.27 (0.99–5.22) | 4 (14.3) | 1.18 (0.28–4.96) | 1.05 (0.24–4.65) |
| Dominant model |       | 52 (66.7) | 1.22 (0.69–2.15) | 15 (53.6) | 0.80 (0.32–2.01) | 0.94 (0.36–2.44) |
| Recessive model |      | 17 (21.8) | 1.08 (0.56–2.10) | 8 (28.6) | 1.64 (0.57–4.69) | 1.67 (0.55–5.02) |
| MLH1 rs175080 |                       |                |                        |        |                    |                          |
| Pro/Pro  | 19 (24.4)           | 1 (reference) | 1 (reference)          | 3 (10.7) | 1 (reference) | 1 (reference)          |
| Pro/Leu  | 42 (53.8)           | 1.13 (0.59–2.19) | 17 (60.7) | 3.78 (0.97–14.79) | 3.61 (0.88–14.85) |
| Leu/Leu  | 17 (21.8)           | 1.21 (0.53–2.61) | 8 (28.6) | 4.36 (0.95–20.04) | 4.29 (0.89–20.78) |
| Dominant model |       | 59 (75.6) | 1.14 (0.61–2.14) | 25 (89.3) | **3.95 (1.05–14.81)** b | 3.81 (0.97–14.95) |
| Recessive model |      | 17 (21.8) | 1.08 (0.56–2.10) | 8 (28.6) | 1.64 (0.57–4.69) | 1.67 (0.55–5.02) |
| MSH6 rs1042821 |                       |                |                        |        |                    |                          |
| Gly/Gly  | 49 (62.8)           | 1 (reference) | 1 (reference)          | 19 (67.9) | 1 (reference) | 1 (reference)          |
| Gly/Glu  | 24 (30.8)           | 0.74 (0.41–1.32) | 6 (21.4) | 0.65 (0.22–1.91) | 0.76 (0.24–2.35) |
| Glu/Glu  | 6 (7.4)             | 2.30 (0.59–9.85) | 3 (10.7) | 5.84 (0.57–60.03) | 20.96 (1.08–406.53) b |
| Dominant model |       | 29 (37.2) | 0.83 (0.48–1.46) | 9 (32.1) | 0.92 (0.35–2.43) | 1.10 (0.39–3.07) |
| Recessive model |      | 5 (6.4) | 2.57 (0.67–9.85) | 3 (10.7) | 6.60 (0.65–66.63) | **23.70 (1.25–449.32)** b |
Table 4. Cont.

| Genotype | Male                          | Female                        |
|----------|-------------------------------|-------------------------------|
|          | Genotype | OR (95% CI) | Adjusted OR | Genotype | OR (95% CI) | Adjusted OR |
| NBN rs1805794 | 78 (100) | 1 (reference) | 1 (reference) | 28 (100) | 1 (reference) | 1 (reference) |
| GLU/Glu    | 42 (53.8) | 1.17 (0.66–2.07) | 0.90 (0.33–2.41) | 0.72 (0.25–2.05) |
| GLU/Gln    | 3 (3.8) | 0.94 (0.54–1.63) | 1.15 (0.47–2.86) | 0.90 (0.34–2.39) |
| Dominant model | 36 (46.2) | 0.95 (0.55–1.64) | 1.15 (0.47–2.86) | 0.90 (0.34–2.39) |
| Recessive model | 3 (3.8) | 0.29 (0.08–1.01) | 2.69 (0.62–11.71) | 2.23 (0.44–11.18) |
| XRCC2 rs3218536 | 78 (100) | 1 (reference) | 1 (reference) | 28 (100) | 1 (reference) | 1 (reference) |
| Arg/Arg    | 66 (84.6) | 1.17 (0.54–2.52) | 2.19 (0.55–9.78) | 0.20 (0.04–1.05) |
| Arg/His    | 12 (15.4) | 1.17 (0.54–2.52) | 2.19 (0.55–9.78) | 0.20 (0.04–1.05) |
| His/His    | 0 (0.0) | – | – | – |
| Dominant model | 12 (15.4) | 1.17 (0.54–2.52) | 2.19 (0.55–9.78) | 0.20 (0.04–1.05) |
| Recessive model | 0 (0.0) | – | – | – |
| XRCC3 rs861539 | 78 (100) | 1 (reference) | 1.00 (0.39–2.58) | 0.78 (0.27–2.24) |
| Thr/Thr    | 26 (33.3) | 0.75 (0.40–1.40) | 0.71 (0.36–1.43) | 0.71 (0.36–1.43) |
| Thr/Met    | 31 (39.7) | 0.81 (0.30–2.20) | 0.81 (0.30–2.20) | 0.78 (0.27–2.24) |
| Met/Met    | 21 (26.9) | 2.50 (0.55–11.41) | 2.50 (0.55–11.41) | 2.72 (0.54–13.60) |
| Dominant model | 52 (66.7) | 0.97 (0.54–1.73) | 0.97 (0.54–1.73) | 1.00 (0.37–2.69) |
| Recessive model | 21 (26.9) | 2.08 (1.07–4.06) | 2.08 (1.07–4.06) | 1.00 (0.37–2.69) |
| CCN1 rs2230641 | 16 (100) | 1 (reference) | 1 (reference) | 90 (100) | 1 (reference) | 1 (reference) |
| Val/Val    | 7 (43.8) | 1.67 (0.44–6.34) | 1.26 (0.30–5.20) | 1.26 (0.30–5.20) |
| Val/Ala    | 9 (56.3) | 1.67 (0.44–6.34) | 1.26 (0.30–5.20) | 1.26 (0.30–5.20) |
| Ala/Ala    | 0 (0.0) | – | – | – |
| Dominant model | 9 (56.3) | 1.21 (0.36–4.06) | 1.21 (0.36–4.06) | 0.34 (0.11–1.01) |
| Recessive model | 0 (0.0) | – | – | – |
| ERCC5 rs2227869 | 16 (100) | 1 (reference) | 1 (reference) | 90 (100) | 1 (reference) | 1 (reference) |
| Cys/Cys    | 13 (81.3) | 1.40 (0.38–5.17) | 1.01 (0.25–4.12) | 1.01 (0.25–4.12) |
| Cys/Ser    | 3 (18.8) | 0.94 (0.19–4.62) | 0.94 (0.19–4.62) | 0.94 (0.19–4.62) |
| Ser/Ser    | 0 (0.0) | – | – | – |
| Dominant model | 3 (18.8) | 0.94 (0.19–4.62) | 0.94 (0.19–4.62) | 0.94 (0.19–4.62) |
| Recessive model | 0 (0.0) | – | – | – |
| Genotype       | n (%) | OR (95% CI)       | Adjusted OR (95% CI) | n (%) | OR (95% CI)       | Adjusted OR (95% CI) |
|----------------|-------|-------------------|----------------------|-------|-------------------|----------------------|
|                | <50 years |                  |                      | ≥50 years |                  |                      |
| ERCC5 rs17655  |       |                   |                      |        |                   |                      |
| Asp/Asp        | 16 (100) | 1 (reference)    | 1 (reference)        | 89 (100) | 1 (reference)    | 1 (reference)        |
| Asp/His        | 10 (62.5) | 0.61 (0.17-2.20)  | 0.63 (0.17-2.34)     | 45 (50.6) | 1.38 (0.81-2.33)  | 1.36 (0.80-2.30)     |
| His/His        | 1 (63)  | 0.31 (0.09-1.10)  | 0.32 (0.09-1.14)     | 3 (3.4)  |                   |                      |
| Dominant model | 6 (37.5) | 0.76 (0.22-2.67)  | 1.13 (0.68-1.88)     | 48 (53.9) | 1.13 (0.68-1.89)  |                      |
| Recessive model| 1 (63)  | 3 (3.4)           |                      | 0.27 (0.08-0.92) | 0.27 (0.08-0.95) |                      |
| XPC rs2228001  |       |                   |                      |        |                   |                      |
| Lys/Lys        | 9 (56.3) | 1 (reference)    | 1 (reference)        | 30 (33.3) | 1 (reference)    | 1 (reference)        |
| Lys/Gln        | 6 (37.5) | 0.58 (0.17-2.05)  | 0.59 (0.16-2.20)     | 41 (45.6) | 1.01 (0.57-1.78)  | 1.05 (0.59-1.86)     |
| Gin/Gln        | 1 (63)  | 1.22 (0.06-23.8)  | 19 (21.1)            | 2.05 (0.96-4.36) | 2.05 (0.96-4.38)   |                      |
| Dominant model | 7 (43.8) | 0.63 (0.18-2.27)  | 60 (66.7)            | 1.20 (0.71-2.05) | 1.24 (0.72-2.12)  |                      |
| Recessive model| 1 (63)  | 3 (3.4)           |                      | 0.27 (0.08-0.92) | 0.27 (0.08-0.95) |                      |
| MSH6 rs1042821 |       |                   |                      |        |                   |                      |
| Gly/Gly        | 11 (68.8) | 1 (reference)    | 1 (reference)        | 57 (63.3) | 1 (reference)    | 1 (reference)        |
| Gly/Glu        | 4 (25.0) | 0.86 (0.24-3.54)  | 26 (28.9)            | 0.70 (0.41-1.22) | 0.70 (0.40-1.22) |                      |
| Glu/Glu        | 1 (63)  | 1.08 (0.07-16.53) | 7 (7.8)              | 4.42 (1.10-17.75) | 4.78 (1.17-19.56) |                      |
| Dominant model | 5 (31.2) | 0.98 (0.24-4.24)  | 33 (36.7)            | 0.86 (0.51-1.44) | 0.86 (0.51-1.45) |                      |
| Recessive model| 1 (63)  | 1.09 (0.06-15.6)  | 7 (7.8)              | 5.00 (1.26-19.94) | 5.42 (1.34-21.92) |                      |
| XRCC3 rs861539 |       |                   |                      |        |                   |                      |
| Thr/Thr        | 8 (50.0) | 1 (reference)    | 1 (reference)        | 38 (42.2) | 0.80 (0.44-1.76)  | 0.81 (0.45-1.46)     |
| Thr/Met        | 6 (37.5) | 0.47 (0.07-3.28)  | 24 (26.7)            | 2.26 (1.09-4.71) | 2.36 (1.12-4.97)  |                      |
| Met/Met        | 2 (12.5) | 0.58 (0.16-2.08)  | 62 (68.9)            | 1.06 (0.62-1.83) | 1.08 (0.63-1.88)  |                      |
| Dominant model | 8 (50.0) | 0.60 (0.10-3.67)  | 24 (26.7)            | 2.60 (1.36-4.95) | 2.68 (1.39-5.18)  |                      |
| Recessive model| 2 (12.5) | 28 (31.1)         |                      | 0.81 (0.45-1.46) | 0.81 (0.45-1.46) |                      |

**Table 4. Cont.**

| Genotype       | n (%) | OR (95% CI)       | Adjusted OR (95% CI) | n (%) | OR (95% CI)       | Adjusted OR (95% CI) |
|----------------|-------|-------------------|----------------------|-------|-------------------|----------------------|
|                | CCNH rs2230641 |                  |                      |       |                   |                      |
| Val/Val        | 16 (100) | 1 (reference)    | 1 (reference)        | 64 (100) | 1 (reference)    | 1 (reference)        |
| Val/Ala        | 27 (64.3) | 0.96 (0.43-2.13)  | 29 (45.3)            | 2.97 (1.55-5.68) | 2.91 (1.51-5.60)  |                      |
| Ala/Ala        | 1 (2.4)  | 0.27 (0.03-2.33)  | 2 (3.1)              | 5.94 (0.52-67.64) | 8.01 (0.62-102.77) |                      |
| Dominant model | 15 (37.7) | 0.79 (0.36-1.75)  | 31 (48.4)            | 3.07 (1.62-5.81) | 3.04 (1.59-5.81)  |                      |
| Recessive model| 1 (2.4)  | 2.60 (1.36-4.95)  | 2.68 (1.39-5.18)     |       |                   |                      |
### Table 4. Cont.

| Gene   | rs Number | Genotype | Case (n=42) | Control (n=62) | p-Value | 95% CI       | p-Value | 95% CI       |
|--------|-----------|----------|-------------|----------------|---------|--------------|---------|--------------|
| ERCC6  | rs2228529 | Gln/Gln  | 20 (47.6)   | 1 (reference)  | 46 (74.2) | 0.49 (0.25-0.98) | 0.48 (0.24-0.97) |
|        |           | Gln/Arg  | 20 (47.6)   | 1.19 (0.56-2.54) | 15 (24.2) | 0.032 (0.04-2.84) | 0.03 (0.03-2.63) |
|        |           | Arg/Arg  | 2 (4.8)     | 2.20 (0.29-16.75) | 1 (1.6)   | 1.09 (0.53-2.44) | 1.09 (0.50-2.36) |
|        |           |          |             |                |         |              |         |              |
|        |           | Dominant | 22 (52.4)   | 1.24 (0.59-2.61) | 16 (25.8) | 0.48 (0.24-0.93) | 0.47 (0.24-0.92) |
|        |           | Recessive| 2 (4.8)     | 2.03 (0.28-14.91) | 1 (1.6)   | 1.09 (0.53-2.44) | 1.09 (0.50-2.36) |
|        |           |          |             |                |         |              |         |              |
| XPC    | rs2228001 | Lys/Lys  | 17 (40.5)   | 1 (reference)  | 22 (34.4) | 1.09 (0.53-2.44) | 1.09 (0.50-2.36) |
|        |           | Lys/Gln  | 15 (35.7)   | 0.58 (0.25-1.32) | 32 (50.0) | 1.22 (0.63-2.35) | 1.27 (0.66-2.48) |
|        |           | Gln/Gln  | 10 (23.8)   | 2.21 (0.73-6.65) | 10 (15.6) | 1.69 (0.65-4.38) | 1.74 (0.66-4.57) |
|        |           |          |             |                |         |              |         |              |
|        |           | Dominant | 25 (59.5)   | 0.82 (0.36-1.75) | 42 (65.6) | 1.31 (0.70-2.44) | 1.36 (0.72-2.56) |
|        |           | Recessive| 10 (23.8)   | 2.03 (0.28-14.91) | 10 (15.6) | 1.51 (0.63-3.61) | 1.52 (0.63-3.67) |
| RAD51  | rs1801321 | G/G      | 14 (33.3)   | 1 (reference)  | 24 (39.7) | 2.90 (1.23-6.83) | 2.99 (1.25-7.14) |
|        |           | G/T      | 19 (45.2)   | 0.95 (0.41-2.24) | 31 (48.4) | 1.76 (0.84-3.69) | 1.83 (0.87-3.86) |
|        |           | T/T      | 9 (21.4)    | 0.80 (0.29-2.20) | 19 (29.7) | 1.75 (0.72-4.08) | 1.76 (0.72-4.08) |
|        |           |          |             |                |         |              |         |              |
|        |           | Dominant | 28 (66.7)   | 0.90 (0.41-1.98) | 50 (78.1) | 2.07 (1.04-4.14) | 2.14 (1.06-4.32) |
|        |           | Recessive| 9 (21.4)    | 0.82 (0.34-1.99) | 19 (29.7) | 2.03 (1.00-4.12) | 2.05 (1.00-4.21) |
| XRCC3  | rs861539  | Thr/Thr  | 15 (35.7)   | 1 (reference)  | 21 (32.8) | 1.09 (0.53-2.17) | 1.10 (0.55-2.27) |
|        |           | Thr/Met  | 16 (38.1)   | 0.65 (0.27-1.52) | 28 (43.8) | 0.85 (0.43-1.78) | 0.87 (0.44-1.73) |
|        |           | Met/Met  | 11 (26.2)   | 1.47 (0.53-4.08) | 15 (23.4) | 2.25 (0.92-5.49) | 2.42 (0.97-6.03) |
|        |           |          |             |                |         |              |         |              |
|        |           | Dominant | 27 (64.3)   | 0.84 (0.38-1.83) | 43 (67.2) | 1.09 (0.57-2.05) | 1.12 (0.59-2.14) |
|        |           | Recessive| 11 (26.2)   | 1.88 (0.76-4.67) | 15 (23.4) | 2.47 (1.11-5.41) | 2.63 (1.16-5.97) |
| XRCC5  | rs24440   | C/C      | 8 (19.0)    | 1 (reference)  | 22 (35.5) | 1.00 (0.52-1.95) | 0.97 (0.50-1.90) |
|        |           | C/T      | 23 (54.8)   | 2.25 (0.88-5.77) | 31 (50.0) | 1.00 (0.52-1.95) | 0.97 (0.50-1.90) |
|        |           | T/T      | 11 (26.2)   | 2.35 (0.79-6.98) | 9 (14.5)  | 1.28 (0.49-3.38) | 1.29 (0.48-3.45) |
|        |           |          |             |                |         |              |         |              |
|        |           | Dominant | 34 (81.0)   | 2.28 (0.94-5.57) | 40 (64.5) | 1.06 (0.56-1.99) | 1.03 (0.54-1.95) |
|        |           | Recessive| 11 (26.2)   | 1.38 (0.58-3.29) | 9 (14.5)  | 1.28 (0.53-3.11) | 1.31 (0.53-3.23) |

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a ORs were adjusted for gender (male and female), age (<30, 30–49, 50–69, and ≥70 years), and smoking status (non-smoker and smoker). b Significant results (p < 0.05) highlighted in bold.

Abbreviations: DTC, well-differentiated thyroid cancer; SNP, single nucleotide polymorphism; OR, odds ratio; CI, confidence interval.
3.4. Combined Genotypes

In order to investigate the joint effect of multiple SNPs on DTC risk, genetic risk scores (RS) were calculated for each study participant, considering only significant findings on single SNP analysis. As depicted in Table 5, after adjusting for covariates, DTC risk was more than two and five times higher in individuals bearing, respectively, 2 (adjusted OR = 2.68, 95% CI: 1.56–4.59, \( p < 0.001 \)) and 3 or more (adjusted OR = 5.02, 95% CI: 2.24–11.24, \( p = 0.001 \)) risk genotypes (CCNH rs2230641 Val/Ala or Ala/Ala; ERCC5 rs2227869 Cys/Cys or Ser/Ser; XPC rs2228001 Gln/Gln; MSH6 rs1042821 Glu/Glu; XRCC3 rs861539 Met/Met), when compared to individuals bearing none or only one of such risk genotypes. Similar associations between RS and TC risk were also observed on stratification according to histological, gender or age criteria, after adapting RS calculations to the SNPs significant for each strata (Table 5). A high significance level was observed in most cases (\( p < 0.001 \) in approximately 50% of RS categories) and was even greater if higher RS categories were merged together (results not shown).

Also, in order to investigate the combined effect of different pairs of SNPs on DTC risk, we performed a paired SNP analysis considering all possible 2 × 2 combinations of the DNA repair SNPs included in this study. Overall, 595 SNP-SNP combinations were tested, 114 (approximately 20%) of which yielded significant results at a 0.05 significance level (results not shown). Considering that such a high number of hypothesis being tested may result in a considerable number of false positive findings, a more stringent significance level (\( p < 0.01 \)) was employed in this analysis, limiting the number of SNP pairs with significant findings to 15 (approximately 2.5% of all possible combinations). Such significant findings are depicted in Table 6 and also in Figure 1. CCNH rs2230641 emerges from Figure 1 as the DNA repair SNP most frequently represented in significant SNP-SNP combinations, both at 0.01 and 0.05 significance levels, followed by RAD51 rs1801321, MLH3 rs175080 and MSH4 rs5745549 (0.01 significance level) or RAD51 rs1801321 and XRCC3 rs861539 (0.05 significance level). MMR variants were the most frequently involved as they were present in 9 of the 15 SNP-SNP combinations that were significant. Also, among significant findings, 3 intra-pathway SNP combinations were detected: RAD51 rs1801321–XRCC3 rs861539 (HR pathway), MLH3 rs175080–MSH6 rs1042821 and MSH4 rs5745549–MSH6 rs1042821 (MMR pathway).

Finally, haplotype analysis was applied to SNPs located in the same chromosome arm, since these are likely to segregate together. According to such criteria, it was possible to establish 8 blocks of DNA repair SNPs, of which only one, located on chromosome 5q and comprising 6 SNPs (CCNH rs2230641, CDK7 rs2972388, MSH3 rs26279, MSH3 rs184967, XRCC4 rs1805377 and XRCC4 rs28360135), revealed significant associations with DTC (Table 7): two different allele combinations were associated with a significantly decreased DTC risk, when compared to the most frequent combination of chromosome 5q SNPs (adjusted OR1 = 0.26, 95% CI: 0.08–0.87, \( p = 0.030 \); adjusted OR2 = 0.15, 95% CI: 0.03–0.72, \( p = 0.019 \)). Haplogroup analysis comprising all SNPs under study could also prove useful to understand the joint effect of the variants since it would better reflect the real context situation (where different DNA repair proteins interact with each other) but could not be performed because, considering the high number of SNPs under study, the frequency of each specific allele combination would be too low for meaningful results to be obtained.
Figure 1. SNP frequency (%) in SNP-SNP pairs showing significant results at $p < 0.01$ and $p < 0.05$ levels. Only SNPs presenting significant results ($p < 0.05$) on combined genotype analysis are shown.

Table 5. Risk score (RS) in case and control populations and associated DTC risk (crude and adjusted ORs). Risk scores calculated from significant results on single SNP analysis.

| Risk Score (RS) * | Frequency | p-Value b | OR (95% CI) | p-Value | Adjusted OR (95% CI) c | p-Value b |
|-------------------|-----------|-----------|-------------|---------|------------------------|---------|
| DTC (all cases)   | 191 (100) | 0.05      | 1 (Reference) |        | 1 (Reference)          |<0.001 d |
| 0-1               | 114 (59.7) | 1 (Reference) |        | <0.001 | 1 (Reference)          |<0.001 d |
| 2                 | 64 (33.5)  | 5.16 (2.33–11.44) | <0.001 |<0.001 | 5.02 (2.24–11.24)      |<0.001 d |
| 3/4               | 13 (6.8)   | 2.72 (1.60–4.63)  | <0.001 |<0.001 | 2.68 (1.56–4.59)       |<0.001 d |
| Papillary TC      | 152 (100) | 4.58 (2.36–8.89)  | <0.001 |<0.001 | 4.55 (2.34–8.84)       |<0.001 d |
| 0–2               | 85 (55.9)  | 4.47 (1.94–10.32) | <0.001 |<0.001 | 4.46 (1.92–10.36)      |<0.001 d |
| 3                 | 48 (31.6)  | 1 (Reference)    |        |        | 1 (Reference)          |<0.001 d |
| 4/+               | 19 (12.5)  | 1 (Reference)    |        |        | 1 (Reference)          |<0.001 d |
| Follicular TC     | 56 (100)  | 3.45 (1.15–10.39) | 0.028  |<0.001 | 3.52 (1.12–11.07)      | 0.032 d |
| 0–1               | 24 (42.9)  | 1 (Reference)    |        |        | 1 (Reference)          |<0.001 d |
| 2/+               | 32 (57.1)  | 3.43 (1.92–6.13)  | <0.001 |<0.001 | 3.42 (1.90–6.14)       |<0.001 d |
| Gender            |           |             |             |         |                        |         |
| Female            | 174 (100) | 8.14 (3.31–20.94) | <0.001 |<0.001 | 8.01 (3.22–19.92)      |<0.001 d |
| 0–2               | 114 (65.5) | 1 (Reference)    |        |        | 1 (Reference)          |<0.001 d |
| 3                 | 51 (29.3)  | 3.32 (1.31–8.01)  | <0.001 |<0.001 | 3.23 (1.24–8.04)       |<0.001 d |
| 4/+               | 9 (5.2)    | 1 (Reference)    |        |        | 1 (Reference)          |<0.001 d |
| Age               |           |             |             |         |                        |         |
| <50 years         | 83 (100)  | 3.43 (1.92–6.13)  | <0.001 |<0.001 | 3.42 (1.90–6.14)       |<0.001 d |
| 0                 | 26 (31.3)  | 2.33 (0.66–8.54)  | 0.097  |        | 2.32 (0.92–6.94)       |0.073 d  |
| 1                 | 52 (62.7)  | 6.93 (1.64–28.89) | 0.008  |        | 7.34 (1.72–31.24)      | 0.007 d  |
| 2                 | 5 (6.0)    | 3.43 (1.92–6.13)  | <0.001 |<0.001 | 3.23 (1.24–8.04)       |<0.001 d |
| ≥50 years         | 127 (100) | 1 (Reference)    |        |        | 1 (Reference)          |<0.001 d |
| 0–1               | 60 (47.2)  | 1 (Reference)    |        |        | 1 (Reference)          |<0.001 d |
| 2                 | 51 (40.2)  | 2.55 (1.17–5.61)  | 0.019  |        | 2.66 (1.21–5.83)       |0.015 d  |
| 3/4               | 16 (12.6)  | 7.50 (3.09–19.18) | <0.001 |<0.001 | 7.90 (2.21–19.45)      |<0.001 d |

* For the purpose of risk score calculations, genotypes presenting significant results on single SNP analysis were attributed a +1 score, risk score for each participant corresponding to the sum of such scores (+1 in all cases: CCNH rs2230641 Val/Ala or Ala/Ala + ERCC5 rs2227869 Cys/Cys or Ser/Ser + XPC rs2228001 Gln/Gln + MSH6 rs1042821 Glu/Glu + XRC3 rs861539 Met/Met; +1 in papillary TC: MTHYH rs3219489 Gln/Gln + ERCC5 rs2227869 Cys/Cys + XPC rs2228001 Gln/Gln + NBN rs1805794 Glu/Glu or Gln/Gln + XRC3 rs861539 Met/Met; +1 in follicular TC: MLH3 rs175808 Pro/Leu or Leu/Leu + MSH6 rs1042821 Gln/Glu + XRC2 rs3218536 Arg/Arg; +1 in female participants: CCNH rs2230641 Val/Ala or Ala/Ala + ERCC5 rs2227869 Cys/Cys + ERCC5 rs17655 Asp/Asp or Asp/His + XPC rs2228001 Gln/Gln + MSH6 rs1042821 Gln/Glu + XRC3 rs861539 Met/Met; +1 in participants with age <50 years: XPC rs2228001 Gln/Gln + XRC5 rs2440 C/T or T/T; +1 in participants with age ≥50 years: CCNH rs2230641 Val/Ala or Ala/Ala + ERCC5 rs2228001 Gln/Gln + RAD51 rs1801321 G/G or T/T + XRC3 rs861539 Met/Met). b p-value for cases versus control group determined by two-sided Fisher’s exact test (whenever $2 \times 2$ contingency tables are possible) or $\chi^2$ test (remaining cases). c ORs were adjusted for gender (male and female), age (<30, 30–49, 50–69, ≥70 years) and smoking status (non-smoker and smoker). d $p < 0.05$. Abbreviations: DTC, well-differentiated thyroid cancer; MAF, minor allele frequency; OR, odds ratio; CI, confidence interval.
**Table 6.** Two-way SNP interactions among DNA repair genes: distribution of combined genotypes in enrolled populations and associated DTC risk (adjusted ORs). Only SNPs presenting significant findings ($p < 0.01$) are shown.

| Combined Genotype | Frequency | DTC Risk |  |
|-------------------|-----------|----------|---|
|                   | Controls n (%) | Cases n (%) | $p$-Value | Adjusted OR (95% CI) | $p$-Value |
| **CCNH rs2230641 – RAD51 rs1801321** | | | | |
| Val/Val – G/G | 58 (27.4) | 13 (12.3) | 0.037 | 1 (Reference) | |
| Val/Val – G/T | 64 (30.2) | 29 (27.4) | 2.10 (0.99–4.45) | 0.052 | |
| Val/Ala – G/G | 15 (7.1) | 13 (12.3) | 3.77 (1.44–9.87) | 0.007 | |
| Val/Ala – G/T | 27 (12.7) | 20 (18.9) | 3.43 (1.46–8.06) | 0.005 | |
| Val/Val – T/T | 26 (12.3) | 18 (17.0) | 3.05 (1.29–7.19) | 0.011 | |
| Ala/Ala – G/G | 14 (6.6) | 10 (9.4) | 3.22 (1.17–8.06) | 0.024 | |
| Ala/Ala – G/T | 8 (3.8) | 3 (2.8) | 1.86 (0.42–8.18) | 0.414 | |
| Ala/Ala – T/T | 212 (100) | 106 (100) | | | |
| **MUTYH rs3219489 – CCNH rs2230641** | | | | |
| Gin/Gln – Val/Val | 77 (36.5) | 35 (33.0) | 0.018 | 1 (Reference) | |
| Gin/Gln – Val/Ala | 22 (10.4) | 26 (24.5) | 2.68 (1.32–5.42) | 0.006 | |
| Gin/His – Val/Val | 66 (31.3) | 23 (21.7) | 0.81 (0.43–1.51) | 0.500 | |
| Gin/His – Val/Ala | 30 (14.2) | 14 (13.2) | 1.05 (0.49–2.23) | 0.904 | |
| Gin/Gln – Ala/Ala | 8 (3.8) | 3 (2.8) | 1.86 (0.42–8.18) | 0.414 | |
| Gin/Gln – Ala/Ala | 211 (100) | 106 (100) | | | |
| **CCNH rs2230641 – MLH3 rs175080** | | | | |
| Val/Val – Pro/Pro | 40 (20.5) | 11 (10.4) | 0.097 | 1 (Reference) | |
| Val/Val – Pro/Leu | 77 (39.5) | 36 (34.0) | 1.76 (0.80–3.87) | 0.162 | |
| Val/Ala – Pro/Pro | 14 (7.2) | 11 (10.4) | 2.60 (0.91–7.41) | 0.074 | |
| Val/Ala – Pro/Leu | 41 (21.0) | 38 (35.8) | 2.27 (1.30–3.96) | 0.004 | |
| Val/Val – Leu/Leu | 5 (2.6) | 3 (2.8) | 2.44 (0.48–12.45) | 0.284 | |
| Val/Ala – Leu/Leu | 132 (67.7) | 51 (48.1) | 0.009 | 1 (Reference) | |
| Val/Val – Ser/Ser | 10 (5.1) | 9 (8.5) | 2.45 (0.93–6.43) | 0.070 | |
| Val/Ala – Ser/Ser | 41 (21.0) | 38 (35.8) | 2.27 (1.30–3.96) | 0.004 | |
| Val/Val – Leu/Leu | 12 (6.2) | 8 (7.5) | 1.87 (0.71–4.92) | 0.207 | |
| Val/Ala – Leu/Leu | 195 (100) | 106 (100) | | | |
| **CCNH rs2230641 – MSH4 rs5745549** | | | | |
| Val/Val – Ser/Asn | 132 (67.7) | 51 (48.1) | 0.009 | 1 (Reference) | |
| Val/Val – Ser/Asn | 10 (5.1) | 9 (8.5) | 2.45 (0.93–6.43) | 0.070 | |
| Val/Ala – Ser/Asn | 41 (21.0) | 38 (35.8) | 2.27 (1.30–3.96) | 0.004 | |
| Val/Val – Leu/Leu | 12 (6.2) | 8 (7.5) | 1.87 (0.71–4.92) | 0.207 | |
| Val/Ala – Leu/Leu | 195 (100) | 106 (100) | | | |
| **MLH3 rs175080 – RAD51 rs1801321** | | | | |
| Pro/Pro – G/G | 23 (11.8) | 4 (3.8) | 0.288 | 1 (Reference) | |
| Pro/Pro – G/T | 24 (12.3) | 10 (9.4) | 3.98 | 1.78 (0.77–10.78) | 0.117 | |
| Pro/Leu – G/G | 32 (16.4) | 18 (17.0) | 3.98 | (1.14–13.89) | 0.031 | |
| Pro/Leu – G/T | 46 (23.6) | 25 (23.6) | 3.98 | (1.09–11.81) | 0.035 | |
| Pro/Pro – T/T | 9 (4.6) | 8 (7.5) | 3.98 | (1.23–23.88) | 0.025 | |
| Leu/Leu – G/G | 14 (7.2) | 6 (5.7) | 2.92 | (0.66–12.57) | 0.151 | |
| Leu/Leu – G/T | 23 (11.8) | 16 (15.1) | 4.66 | (1.32–16.45) | 0.017 | |
| Leu/Leu – T/T | 16 (8.2) | 15 (14.2) | 6.22 | (1.70–22.78) | 0.006 | |
| Leu/Leu – T/T | 8 (4.1) | 4 (3.8) | 3.55 | (0.69–18.15) | 0.128 | |
Table 6. Cont.

| Combined Genotype | Frequency | DTC Risk |
|-------------------|-----------|----------|
|                   | Controls n (%) | Cases n (%) | p-Value * | Adjusted OR (95% CI) | p-Value * |
| **ERCC6** rs4253211 – **RAD51** rs1801321 | | | | | |
| Arg/Arg – G/G     | 65 (30.8) | 16 (15.7) | 0.026 6 | 1 (Reference) | 0.007 4 |
| Arg/Arg – G/T     | 72 (34.1) | 42 (41.2) | 2.51 (1.28–4.94) | 0.014 6 |
| Arg/Pro – G/T     | 21 (10.0) | 7 (6.9) | 1.53 (0.54–4.29) | 0.423 |
| Arg/Arg – T/T     | 33 (15.6) | 21 (20.6) | 2.67 (1.22–5.85) | 0.014 6 |
| Pro/Pro – G/G     | 20 (9.5) | 16 (15.7) | 3.65 (1.52–8.78) | 0.004 4 |
| **MLH3** rs175080 – **MSH6** rs1042821 | | | | | |
| Pro/Pro – Gly/Gly | 32 (15.2) | 19 (17.9) | 0.032 4 | 1 (Reference) | 0.561 |
| Pro/Pro – Gly/Glu | 26 (12.4) | 2 (1.9) | 0.11 (0.02–0.53) | 0.006 4 |
| Pro/Leu – Gly/Gly | 71 (33.8) | 36 (34.0) | 0.81 (0.40–1.65) | 0.878 |
| Pro/Leu – Gly/Glu | 35 (16.7) | 19 (17.9) | 0.94 (0.41–2.13) | 0.819 |
| Leu/Leu – Gly/Gly | 24 (11.4) | 13 (12.3) | 0.83 (0.34–2.03) | 0.660 |
| Leu/Leu – Gly/Glu | 17 (8.1)  | 9 (8.5)  | 0.89 (0.33–2.43) | 0.680 |
| **MSH4** rs5745549 – **MSH6** rs1042821 | | | | | |
| Ser/Ser – Gly/Gly | 124 (59.0) | 60 (56.6) | 0.004 4 | 1 (Reference) | 0.467 |
| Ser/Ser – Gly/Glu | 63 (30.0) | 24 (22.6) | 0.83 (0.30–2.28) | 0.720 |
| Ser/Asn – Gly/Gly | 15 (7.1)  | 6 (5.7)  | 0.88 (0.33–2.43) | 0.680 |
| Ser/Asn – Gly/Glu | 8 (3.8)   | 16 (15.1) | 4.63 (1.83–11.69) | 0.001 4 |
| **ERCC6** rs4253211 – **MLH3** rs175080 | | | | | |
| Arg/Arg – Pro/Pro | 51 (26.2) | 13 (12.7) | 0.067 4 | 1 (Reference) | 0.113 |
| Arg/Arg – Pro/Leu | 78 (40.0) | 45 (44.1) | 2.43 (1.18–5.04) | 0.017 4 |
| Arg/Pro – Pro/Leu | 21 (10.8) | 10 (9.8)  | 2.25 (0.83–6.14) | 0.113 |
| Arg/Leu – Pro/Pro | 30 (15.4) | 21 (20.6) | 2.96 (1.28–6.88) | 0.012 4 |
| **RAD51** rs1801321 – **XRCC3** rs861539 | | | | | |
| G/G – Thr/Thr     | 26 (12.4) | 7 (6.6)  | 0.006 4 | 1 (Reference) | 0.381 |
| G/G – Thr/Met     | 35 (16.7) | 15 (14.2) | 1.59 (0.56–4.49) | 0.423 |
| G/T – Thr/Thr     | 29 (13.9) | 24 (22.6) | 3.10 (1.14–8.44) | 0.029 8 |
| G/T – Thr/Met     | 55 (26.3) | 14 (13.2) | 0.98 (0.35–2.73) | 0.967 |
| G/G – Met/Met     | 11 (5.3)  | 6 (5.7)  | 1.99 (0.54–7.41) | 0.304 |
| T/T – Thr/Thr     | 15 (7.2)  | 5 (4.7)  | 1.23 (0.33–4.61) | 0.759 |
| G/T – Met/Met     | 12 (5.7)  | 12 (11.3) | 0.577 | 0.026 8 |
| T/T – Thr/Met     | 22 (10.5) | 15 (14.2) | 2.41 (0.83–7.05) | 0.108 |
| T/T – Met/Met     | 4 (1.9)   | 8 (7.5)  | 7.90 (1.60–34.74) | 0.006 4 |
| **ERCC6** rs2228529 – **MSH4** rs5745549 | | | | | |
| Gln/Gln – Ser/Ser | 102 (52.3) | 53 (51.0) | 0.009 4 | 1 (Reference) | 0.77 |
| Gln/Gln – Ser/Asn | 6 (3.1)   | 13 (12.5) | 4.77 (1.67–13.61) | 0.003 4 |
| Gln/Arg – Ser/Ser | 71 (36.4) | 34 (32.7) | 0.82 (0.48–1.43) | 0.489 |
| Gln/Arg – Ser/Asn | Arg/Arg – Ser/Ser | 16 (8.2) | 4 (3.8) | 0.46 (0.14–1.47) | 0.190 |
Table 6. Cont.

| Combined Genotype | Frequency | DTC Risk |
|-------------------|-----------|----------|
|                   | Controls n (%) | Cases n (%) | p-Value | Adjusted OR (95% CI) | p-Value |
| **MSH4 rs5745549 – XRCC5 rs2440** | | | | |
| Ser/Ser – C/C     | 67 (34.4) | 24 (23.1) | 0.049 | 1 (Reference) | |
| Ser/Ser – C/T     | 84 (43.1) | 50 (48.1) | 1.76 (0.97-3.19) | 0.063 | |
| Ser/Asn – C/T     | 12 (6.2) | 4 (3.8) | 1.02 (0.29-3.56) | 0.972 | |
| Ser/Asn – C/C     | 27 (13.8) | 17 (16.3) | 1.86 (0.84-4.12) | 0.124 | |
| Ser/Asn – T/T     | 5 (2.6) | 9 (8.7) | 6.18 (1.83-20.86) | 0.003 | |
| **MUTYH rs3219489 – XPC rs2228001** | | | | |
| Gln/Gln – Lys/Lys | 38 (18.0) | 28 (26.4) | 0.037 | 1 (Reference) | |
| Gln/His – Lys/Lys | 41 (19.4) | 9 (8.5) | 0.31 (0.13-0.73) | 0.008 | |
| Gln/His – Lys/Gln | 48 (22.7) | 18 (17.0) | 0.55 (0.26-1.16) | 0.117 | |
| Gln/Gln – Gln/Gln | 13 (6.2) | 8 (7.5) | 0.81 (0.29-2.25) | 0.689 | |
| Gln/His – Gln/Gln | 9 (4.3) | 11 (10.4) | 1.70 (0.61-4.77) | 0.311 | |
| **MSH3 rs184967 – XRCC5 rs1051685** | | | | |
| Arg/Arg – A/A     | 99 (50.8) | 70 (66.0) | 0.001 | 1 (Reference) | |
| Arg/Arg – A/G     | 32 (16.4) | 8 (7.5) | 0.34 (0.15-0.80) | 0.013 | |
| Arg/Gln – A/A     | 52 (26.7) | 14 (13.2) | 0.36 (0.18-0.71) | 0.003 | |
| Arg/Gln – A/G     | 12 (6.2) | 14 (13.2) | 1.46 (0.62-3.40) | 0.387 | |
| **CCNH rs2230641 – LIG4 rs1805386** | | | | |
| Val/Val – Thr/Thr | 112 (52.8) | 42 (39.6) | 0.015 | 1 (Reference) | |
| Val/Val – Thr/Ile | 32 (15.1) | 16 (15.1) | 1.36 (0.67-2.75) | 0.396 | |
| Val/Ala – Thr/Thr | 37 (17.5) | 36 (34.0) | 2.62 (1.45-4.71) | 0.001 | |
| Val/Ala – Thr/Ile | 18 (8.5) | 5 (4.7) | 0.73 (0.25-2.11) | 0.355 | |

* p value for cases versus control group determined by two–sided Fisher’s exact test (whenever 2x2 contingency tables are possible) or χ² test (remaining cases). ‡ ORs were adjusted for gender (male and female), age (<30, 30–49, 50–69, ≥70 years) and smoking status (non-smoker and smoker). ∗ p<0.05. †p < 0.01.

Table 7. Haplotypes comprising SNPs located in the same chromosome arm and corresponding DTC risk (adjusted ORs). Only haplotypes presenting significant results are shown.

| Haplotype | Adj. OR (95% CI) | p-Value *
|-----------|-----------------|----------|
| **Chromosome 5q** | | |
| **CCNH rs2230641** | | |
| Val/Val – Thr/Thr | 1.00 | 0.015 |
| Val/Val – Thr/Ile | 1.36 (0.67-2.75) | 0.396 |
| Val/Ala – Thr/Thr | 2.62 (1.45-4.71) | 0.001 |
| Val/Ala – Thr/Ile | 0.73 (0.25-2.11) | 0.355 |
| **CDK7 rs26972388** | | |
| Val/Val – Arg/Gln | 1.00 | (Reference) |
| Val/Val – Ala/Ile | 0.26 (0.08-0.87) | 0.03 |
| Val/Val – Thr/Ile | 0.15 (0.03-0.72) | 0.019 |

* p < 0.05. Abbreviations: DTC, well-differentiated thyroid cancer; OR, odds ratio; CI, confidence interval.
4. Discussion

In order to further characterize the potential contribution of DNA repair SNPs to DTC susceptibility, we aggregated and reanalysed the data from our previously published case-control studies [14–18] performed on a Caucasian Portuguese population.

A significant risk increase was observed, after adjustment for age, gender and smoking status, in CCNH rs2230641 heterozygotes and variant allele carriers, in MSH6 rs1042821 variant allele homozygotes (codominant and recessive model), in XRCC3 rs861539 variant allele homozygotes (recessive model) and in XPC rs2228001 variant allele homozygotes (recessive model), while the heterozygous ERCC5 rs2227869 genotype was associated with a borderline risk reduction. Except for XPC rs2228001, which is a new finding emerging from this reanalysis because the recessive model of inheritance had not been applied in the original study, such results are fundamentally similar to those reported on the original studies despite, on reanalysis, data was restricted to DTC cases and corresponding controls. A role for these variants specifically on well-differentiated forms of TC is thus apparent from this reanalysis. As these findings have been discussed in detail in the original studies, they will be discussed here only briefly, with emphasis on new data published since then.

XRCC3 participates in HR to maintain chromosome stability and repair DNA damage and is therefore a highly suspected candidate gene for cancer susceptibility. The XRCC3 rs861539 has been the most studied genetic variant of XRCC3 gene, especially because is located in a functional relevant domain of the protein, in an interaction region with other proteins such as RAD51 [22,32]. The presence of this variant may affect the structure of this DNA repair protein and lead to a deficiency in the HR pathway. As a result, the HR pathway may be compromised, shifting the repair mechanism to NHEJ, promoting chromosome instability and disturbing the cellular repair capacity [33]. The potential contribution of XRCC3 rs861539 to cancer susceptibility has been widely addressed: while conflicting evidence exists, several large meta-analyses strongly support a positive association with cancer susceptibility, namely breast [34–36] and bladder cancer [36–38], among others. In the particular context of thyroid cancer, interestingly, multiple studies [22,39–43], including a meta-analysis [44], have suggested the XRCC3 rs861539 variant T allele and/or, in particular, the TT homozygous genotype to be associated with increased risk of TC or, more specifically, PTC. In another meta-analysis [45] such association was also detected but only in Caucasian populations. Therefore, despite studies reporting no significant association also exist [46,47], the vast majority of available evidence supports our results and suggests a role for XRCC3 rs861539 in DTC susceptibility.

To the best of our knowledge, none of the remaining SNPs presenting significant results on overall analysis has been evaluated in the context of DTC (or TC) susceptibility.

XPC codes for a DNA binding protein that acts forming the distortion-sensing component of NER by binding tightly with another important NER protein, HR23B, to form a stable XPC-HR23B complex, thus playing a central role in the process of early damage recognition [48,49]. XPC-HR23B complex can recognize a variety of DNA adducts formed by exogenous carcinogens and binds to the DNA damage sites. Therefore, it may play a role in decreasing the toxic effects of such carcinogens and its deficiency may interact with carcinogen exposure [50]. XPC is also involved in DNA damage-induced cell cycle checkpoint regulation and apoptosis, removal of oxidative DNA damage and redox homeostasis [49,51]. XPC rs2228001 (an A-to-C transition in exon 15) leads to a substitution of glutamine for lysine in codon 939 (Lys939Gln) and is located in the domain interacting with the transcription factor IIH (TFIIH) complex [50,52–55], initiating the global genome NER pathway. XPC rs2228001 is one of the most extensively studied NER pathway SNPs, as numerous case-control association studies and meta-analyses have been performed to investigate its potential role on cancer predisposition. In line with our data for DTC, a modest but consistent association of the Gln/Gln homozygous genotype with overall cancer risk is apparent from two of the three meta-analysis that pool data from different cancer types [56–58]. Evidence from these and other cancer site-specific meta-analyses is stronger for lung [53,56–60], bladder [54,56,61,62] and colorectal cancer (CRC) [56,58] [63,64], but also exists for other cancer types such as upper digestive system cancer [65] and hepatocellular carcinoma [50,66].
XPC rs2228001 genotype has also been found to correlate with survival of hepatocellular patients [66], with XPC mRNA expression levels [60,66,67], with drug-induced toxicity in cancer patients treated with platinum-based chemotherapeutic agents (e.g., cisplatin) [68,69], with sensitivity of lung squamous cell carcinoma patients to chemotherapy [67] and to interfere with the capacity to repair DNA lesions induced by, e.g., benzo(a)pyrene [70–72], gamma-radiation [70], X-rays [73], UV radiation [74], aflatoxin B1 [50] and meat-derived carcinogens [75]. Overall, evidence strongly suggests that XPC rs2228001 genotype is associated with altered DNA repair capacity, establishing ground for a putative role of this SNP in cancer susceptibility.

The MSH6 gene (mutS homolog 6) is a member of a set of genes known as the mismatch repair (MMR) genes. MSH6 integrates the MutS\(\alpha\) complex, a sensor of genetic damage that, besides its role in the repair of replication errors, cooperates with other DNA repair and damage-response signalling pathways to allow for cell cycle arrest, DNA repair and/or apoptosis of genetically damaged cells. Several MSH6 mutations have been identified and suggested as causative in Lynch syndrome (LS) patients [76–80]. Despite TC is not part of the usual LS spectrum, the effect of MSH6 in TC susceptibility has previously been explored [81,82]. MSH6 rs1042821 has also been frequently investigated in the context of cancer susceptibility, mostly with inconclusive findings [83–90]. Consistent with our results, MSH6 rs1042821 has previously been associated with increased CRC risk [91–93], highly malignant bladder cancer [94], pancreatic cancer [95] and triple negative breast cancer (TNBC) [96]. On the contrary, the T allele [97] and the CT heterozygous genotype [98] have been associated with decreased colorectal and hepatocellular carcinoma, respectively. The only meta-analysis concerning the role of MSH6 rs1042821 on cancer predisposition that we are aware of is also inconclusive [99]. Despite plausible, a potential role for MSH6 rs1042821 on cancer predisposition (DTC, in particular) remains elusive. Further well-powered studies are needed to clarify this issue.

The role of CCNH rs2230641 on cancer predisposition has only seldom been evaluated: in agreement with our results, a significantly increased bladder cancer risk in ever smokers has been reported for C allele carriers [100] but, on the contrary, such genotype has also been associated with a significantly decreased risk of chronic leukaemia [101]. Most other studies, namely in oesophageal [102], bladder [103], biliary tract [104] and renal cell carcinoma [105], as well as in oral premalignant lesions [106] have been inconclusive. Interestingly, the pharmacogenomic implications of CCNH rs2230641 on the outcome of platinum-based chemotherapy have also been evaluated, results supporting a role for CCNH rs2230641 on the response to DNA damaging agents: the presence of the CCNH rs2230641 variant C allele has been associated with longer survival in NLCSC patients receiving platinum-based chemotherapy [107] and with increased incidence and severity of oxaliplatin-induced acute peripheral neuropathy in digestive tract cancer patients undergoing with the oxaliplatin-based chemotherapy [108]. Similarly, increased risk of severe oxaliplatin-induced acute peripheral neuropathy was observed by Custodio et al. [109] in high-risk stage II and stage III colon cancer patients homozygous for the C allele, submitted to oxaliplatin-based adjuvant chemotherapy. CCNH codes for a highly conserved cyclin protein that participates in several cellular processes such as the NER pathway, cell cycle regulation and receptor phosphorylation, among others [48,110]. Although data on the functional relevance of rs2230641 is lacking, the pleiotropic effects of CCNH confer biological plausibility to our hypothesis that CCNH variants may be involved in cancer susceptibility.

Finally, ERCC5, also known as XPG, is located on chromosome 13q22–q33 [111] and comprises 15 exons [112,113]. It encodes a structure-specific endonuclease that has multiple functions during NER [114], reason why defects in this gene can impair DNA repair resulting in genomic instability and carcinogenesis [115]. In fact, only a few studies have considered the putative contribution of ERCC5 rs2227869 to cancer susceptibility, most being inconclusive. Interestingly, the only significant findings reported thus far are in line with those reported here, suggesting a protective role for the heterozygous genotype: Hussain et al. [116] reported a significant reduction in stomach cancer risk in heterozygous genotype individuals and a similar, despite nonsignificant, trend has also been independently observed for melanoma [117] and for squamous cell carcinoma of the head and neck (SCCHN) [118]. More
importantly, in the only meta-analysis performed to date [119], a decrease in cancer risk in ERCC5 rs2227869 heterozygotes (and for the C allele) has also been reported.

Many of these (and other) SNPs also presented significant findings on stratifying data according to hystotype, gender and age: on histological stratification, significant associations were observed between XRCC3 rs861539, XPC rs2228001, ERCC5 rs2227869, MUTYH rs3219489 and NBN rs1805794 and papillary TC, while MSH6 rs1042821, MLH3 rs175080 and XRCC2 rs3218536 were associated with follicular TC. XRCC3 rs861539, XPC rs2228001, MSH6 rs1042821, CCNH rs2230641, ERCC5 rs2227869 and ERCC5 rs17655 were associated with DTC in the female subset while no association was observed in males. Finally, XPC rs2228001 and XRCC5 rs2440 were associated with DTC in participants younger than 50 years, while, in participants aged 50 or more years, the DTC-associated SNPs included XRCC3 rs861539, CCNH rs2230641, ERCC5 rs2228529 and RAD51 rs1801321.

It is unclear whether these findings (and which among these) truly represent group-specific effects or whether they simply reflect the overall effect on the largest groups (i.e., when group sizes are unbalanced, e.g., papillary TC vs follicular TC, female vs male) and the corresponding lack of power to detect an effect on the smallest groups. Also, due to the low sample size on each strata, some of these results may simply represent incident findings (type I errors). XRCC3 rs861539, for example, has been previously associated with papillary TC [22,39,40]—in line with our results—but not with follicular TC. An effect of XRCC3 rs861539 genotype in follicular TC cannot, however, be excluded since follicular TC is much less frequent than papillary TC and these studies may have been underpowered to detect such effect. Also, Su et al. [120] have demonstrated the homozygous genotype of this SNP to be associated with breast cancer; the association being stronger in women younger than 55 years, with earlier first menarche or with latter menopause. This suggests an oestrogen-potentiated genetic effect, compatible with our own observation of increased DTC risk in XRCC3 rs861539 TT homozygotes among females but not among males. Further, the involvement of CCNH, through a cyclin-activated kinase complex, in oestrogen receptor phosphorylation [48] provides a possible rationale for our own observation of an association of the CCNH rs2230641 genotype with DTC among females but not among males. Finally, the association of MSH6 rs1042821 with DTC, observed in this study for female but not male individuals, is compatible with the growing evidence placing DTC as an oestrogen-associated cancer [121–124] and implicating MSH6 in such cancers [78,125–129].

These selected examples highlight the plausibility of the existence of group-specific genetic effects. Overall, such hystotype, gender and age specifies in DTC susceptibility are likely since (1) papillary and follicular TC represent distinct entities, with hystotype-specific molecular profiles (e.g., BRAF mutations and RET/PTC rearrangements in PTC, RAS mutations and PAX8/PPARY translocations in FTC) [130]; (2) important gender differences exist in the incidence of DTC (i.e., DTC is, as previously stated two to four times more frequent in women than in men) [1,2]; and (3) DTC presents some age specificities, uncommon in other types of cancer (DTC is one of the most common malignancies in adolescent and young adults, the median age at diagnosis being lower than that for most other types of cancer) [1,2]. Further well-powered studies are urgently needed to clarify these results and thus establish which of these SNPs, if any, represents true group-specific susceptibility biomarkers.

Considering the multifactorial nature of DTC aetiology and the probable involvement of multiple genetic factors, alone or in combination, in DTC susceptibility, we undertook a combined genotype analyses to investigate the joint effect of multiple SNPs on DTC risk. When combining all risk genotypes significant at single SNP analysis into a unique unbalanced risk score, a clear-cut gene-dosage effect between the number of risk genotypes (unbalanced risk score) and DTC risk was observed, both on global analysis (considering all DTC cases and corresponding controls) and after stratification according to histological, gender and age criteria. This is biologically plausible since the different DNA repair proteins physically and functionally interact with each other, within the same or different DNA repair pathways, establishing ground for additive or even multiplicative effects of different SNPs on DNA repair activity and, hence, cancer risk. Such polygenic approach to assess the cumulative effects
of multiple genetic variants on cancer risk has previously been employed [27,107,131,132], supporting its usefulness and clinical potential.

To investigate the effect of specific DNA repair SNP combinations on DTC risk, all possible $2 \times 2$ combinations were tested on paired SNP analysis, yielding fifteen SNP pairs with $p < 0.01$. Multiple interactions between SNPs from different DNA repair pathways and, even, other DNA damage response proteins have previously been reported [39,42,66,87], providing a rationale for such approach. Of notice, CCNH rs2230641 was the most frequently represented DNA repair SNP in such significant combinations, both at 0.01 and 0.05 significance levels, a finding that is compatible with the pleiotropic role of CCNH in DNA damage repair, cell cycle regulation and receptor phosphorylation [48,110]. More importantly, the contribution of MMR variants to the joint effect of DNA repair SNPs on DTC risk is evident from our results, as they were present in 9 of the 15 SNP pairs presenting significant findings. Besides its critical role in post-replication repair (through recognition and repair of base-base mispairs and insertion/deletion loops that arise during replication), the MMR pathway cooperates with other repair pathways in the recognition and subsequent repair of DNA damage induced by IR, UV light, oxidative stress or genotoxic chemicals (e.g., oxidative lesions, double strand breaks, pyrimidine dimers and inter-strand crosslinks) and contributes to damage-induced cytotoxicity through downstream signalling for cell cycle arrest and apoptosis [133–135]. Therefore, considering the large spectre of action of the MMR pathway, an elevated number of interactions between MMR and other DNA repair SNPs is expected. Such hypothesis, in line with our findings, has been recently strengthened by a report [136] associating SNPs from different DNA repair pathways with CRC in Lynch syndrome patients, a cancer predisposition condition originated by germline MMR mutations. Finally, among SNP pairs presenting significant findings in this study, three are intra-pathway combinations involving either HR or MMR pathway SNPs. The joint effects of MLH3 rs175080 – MSH6 rs1042821 and MSH4 rs5745549 – MSH6 rs1042821 (MMR pathway) SNP combinations were reported and discussed in our original study [18]. The joint effect of RAD51 rs1801321 and XRCC3 rs861539 (HR pathway) on cancer risk has been previously reported for breast cancer [137], in line with our results, and may be of particular relevance for DTC since the formation of radiation damage-induced RAD51 foci requires functional XRCC3 [138].

Finally, on applying haplotype analysis to SNPs that are located in the same chromosome arm (thus likely to segregate together), one block of DNA repair SNPs located on chromosome 5q (comprising CCNH rs2230641, CDK7 rs2972388, MSH3 rs26279, MSH3 rs184967, XRCC4 rs1805377 and XRCC4 rs28360135) was associated with DTC risk in our study. Such results further suggest an independent or interactive effect of these SNPs on DTC predisposition.

Overall, our results suggest that DNA repair SNPs across different pathways and may contribute to DTC predisposition, possibly exerting cumulative effects. This is of relevance since the estimated high heritability of DTC is only partially explained, even when considering the contribution of several GWAS recently performed. Gene-gene and gene-environment interactions have been hypothesised to play an important role so their identification and in-depth study is highly desirable to explain the “missing” heritability of DTC. However, the results presented here should be regarded only as proof of concept and must therefore be validated through replication in larger independent populations. Future studies should also be designed with the intention of accounting for environmental factors such as IR exposure and iodine deficiency (and their potential interaction with genetic factors). In addition, they should be sufficiently powered to allow other, less frequent but potentially relevant SNPs, to be studied and to allow more sophisticated and conclusive gene-gene interaction analysis to be performed. Finally, in order to strengthen our preliminary findings, the functional significance of these SNPs should be further investigated as well as their potential association with mutational events involved in DTC carcinogenesis (e.g., BRAF mutations and RET/PTC rearrangements).

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restrict collaboration by O.M.G., T.C.F. and A.P.A.; Data Curation, O.M.G., T.C.F. and E.L.; Writing—Original Draft Preparation, L.S.S.; Writing – Review & Editing, B.C.G., H.N.B., O.M.G., A.P.A., S.N.S., J.R.; Visualization has been prepared by L.S.S., and S.N.S.; Supervision of this project, J.R.; Project Administration, J.R. and E.L.; Funding Acquisition, J.R.

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**References**

1. Ferlay, J.; Ervik, M.; Lam, F.; Colombet, M.; Mery, L.; Piñeros, M.; Znaor, A.; Soerjomataram, I.; Bray, F. Global Cancer Observatory: Cancer Today. International Agency for Research on Cancer, 2018. Available online: https://gco.iarc.fr/today (accessed on 28 May 2019).

2. Kitahara, C.M.; Sosa, J.A. The changing incidence of thyroid cancer. *Nat. Rev. Endocrinol.* **2016**, 12, 646. [CrossRef] [PubMed]

3. Lloyd, R.V.; Osamura, R.Y.; Klöppel, G.N.; Rosai, J. *WHO Classification of Tumours of Endocrine Organs*, 4th ed.; International Agency for Research on Cancer (IARC) Press: Lyon, France, 2017; Volume 10, pp. 65–144.

4. Khosravi, M.H.; Kouhi, A.; Saeedi, M.; Bagheriagh, A.; Amirzade-Iranjaq, M.H. Thyroid Cancers: Considerations, Classifications, and Managements. In *Diagnosis and Management of Head and Neck Cancer*; Akarslan, Z., Ed.; IntechOpen: London, UK, 2017; pp. 57–82. [CrossRef]

5. Marcello, M.A.; Malandrino, P.; Almeida, J.F.M.; Martins, M.B.; Cunha, L.L.; Bufalo, N.E.; Pellegriti, G.; Ward, L.S. The influence of the environment on the development of thyroid tumors: A new appraisal. *Endocr. Relat. Cancer* **2014**, 21, T235–T254. [CrossRef] [PubMed]

6. Saenko, V.A.; Rogounovitch, T.I. Genetic Polymorphism Predisposing to Differentiated Thyroid Cancer: A Review of Major Findings of the Genome-Wide Association Studies. *Endocrinol. Metab.* **2018**, 33, 164–174. [CrossRef]

7. Hwangbo, Y.; Park, Y.J. Genome-Wide Association Studies of Autoimmune Thyroid Diseases, Thyroid Function, and Thyroid Cancer. *Endocrinol. Metab.* **2018**, 33, 175–184. [CrossRef] [PubMed]

8. Adjadj, E.; Schlumberger, M.; de Vathaire, F. Germ-line DNA polymorphisms and susceptibility to differentiated thyroid cancer. *Lancet. Oncol.* **2009**, 10, 181–190. [CrossRef]

9. Iñigo, L.; Mercedes, R. Association studies in thyroid cancer susceptibility: Are we on the right track? *J. Mol. Endocrinol.* **2011**, 47, R43–R58. [CrossRef]

10. Figlioli, G.; Elisei, R.; Romei, C.; Melaiu, O.; Cippolini, M.; Bambi, F.; Chen, B.; Köhler, A.; Cristaudo, A.; Hemminki, K.; et al. A Comprehensive Meta-analysis of Case–Control Association Studies to Evaluate Polymorphisms Associated with the Risk of Differentiated Thyroid Carcinoma. *Cancer Epidemiol. Biomark. Prev.* **2016**, 25, 700–713. [CrossRef] [PubMed]

11. Chatterjee, N.; Walker, G.C. Mechanisms of DNA damage, repair, and mutagenesis. *Environ. Mol. Mutagen.* **2017**, 58, 235–263. [CrossRef]

12. Gatzidou, E.; Michailidi, C.; Tseleni-Balafouta, S.; Theocharis, S. An epitome of DNA repair related genes and mechanisms in thyroid carcinoma. *Cancer Lett.* **2010**, 290, 139–147. [CrossRef]

13. Köberle, B.; Koch, B.; Fischer, B.M.; Hartwig, A. Single nucleotide polymorphisms in DNA repair genes and putative cancer risk. *Arch. Toxicol.* **2016**, 90, 2369–2388. [CrossRef]

14. Bastos, H.N.; Antão, M.R.; Silva, S.N.; Azevedo, A.P.; Manita, I.; Teixeira, V.; Pina, J.E.; Gil, O.M.; Ferreira, T.C.; Limbert, E.; et al. Association of Polymorphisms in Genes of the Homologous Recombination DNA Repair Pathway and Thyroid Cancer Risk. *Thyroid* **2009**, 19, 1067–1075. [CrossRef]

15. Gomes, B.C.; Silva, S.N.; Azevedo, A.P.; Manita, I.; Gil, O.M.; Ferreira, T.C.; Limbert, E.; Rueff, J.; Gaspar, J.F. The role of common variants of non-homologous end-joining repair genes XRCC4, LIG4 and Ku80 in thyroid cancer risk. *Oncof. Rep.* **2010**, 24, 1079–1085.
16. Santos, L.S.; Branco, S.C.; Silva, S.N.; Azevedo, A.P.; Gil, O.M.; Manita, I.; Ferreira, T.C.; Limbert, E.; Rueff, J.; Gaspar, J.F. Polymorphisms in base excision repair genes and thyroid cancer risk. *Oncol. Rep.* 2012, 28, 1859–1868. [CrossRef]

17. Santos, L.S.; Gomes, B.C.; Gouveia, R.; Silva, S.N.; Azevedo, A.P.; Camacho, V.; Manita, I.; Gil, O.M.; Ferreira, T.C.; Limbert, E.; et al. The role of CCNH Val270Ala (rs2230641) and other nucleotide excision repair polymorphisms in individual susceptibility to well-differentiated thyroid cancer. *Oncol. Rep.* 2013, 30, 2458–2466. [CrossRef]

18. Bashir, K.; Sarwar, R.; Fatima, S.; Saeed, S.; Mahjabeen, I.; Akhtar Kayani, M. Haplotype analysis of XRCCI gene polymorphisms and the risk of thyroid carcinoma. *J. BUON* 2018, 23, 234–243.

19. Bashir, K.; Sarwar, R.; Saeed, S.; Mahjabeen, I.; Kayani, M.A. Interaction among susceptibility genotypes of PARP1 SNPs in thyroid carcinoma. *PLoS ONE* 2018, 13, e0199007. [CrossRef]

20. Sandler, J.E.; Huang, H.; Zhao, N.; Wu, W.; Liu, F.; Ma, S.; Udelsman, R.; Zhang, Y. Germline Variants in DNA Repair Genes, Diagnostic Radiation, and Risk of Thyroid Cancer. *Cancer Epidemiol. Biomark. Prev.* 2018, 27, 285–294. [CrossRef]

21. Santos, L.S.; Silva, S.N.; Gil, O.M.; Ferreira, T.C.; Limbert, E.; Rueff, J. Mismatch repair single nucleotide polymorphisms and thyroid cancer susceptibility. *Oncol. Lett.* 2015, 15, 6715–6726. [CrossRef]

22. Sarwar, R.; Mahjabeen, I.; Bashir, K.; Saeed, S.; Kayani, M.A. Haplotype Based Analysis of XRCC3 Gene Polymorphisms in Thyroid Cancer. *Cell Physiol. Biochem.* 2017, 42, 22–33. [CrossRef]

23. Liyanarachchi, S.; Wojcicka, A.; Li, W.; Czetwertynska, M.; Stachlewska, E.; Nagy, R.; Hoag, K.; Wen, B.; Ruiz-Llorente, S.; Schiavi, F.; Marcos, R.; et al. An Epistatic Interaction between the PAX8 and STK17B Genes in Papillary Thyroid Cancer Susceptibility. *Oncol. Lett.* 2015, 8, 758. [CrossRef]

24. Isakova, J.; Talaibekova, E.; Aldasheva, N.; Vinnikov, D.; Aldashev, A. The association of polymorphic markers Arg399Gln of XRCCI gene, Arg72Pro of TP53 gene and T309G of MDM2 gene with breast cancer in Kyrgyz females. *BMC Cancer* 2017, 17, 758. [CrossRef]

25. Mordukhovich, I.; Beyea, J.; Herring, A.H.; Hatch, M.; Stellman, S.D.; Teitellbaum, S.L.; Richardson, D.B.; Millikan, R.C.; Engel, L.S.; Shantakumar, S.; et al. Polymorphisms in DNA repair genes, traffic-related polycyclic aromatic hydrocarbon exposure and breast cancer incidence. *Int. J. Cancer* 2016, 139, 310–321. [CrossRef]

26. Solé, X.; Guinó, E.; Valls, J.; Iniesta, R.; Moreno, V. SNPStats: A web tool for the analysis of association studies. *Bioinformatics* 2006, 22, 1928–1929. [CrossRef]

27. Shakeri, M.; Zakeri, F.; Changizi, V.; Rajajbopr, M.R.; Farshidpour, M.R. Cytogenetic effects of radiation and genetic polymorphisms of the XRCCI1 and XRCC3 repair genes in industrial radiographers. *Radiat. Environ. Biophys.* 2019, 58, 247–255. [CrossRef]

28. Song, Y.-Z.; Han, F.-J.; Liu, M.; Xia, C.-C.; Shi, W.-Y.; Dong, L.-H. Association between Single Nucleotide Polymorphisms in XRCC3 and Radiation-Induced Adverse Effects on Normal Tissue: A Meta-Analysis. *PLoS ONE* 2015, 10, e0130388. [CrossRef]
34. Chai, F.; Liang, Y.; Chen, L.; Zhang, F.; Jiang, J. Association between XRCC3 Thr241Met Polymorphism and Risk of Breast Cancer: Meta-Analysis of 23 Case-Control Studies. *Med. Sci. Monit.* **2015**, *21*, 3231–3240. [CrossRef]

35. Zhang, B.; Beeghly-Fadiel, A.; Long, J.; Zheng, W. Genetic variants associated with breast-cancer risk: Comprehensive research synthesis, meta-analysis, and epidemiological evidence. *Lancet Oncol.* **2011**, *12*, 477–488. [CrossRef]

36. He, X.-F.; Wei, W.; Li, J.-L.; Shen, X.-L.; Ding, D.-P.; Wang, S.-L.; Liu, Z.-Z.; Qin, J.-B.; Wu, L.-X.; Xie, D.-L. Association between the XRCC3 T241M polymorphism and risk of cancer: Evidence from 157 case–control studies. *Gene* **2013**, *523*, 10–19. [CrossRef]

37. Peng, Q.; Mo, C.; Tang, W.; Chen, Z.; Li, R.; Zhai, L.; Yang, S.; Wu, J.; Sui, J.; Li, S.; et al. DNA repair gene XRCC3 polymorphisms and bladder cancer risk: A meta-analysis. *Tumor Biol.* **2014**, *35*, 1933–1944. [CrossRef]

38. Ma, Q.; Zhao, Y.; Wang, S.; Zhang, X.; Zhang, J.; Du, M.; Li, L.; Zhang, Y. Genetic polymorphisms of XRCC3 Thr241Met (C18067T, rs861539) and bladder cancer risk: A meta-analysis of 18 research studies. *Tumor Biol.* **2014**, *35*, 1473–1480. [CrossRef]

39. Yan, L.; Li, Q.; Li, X.; Ji, H.; Zhang, L. Association Studies between XRCC1, XRCC2, XRCC3 Polymorphisms and Differentiated Thyroid Carcinoma. *Cell Physiol. Biochem.* **2016**, *38*, 1075–1084. [CrossRef]

40. Yuan, K.; Huo, M.; Sun, Y.; Wu, H.; Chen, H.; Wang, Y.; Fu, R. Association between x-ray repair cross-complementing group 3 (XRCC3) genetic polymorphisms and papillary thyroid cancer susceptibility in a Chinese Han population. *Tumor Biol.* **2016**, *37*, 997–987. [CrossRef]

41. Wang, X.; Zhang, K.; Liu, X.; Liu, B.; Wang, Z. Association between XRCC1 and XRCC3 gene polymorphisms and risk of thyroid cancer. *Int. J. Clin. Exp. Pathol.* **2015**, *8*, 3160–3167.

42. Fayaz, S.; Karimmirza, M.; Tanhaei, S.; Fathi, M.; Torbati, P.M.; Fard-Esfahani, P. Increased risk of differentiated thyroid carcinoma with combined effects of homologous recombination repair gene polymorphisms in an Iranian population. *Asian Pac. J. Cancer Prev.* **2014**, *14*, 6727–6731. [CrossRef]

43. Sturgis, E.M.; Zhao, C.; Zheng, R.; Wei, Q. Radiation Response Genotype and Risk of Differentiated Thyroid Cancer: A Case-Control Analysis. *Laryngoscope* **2005**, *115*, 938–945. [CrossRef]

44. Lu, W.; Wu, G.; Zhang, B. Association Between X-Ray Cross-complementing Group 3 (XRCC3) Thr241Met Polymorphism and Risk of Thyroid Cancer: A Meta-Analysis. *Med. Sci. Monit.* **2015**, *21*, 3978–3985. [CrossRef]

45. Yu, X.-L.; Liu, H.; Wang, B.; Fu, Z.-J.; Yuan, Y.; Yan, S.-L.; Zhao, W.-J.; Wang, Y.-G.; Cai, J. Significant associations between X-ray repair cross-complementing group 3 genetic polymorphisms and thyroid cancer risk. *Tumor Biol.* **2014**, *35*, 2009–2015. [CrossRef]

46. Natalia, M.A.; Vladimir, A.S.; Tatiana, I.R.; Valentina, M.D.; Eugeny, F.L.; Victor, K.I.; Norisato, M.; Ryo, K.; Shunichi, Y. Polymorphisms of DNA damage response genes in radiation-related and sporadic papillary thyroid carcinoma. *Endocr. Relat. Cancer* **2009**, *16*, 491–503. [CrossRef]

47. Siraj, A.K.; Al-Rasheed, M.; Ibrahim, M.; Siddiqui, K.; Al-Dayel, F.; Al-Sanea, O.; Uddin, S.; Al-Kuraya, K. RAD52 polymorphisms contribute to the development of papillary thyroid cancer susceptibility in Middle Eastern population. *J. Endocrinol. Investig.* **2008**, *31*, 893–899. [CrossRef]

48. Nouspikel, T. DNA Repair in Mammalian Cells. *Cell Mol. Life Sci.* **2009**, *66*, 994–1009. [CrossRef]

49. Melis, J.P.M.; Luijten, M.; Mullenders, L.H.F.; van Steeg, H. The role of XPC: Implications in cancer and oxidative DNA damage. *Mutat. Res. Rev. Mutat.* **2011**, *728*, 107–117. [CrossRef]

50. Yao, J.-G.; Huang, X.-Y.; Long, X.-D. Interaction of DNA repair gene polymorphisms and aflatoxin B1 in the risk of hepatocellular carcinoma. *Int. J. Clin. Exp. Pathol.* **2014**, *7*, 6231–6244.

51. Chen, Z.; Yang, J.; Wang, G.; Song, B.; Li, J.; Xu, Z. Attenuated Expression of Xeroderma Pigmentosum Group C Is Associated with Critical Events in Human Bladder Cancer Carcinogenesis and Progression. *Cancer Res.* **2007**, *67*, 4578–4585. [CrossRef]

52. Park, J.Y.; Huang, Y.; Sellers, T.A. Single Nucleotide Polymorphisms in DNA Repair Genes and Prostate Cancer Risk. *In Cancer Epidemiology. Methods in Molecular Biology; Verma, M., Ed.; Humana Press: Totowa, NJ, USA, 2009; Volume 471, pp. 361–385. [CrossRef]

53. Mei, C.-R.; Luo, M.; Li, H.-M.; Deng, W.-J.; Zhou, Q.-H. DNA repair gene polymorphisms in the nucleotide excision repair pathway and lung cancer risk: A meta-analysis. *Chin. J. Cancer Res.* **2011**, *23*, 79. [CrossRef]
54. Zhang, Y.; Wang, X.; Zhang, W.; Gong, S. An association between XPC Lys939Gln polymorphism and the risk of bladder cancer: A meta-analysis. *Tumor Biol.* 2013, 34, 973–982. [CrossRef]

55. Hua, R.-X.; Zhu, J.; Jiang, D.-H.; Zhang, S.-D.; Zhang, J.-B.; Xue, W.-Q.; Li, X.-Z.; Zhang, P.-F.; He, J.; Jia, W.-H. Association of XPC Gene Polymorphisms with Colorectal Cancer Risk in a Southern Chinese Population: A Case-Control Study and Meta-Analysis. *Genes* 2016, 7, 73. [CrossRef]

56. He, J.; Shi, T.-Y.; Zhu, M.-L.; Wang, M.-Y.; Li, Q.-X.; Wei, Q.-Y. Associations of Lys939Gln and Ala499Val polymorphisms of the XPC gene with cancer susceptibility: A meta-analysis. *Int. J. Cancer* 2013, 133, 1765–1775. [CrossRef]

57. Qiu, L.; Wang, Z.; Shi, X.; Wang, H.; Han, B. Association of XPC Lys939Gln polymorphism and the risk of cancers: A meta-analysis. *Eur. J. Cancer* 2008, 44, 2241–2253. [CrossRef]

58. Francisco, G.; Ménezes, P.R.; Eluf-Neto, J.; Chammas, R. XPC polymorphisms play a role in tissue-specific carcinogenesis: A meta-analysis. *Eur. J. Hum. Genet.* 2008, 16, 724. [CrossRef]

59. Jin, B.; Dong, Y.; Zhang, X.; Wang, H.; Han, B. Association of XPC and Lung Cancer Risk: A Meta-Analysis. *PLoS ONE* 2014, 9, e93937. [CrossRef]

60. Zhu, M.-L.; Hua, R.-X.; Zheng, L. Associations between polymorphisms of the XPC gene and lung cancer susceptibility: A meta-analysis. *Tumor Biol.* 2014, 35, 2931–2939. [CrossRef]

61. Dai, Q.-S.; Hua, R.-X.; Zeng, R.-F.; Long, J.-T.; Peng, Z.-W. XPC gene polymorphisms contribute to bladder cancer susceptibility: A meta-analysis. *Tumor Biol.* 2014, 35, 447–453. [CrossRef]

62. Dou, K.; Xu, Q.; Han, X. The association between XPC Lys939Gln gene polymorphism and urinary bladder cancer susceptibility: A systematic review and meta-analysis. *Diagn. Pathol.* 2013, 8, 112. [CrossRef]

63. Ma, X.; Zhang, B.; Zheng, W. Genetic variants associated with colorectal cancer risk: Comprehensive research synopsis, meta-analysis, and epidemiological evidence. *Gut* 2014, 63, 326–336. [CrossRef]

64. Peng, Q.; Lao, X.; Tang, W.; Chen, Z.; Li, R.; Qin, X.; Li, S. XPC Lys939Gln polymorphism contributes to colorectal cancer susceptibility: Evidence from a meta-analysis. *Diagn. Pathol.* 2014, 9, 120. [CrossRef]

65. Jiang, X.; Zhou, L.-T.; Zhang, S.-C.; Chen, K. XPC Polymorphism Increases Risk of Digestive System Cancers: Current Evidence from A Meta-Analysis. *Chin. J. Cancer Res.* 2012, 24, 181–189. [CrossRef]

66. Long, X.-D.; Ma, Y.; Zhou, Y.-F.; Ma, A.-M.; Fu, G.-H. Polymorphism in xeroderma pigmentosum complementation group C codon 939 and aflatoxin B1-related hepatocellular carcinoma in the Guangxi population. *Hepatology* 2010, 52, 1301–1309. [CrossRef]

67. Wang, C.; Nie, H.; Li, Y.; Liu, G.; Wang, X.; Xing, S.; Zhang, L.; Chen, X.; Chen, Y.; Li, Y. The study of the relation of DNA repair pathway genes SNPs and the sensitivity to radiotherapy and chemotherapy of NSCLC. *Sci. Rep.* 2016, 6, 26256. [CrossRef]

68. Caronia, D.; Patiño-García, A.; Milné, R.L.; Zalacain-Díez, M.; Pita, G.; Alonso, M.R.; Moreno, L.T.; Sierrasesumaga-Ariznabarreta, L.; Benítez, J.; González-Neira, A. Common variations in ERCC2 are associated with response to cisplatin chemotherapy and clinical outcome in osteosarcoma patients. *Pharmacogenom. J.* 2009, 9, 347. [CrossRef]

69. Sakano, S.; Hinoda, Y.; Sasaki, M.; Wada, T.; Matsumoto, H.; Eguchi, S.; Shinohara, A.; Kawai, Y.; Hara, T.; Nagao, K.; et al. Nucleotide excision repair gene polymorphisms may predict acute toxicity in patients treated with chemoradiotherapy for bladder cancer. *Pharmacogenomics* 2010, 11, 1377–1387. [CrossRef]

70. Zhu, Y.; Yang, H.; Chen, Q.; Lin, J.; Grossman, H.B.; Dinney, C.P.; Wu, X.; Gu, J. Modulation of DNA damage/DNA repair capacity by XPC polymorphisms. *DNA Repair* 2008, 7, 141–148. [CrossRef]

71. Pavanello, S.; Pulliero, A.; Siwinska, E.; Mielzynska, D.; Clonfero, E. Reduced nucleotide excision repair and GSTM1-null genotypes influence anti-B[a]PDE–DNA adduct levels in mononuclear white blood cells of highly PAH-exposed coke oven workers. *Carcinogenesis* 2005, 26, 169–175. [CrossRef]

72. Lin, J.; Swan, G.E.; Shields, P.G.; Benowitz, N.L.; Gu, J.; Amos, C.I.; de Andrade, M.; Spitz, M.R.; Wu, X. Mutagen Sensitivity and Genetic Variants in Nucleotide Excision Repair Pathway: Genotype-Phenotype Correlation. *Cancer Epidemiol. Biomark. Prev.* 2007, 16, 2065–2071. [CrossRef]

73. Cornetta, T.; Festa, F.; Testa, A.; Cozzi, R. DNA damage repair and genetic polymorphisms: Assessment of individual sensitivity and repair capacity. *Int. J. Radiat. Oncol.* 2006, 66, 537–545. [CrossRef]

74. Qiao, Y.; Spitz, M.R.; Shen, H.; Guo, Z.; Shete, S.; Hedayati, M.; Grossman, L.; Mohrenweiser, H.; Wei, Q. Modulation of repair of ultraviolet damage in the host-cell reactivation assay by polymorphic XPC and XPD/ERCC2 genotypes. *Carcinogenesis* 2002, 23, 295–299. [CrossRef]
Genes 2019, 10, 586

75. Ho, V.; Brunetti, V.; Peacock, S.; Massey, T.E.; Godshalk, R.W.L.; van Schooten, F.J.; Ashbury, J.E.; Vanner, S.J.; King, W.D. Exposure to meat-derived carcinogens and bulky DNA adduct levels in normal-appearing colon mucosa. Mutat. Res. Genet. Toxicol. Environ. Mutagen. 2017, 821, 5–12. [CrossRef]

76. Broaddus, R.R.; Lynch, P.M.; Lu, K.H.; Luthra, R.; Michelson, S.J. Unusual tumors associated with the hereditary nonpolyposis colorectal cancer syndrome. Mod. Pathol. 2004, 17, 981–989. [CrossRef]

77. Pande, M.; Wei, C.; Chen, J.; Amos, C.I.; Lynch, P.M.; Lu, K.H.; Lucio, L.A.; Boyd-Rogers, S.G.; Bannon, S.A.; Mork, M.E.; et al. Cancer spectrum in DNA mismatch repair gene mutation carriers: Results from a hospital based Lynch syndrome registry. Fam. Cancer 2012, 11, 441–447. [CrossRef]

78. Stulp, R.P.; Herkert, J.C.; Karrenbeld, A.; Mol, B.; Vos, Y.J.; Sijmons, R.H. Thyroid cancer in a patient with a germline MSH2 mutation. Case report and review of the Lynch syndrome expanding tumour spectrum. Hered. Cancer Clin. Pract. 2008, 6, 15–21. [CrossRef]

79. Pelizzo, M.R.; Pennelli, G.; Zane, M.; Galuppini, F.; Colletti, P.M.; Merante Boschin, I.; Rubello, D. Papillary polyps: hMLH1 and CpG island methylator phenotype in colon cancer. Mol. Carcinog. 2009, 48, 989–994. [CrossRef]

80. Johnson, J.M.; Chen, J.; Ali, S.M.; Dardi, I.K.; Tuluc, M.; Cognetti, D.; Campling, B.; Sama, A.R. Molecular Profiling of Synchronous Colon Cancers and Anaplastic Thyroid Cancer in a Patient with Lynch Syndrome. J. Gastrointest. Cancer 2018, 49, 203–206. [CrossRef]

81. Kunstman, J.W.; Juhrin, C.C.; Goh, G.; Brown, T.C.; Stenman, A.; Healy, J.M.; Rubinstein, J.C.; Choi, M.; Kiss, N.; Nelson-Williams, C.; et al. Characterization of the mutational landscape of anaplastic thyroid cancer via whole-exome sequencing. Hum. Mol. Genet. 2015, 24, 2318–2329. [CrossRef]

82. Yu, Y.; Dong, L.; Li, D.; Chuat, S.; Wu, Z.; Zheng, X.; Cheng, Y.; Han, L.; Yu, J.; Gao, M. Targeted DNA Sequencing Detects Mutations Related to Susceptibility among Familial Non-medullary Thyroid Cancer. Sci. Rep. 2015, 5, 16129. [CrossRef]

83. Picelli, S.; Zajac, P.; Zhou, X.-L.; Edler, D.; Lenander, C.; Dalén, J.; Hjern, F.; Lundqvist, N.; Lindforss, U.; Pålman, L.; et al. Common variants in human CRC genes as low-risk alleles. Eur. J. Cancer 2010, 46, 1041–1048. [CrossRef]

84. Giráldez, M.D.; Balaguer, F.; Caldés, T.; Sanchez-de-Abajo, A.; Gómez-Fernández, N.; Ruiz-Ponte, C.; Muñoz, J.; Garre, P.; Gonzalo, V.; Moreira, L.; et al. Association of MUTYH and MSH6 germline mutations in colorectal cancer patients. Fam. Cancer 2009, 8, 525. [CrossRef]

85. Conde, J.; Silva, S.N.; Azevedo, A.P.; Teixeira, V.; Pina, J.E.; Rueff, J.; Gaspar, J.F. Association of common variants in mismatch repair genes and breast cancer susceptibility: A multigene study. BMC Cancer 2009, 9, 344. [CrossRef]

86. Curtin, K.; Samowitz, W.S.; Wolff, R.K.; Caan, B.J.; Ulrich, C.M.; Potter, J.D.; Slattery, M.L. MSH6 G39E polymorphism and CpG island methylator phenotype in colon cancer. Mol. Carcinog. 2009, 48, 989–994. [CrossRef]

87. Hirata, H.; Hinoda, Y.; Kawamoto, K.; Kikuno, N.; Suehiro, Y.; Okayama, N.; Tanaka, Y.; Daihya, R. Mismatch Repair Gene MSH3 Polymorphism is Associated with the Risk of Sporadic Prostate Cancer. J. Urol. 2008, 179, 2020–2024. [CrossRef]

88. Smith, T.R.; Levine, E.A.; Freimanis, R.I.; Akman, S.A.; Allen, G.O.; Hoang, K.N.; Liu-Mares, W.; Hu, J.J. Polygenic model of DNA repair genetic polymorphisms in human breast cancer risk. Carcinogenesis 2008, 29, 2132–2138. [CrossRef]

89. Landi, S.; Gemignani, F.; Canzian, F.; Gaboreau, V.; Barale, R.; Landi, D.; Szaszienia-Dabrowska, N.; Zaridze, D.; Lissowska, J.; Rudnai, P.; et al. DNA Repair and Cell Cycle Control Genes and the Risk of Young-Onset Lung Cancer. Cancer Res. 2006, 66, 11062–11069. [CrossRef]

90. Yu, J.H.; Bgler, J.; Whitton, J.; Potter, J.D.; Ulrich, C.M. Mismatch repair polymorphisms and colorectal polyps: hMLH1-93G>A variant modifies risk associated with smoking. Am. J. Gastroenterol. 2006, 101, 1313–1319. [CrossRef]

91. Zelga, P.; Przybyłow ska-Sygut, K.; Zelga, M.; Dziki, A.; Majsterek, I. The 116G > A MSH6 and IVS1-1121C > T PMS2 Genes Polymorphisms Modulate the Risk of the Sporadic Colorectal Cancer Development in Polish Population. Pathol. Oncol. Res. 2018, 24, 231–235. [CrossRef]

92. Berndt, S.I.; Platz, E.A.; Fallin, M.D.; Thuita, L.W.; Hoffman, S.C.; Helzlsouer, K.J. Mismatch repair polymorphisms and the risk of colorectal cancer. Int. J. Cancer 2007, 120, 1548–1554. [CrossRef]
93. Campbell, P.T.; Curtin, K.; Ulrich, C.M.; Samowitz, W.S.; Bigler, J.; Velicer, C.M.; Caan, B.; Potter, J.D.; Slattery, M.L. Mismatch repair polymorphisms and risk of colon cancer, tumour microsatellite instability and interactions with lifestyle factors. Gut 2009, 58, 661–667. [CrossRef]

94. Sanyal, S.; De Verdier, P.J.; Steinbeck, G.; Larsson, P.; Onelov, E.; Hemminki, K.; Kumar, R. Polymorphisms in XPD, XPC and the risk of death in patients with urinary bladder neoplasms. Acta Oncol. 2007, 46, 31–41. [CrossRef]

95. Dong, X.; Li, Y.; Chang, P.; Hess, K.R.; Abbruzzese, J.L.; Li, D. DNA mismatch repair network gene polymorphism as a susceptibility factor for pancreatic cancer. Mol. Carcinog. 2012, 51, 491–499. [CrossRef]

96. Lee, E.; Levine, E.A.; Franco, V.I.; Allen, G.O.; Gong, F.; Zhang, Y.; Hu, J.J. Combined Genetic and Nutritional Risk Models of Triple Negative Breast Cancer. Nutr. Cancer 2014, 66, 955–963. [CrossRef]

97. Tulupova, E.; Kumar, R.; Hanova, M.; Slyskaova, J.; Pardini, B.; Polakova, V.; Naccarati, A.; Vodickova, L.; Novotny, J.; Halamkova, J.; et al. Do polymorphisms and haplotypes of mismatch repair genes modulate risk of sporadic colorectal cancer? Mutat. Res. 2008, 648, 40–45. [CrossRef]

98. Liu, Y.; Zhang, X.; Jia, J.; Tang, L.; Gao, X.; Yan, L.; Wang, L.; Yu, F.; Ma, N.; Liu, W.; et al. Correlation between polymorphisms in DNA mismatch repair genes and the risk of primary hepatocellular carcinoma for the Han population in northern China. Scand. J. Gastroenterol. 2015, 50, 1404–1410. [CrossRef]

99. Li, Z.; Kong, L.; Yu, L.; Huang, J.; Wang, K.; Chen, S.; Yu, M.; Wei, S. Association between MSH6 G39E polymorphism and cancer susceptibility: A meta-analysis of 7,046 cases and 34,554 controls. Tumour Biol. 2014, 35, 6029–6037. [CrossRef]

100. Chen, M.; Kamat, A.M.; Huang, M.; Grossman, H.B.; Dinney, C.P.; Lerner, S.P.; Wu, X.; Gu, J. High-order interactions among genetic polymorphisms in nucleotide excision repair pathway genes and smoking in modulating bladder cancer risk. Carcinogenesis 2007, 28, 2160–2165. [CrossRef]

101. Enjuanes, A.; Benavente, Y.; Bosch, F.; Martin-Guerrero, I.; Colomer, D.; Pérez-Álvarez, S.; Reina, O.; Ardanaz, M.T.; Jares, P.; García-Orad, A.; et al. Genetic Variants in Apoptosis and Immunoregulation-Related Genes Are Associated with Risk of Chronic Lymphocytic Leukemia. Cancer Res. 2008, 68, 10178–10186. [CrossRef]

102. Pan, J.; Lin, J.; Izzo, J.G.; Liu, Y.; Xing, J.; Huang, M.; Ajani, J.A.; Wu, X. Genetic susceptibility to esophageal cancer: The role of the nucleotide excision repair pathway. Carcinogenesis 2009, 30, 785–792. [CrossRef]

103. Wu, X.; Gu, J.; Grossman, H.B.; Amos, C.I.; Etzel, C.; Huang, M.; Zhang, Q.; Millikan, R.E.; Lerner, S.; Dinney, C.P.; et al. Bladder Cancer Predisposition: A Multigenic Approach to DNA-Repair and Cell-Cycle–Control Genes. Am. J. Hum. Genet. 2006, 78, 464–479. [CrossRef]

104. Zhang, M.; Huang, W.-Y.; Andreotti, G.; Gao, Y.T.; Rashid, A.; Chen, J.; Sakoda, L.C.; Shen, M.-C.; Wang, B.-S.; Chanock, S.; et al. Variants of DNA Repair Genes and the Risk of Biliary Tract Cancers and Stones: A Population-Based Study in China. Cancer Epidemiol. Biomark. Prev. 2008, 17, 2125–2127. [CrossRef]

105. Lin, J.; Pu, X.; Wang, W.; Matin, S.; Tannir, N.M.; Wood, C.G.; Wu, X. Case–control analysis of nucleotide excision repair pathway and the risk of renal cell carcinoma. Carcinogenesis 2008, 29, 2112–2119. [CrossRef]

106. Wang, Y.; Spitz, M.R.; Lee, J.J.; Huang, M.; Lippman, S.M.; Wu, X. Nucleotide Excision Repair Pathway Genes and Oral Premalignant Lesions. Clin. Cancer Res. 2007, 13, 3753–3758. [CrossRef]

107. Wu, X.; Lu, C.; Ye, Y.; Chang, J.; Yang, H.; Lin, J.; Gu, J.; Hong, W.K.; Stewart, D.; Spitz, M.R. Germline genetic variations in drug action pathways predict clinical outcomes in advanced lung cancer treated with platinum-based chemotherapy. Pharmacogenet. Genom. 2008, 18, 955–965. [CrossRef]

108. Palugulla, S.; Devaraju, P.; Kayal, S.; Narayan, S.K.; Mathaiyan, J. Genetic polymorphisms in cyclin H gene are associated with oxaliplatin-induced acute peripheral neuropathy in South Indian digestive tract cancer patients. Cancer Chemother. Pharmacol. 2018, 82, 421–428. [CrossRef]

109. Custodio, A.; Moreno-Rubio, J.; Aparicio, J.; Gallego-Plazas, J.; Yaya, R.; Maurel, J.; Higuera, O.; Burgos, E.; Ramos, D.; Calatrava, A.; et al. Pharmacogenetic predictors of severe peripheral neuropathy in colon cancer patients treated with oxaliplatin-based adjuvant chemotherapy: A GEMCAD group study. Ann. Oncol. 2013, 25, 398–403. [CrossRef]

110. Lolli, G.; Johnson, L.N. CAK-Cyclin-dependent Activating Kinase: A key kinase in cell cycle control and a target for drugs? Cell cycle 2005, 4, 572–577. [CrossRef]

111. Emmert, S.; Schneider, T.D.; Khan, S.G.; Kraemer, K.H. The human XPG gene: Gene architecture, alternative splicing and single nucleotide polymorphisms. Nucleic Acids Res. 2001, 29, 1443–1452. [CrossRef]
112. Sang, L.; Lv, Z.; Sun, L.-P.; Xu, Q.; Yuan, Y. Impact of SNP-SNP interactions of DNA repair gene ERCC5 and metabolic gene GSTP1 on gastric cancer/atrophic gastritis risk in a Chinese population. World J. Gastroenterol. 2018, 24, 602–612. [CrossRef]

113. Negovan, A.; Iancu, M.; Moldovan, V.; Mocan, S.; Banescu, C. The Interaction between GSTT1, GSTM1, and GSTP1 Ile105Val Gene Polymorphisms and Environmental Risk Factors in Premalignant Gastric Lesions Risk. BioMed Res. Int. 2017, 2017, 7365080. [CrossRef]

114. Khabaz, M.N. Polymorphism of the glutathione S-transferase P1 gene (GST-pi) in breast carcinoma. Pol. J. Pathol. 2014, 65, 141–146. [CrossRef]

115. Cheng, L.; Sturgis, E.M.; Eicher, S.A.; Spitz, M.R.; Wei, Q. Expression of nucleotide excision repair genes and the risk for squamous cell carcinoma of the head and neck. Cancer 2002, 94, 393–397. [CrossRef]

116. Hussain, S.K.; Mu, L.-N.; Cai, L.; Chang, S.-C.; Park, S.L.; Oh, S.S.; Wang, Y.; Goldstein, B.Y.; Ding, B.-G.; Jiang, Q.; et al. Genetic Variation in Immune Regulation and DNA Repair Pathways and Stomach Cancer in China. Cancer Epidemiol. Biomark. Prev. 2009, 18, 2304–2309. [CrossRef]

117. Paszkowska-Szcuz, K.; Scott, R.J.; Serrano-Fernandez, P.; Mirecka, A.; Gapska, P.; Gorski, B.; Cybulski, C.; Maleszka, R.; Sulikowski, M.; Nagay, L.; et al. Xeroderma pigmentosum genes and melanoma risk. Int. J. Cancer 2013, 133, 1094–1100. [CrossRef]

118. Ma, H.; Yu, H.; Liu, Z.; Wang, L.-E.; Sturgis, E.M.; Wei, Q. Polymorphisms of XPG/ERCC5 and risk of squamous cell carcinoma of the head and neck. Pharmacogenet. Genom. 2012, 22, 50–57. [CrossRef]

119. Huang, J.; Liu, X.; Fang, L.-L.; Long, J.-T.; Zhu, J.; Hua, R.-X.; Li, J. XRCC5 polymorphisms and cancer susceptibility: Evidence from 47 studies. Oncotarget 2017, 8, 37263–37277. [CrossRef]

120. Su, C.-H.; Chang, W.-S.; Hu, P.-S.; Hsiao, C.-L.; Ji, H.-X.; Liao, C.-H.; Yueh, T.-C.; Chuang, C.-L.; Tsai, C.-W.; Hsu, C.-M.; et al. Contribution of DNA Double-strand Break Repair Gene XRCC3 Genotypes to Triple-negative Breast Cancer Risk. Cancer Genom. Proteom. 2015, 12, 359–367. [CrossRef]

121. Derwahl, M.; Nicula, D. Estrogen and its role in thyroid cancer. Endocr. Relat. Cancer 2014, 21, T273–T283. [CrossRef]

122. Nielsen, S.M.; White, M.G.; Hong, S.; Aschbrook-Kilfoy, B.; Kaplan, E.L.; Angelos, P.; Kulkarni, S.A.; Olopade, O.I.; Grogan, R.H. The Breast–Thyroid Cancer Link: A Systematic Review and Meta-analysis. Cancer Epidemiol. Biomark. Prev. 2016, 25, 231–238. [CrossRef]

123. Chuffa, L.G.; Lupi-Júnior, L.A.; Costa, A.B.; Amorim, J.P.; Seiva, F.R. The role of sex hormones and steroid receptors on female reproductive cancers. Steroids 2017, 118, 93–108. [CrossRef]

124. Cavalieri, E.; Rogn, E. The molecular etiology and prevention of estrogen-initiated cancers: Ockham’s Razor: Pluralitas non est ponenda sine necessitate. Plurality should not be posited without necessity. Mol. Asp. Med. 2013, 36, 1–55. [CrossRef]

125. Vahteristo, P.; Ojala, S.; Tamminen, A.; Tommiska, J.; Sammalkorpi, H.; Kiuru-Kuhlefelt, S.; Eerola, H.; Aaltosen, L.A.; Aittomäki, K.; Nevanlinna, H. No MSH6 germline mutations in breast cancer families with colorectal and/or endometrial cancer. J. Med. Genet. 2005, 42, e22. [CrossRef]

126. Hsieh, P.; Yu, H.; Liu, Z.; Wong, L.-E.; Sturgis, E.M.; Wei, Q. Polymorphisms of GSTT1/ GSTP1 and GSTM1/ GSTM2 on gastric cancer risk. World J. Gastroenterol. 2009, 15, 2304–2309. [CrossRef]

127. Dralle, H.; Machens, A.; et al. Impact of SNP-SNP interactions of DNA repair gene hMSH2 and shMSH6 on gastric cancer/atrophic gastritis risk in a Chinese population. Br. J. Cancer 2005, 92, 2286–2291. [CrossRef]

128. Dralle, H.; Machens, A.; Basa, J.; Fatourechi, V.; Franceschi, S.; Hay, I.D.; Nikiforov, Y.E.; Pacini, F.; Pasieka, J.L.; Sherman, S.I. Follicular cell-derived thyroid cancer. Nat. Rev. Dis. Prim. 2015, 1, 15077. [CrossRef]

129. Li, C.; Yin, M.; Wang, L.-E.; Amos, C.I.; Zhu, D.; Lee, J.E.; Gershenson, J.E.; Grimm, E.A.; Wei, Q. Polymorphisms of Nucleotide Excision Repair Genes Predict Melanoma Survival. J. Investig. Dermatol. 2013, 133, 1813–1821. [CrossRef]

130. Mumbrekar, K.D.; Goutham, H.V.; Vadhiraja, B.M.; Bola Sadashiva, S.R. Polymorphisms in double strand break repair related genes influence radiosensitivity phenotype in lymphocytes from healthy individuals. DNA Repair 2016, 40, 27–34. [CrossRef]
133. Edelbrock, M.A.; Kaliyaperumal, S.; Williams, K.J. Structural, molecular and cellular functions of MSH2 and MSH6 during DNA mismatch repair, damage signaling and other noncanonical activities. *Mutat. Res.* 2013, 743, 53–66. [CrossRef]

134. Bridge, G.; Rashid, S.; Martin, S. DNA Mismatch Repair and Oxidative DNA Damage: Implications for Cancer Biology and Treatment. *Cancers* 2014, 6, 1597–1614. [CrossRef]

135. Li, Z.; Pearlman, A.H.; Hsieh, P. DNA mismatch repair and the DNA damage response. *DNA Repair* 2016, 38, 94–101. [CrossRef]

136. Kamiza, A.B.; Hsieh, L.-L.; Tang, R.; Chien, H.-T.; Lai, C.-H.; Chiu, L.-L.; Lo, T.-P.; Hung, K.-Y.; You, J.-F.; Wang, W.-C.; et al. Polymorphisms of DNA repair genes are associated with colorectal cancer in patients with Lynch syndrome. *Mol. Genet. Genom. Med.* 2018, 6, 533–540. [CrossRef]

137. Vral, A.; Willems, P.; Claes, K.; Poppe, B.; Perletti, G.; Thierens, H. Combined effect of polymorphisms in *Rad51* and *Xrcc3* on breast cancer risk and chromosomal radiosensitivity. *Mol. Med. Rep.* 2011, 4, 901–912. [CrossRef]

138. Bishop, D.K.; Ear, U.; Bhattacharyya, A.; Calderone, C.; Beckett, M.; Weichselbaum, R.R.; Shinohara, A. Xrcc3 Is Required for Assembly of Rad51 Complexes in vivo. *J. Biol. Chem.* 1998, 273, 21482–21488. [CrossRef]

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