Interaction between Dietary and Lifestyle Risk Factors and N-Acetyltransferase Polymorphisms in B-Cell Lymphoma Etiology

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Abstract

Background: Gene-environment interactions are suggested to play a role in lymphomagenesis.

Methods: We tested the interaction between the NAT1/NAT2 phenotype, as inferred by the respective genotypes, and exposure to dietary and lifestyle risk factors, in 199 incident lymphoma cases and 188 population controls. We used unconditional logistic regression to calculate the odds ratio (OR) and its 95% confidence interval for lymphoma (all subtypes combined) and B-cell lymphoma, associated to the rapid NAT1 phenotype and to the intermediate and slow NAT2 phenotype, and to the estimated dietary intake of heterocyclic amines and folate, current smoking, coffee, and use of permanent hair dyes, as well as the respective interaction terms. We adjusted the ORs by age, gender, and education, and we used the likelihood ratio test to test the interaction between the NAT1, NAT2 phenotype and the dietary and lifestyle variables.

Results: We observed an increase in risk of lymphoma (all subtypes combined) and B-cell lymphoma in particular associated with the estimated above median dietary intake of heterocyclic amines (OR = 4.2, 95%CI 1.2 – 14.8) and folate (OR = 4.1, 95%CI 0.7 – 22.4) among subjects with the NAT1 rapid acetylator phenotype, but not independent on the NAT1 phenotype. The test for interaction was significant for heterocyclic amines, but not for folate (p for interaction = 0.262 and 0.076 respectively). Ever use of permanent hair dyes was associated with an elevated risk independent on the NAT1, NAT2 phenotypes; risk decreased to null among intermediate and slow NAT1 acetylators (p for interaction = 0.010).

Conclusions: Our results suggest that NAT1, NAT2 polymorphisms interact with dietary and lifestyle exposures in modulating risk of lymphoma and particularly B-cell lymphoma.

Keywords: Lymphomas; N-acetyl transferase; Molecular epidemiology; Gene-environment interaction; Smoking; Coffee; Folate; Heterocyclic amines

Abbreviations: CI: Confidence Interval; DLBCL: Diffuse Large B-Cell Lymphoma; FL: Follicular Lymphoma; HWE: Hardy–Weinberg Equilibrium; NAT1: N-Acetyl Transferase, Type 1; NAT2: N-Acetyl Transferase, Type 2; NHL: Non Hodgkin Lymphoma; OR: Odds Ratio; SES: Socio-Economic Status

Introduction

Typical substrates of the arylamine N-acetyltransferase 1 and 2 (NAT1, NAT2) enzymes include aromatic amines, putative responsible for the excess of non Hodgkin lymphoma (NHL) risk associated with use of hair dyes [1,2]; heterocyclic amines, known animal carcinogens, found in overcooked red meat but also occurring in well done roasted chicken, salmon, and cigarette smoke condensate [3]; and folate [4,5], which is implicated in one-carbon transfer reactions for DNA synthesis and methylation [6]. Notably, all these agents have been investigated as possible NHL risk factors [1,2,4-12], and, although inconclusively, NAT1, NAT2 gene polymorphisms have also been related to risk of NHL and its major B-cell subtypes, namely diffuse large B-cell lymphoma (DLBCL), and follicular lymphoma (FL) [13-16]. It is plausible that specific nutrients and chemicals from environmental exposures and lifestyle habits might act as carcinogens, and/or they might interfere with epigenetic conditions in silencing or enhancing the expression of genes implicated in the mechanisms leading to the development of specific lymphoma subtypes. Therefore, we explored risk of lymphoma (all subtypes combined), and B-cell lymphoma associated with the genotype-based NAT1/NAT2 polymorphisms and their interaction with dietary and lifestyle risk factors conveying exposure to typical NAT1, NAT2 substrates in a population-based case-control study.

Methods

Details on the study can be found elsewhere [16]. Briefly, in 1999-2003, adult incident cases with a diagnosis of lymphoma were recruited at the A. Businco Oncology Hospital in Cagliari, and the S. Francesco Hospital in Nuoro, two major referral centres for diagnosis and treatment of haematological malignancies in Sardinia, Italy. Population controls were randomly selected within the study area, frequency matched to cases by age (25-74 years by 5-year age-groups), gender, and geographic area of residence. Study participants (88% of eligible cases...
and 69.2% of eligible controls) underwent an in-person interview with a semi-structured questionnaire, including information on smoking, use of hair dyes and diet, and donated a 20 ml blood sample. A DNA sample was available for 248/322 participating lymphoma patients (77%), and, 208/446 (47%) population controls. For the purposes of this analysis, we focused on lymphoma (all subtypes combined) and on the B-cell lymphoma subgroup, as the study size was not sufficient to explore gene-environment interactions by histologically confirmed lymphoma subtype.

As previously described [16], we used multiplex PCR and allele specific primers to detect NAT1 (*4, *10 and *11) and NAT2 (*5A, *5B, *5C, *6 and *7) haplotypes, as suggested by Smith et al. [17]. Laboratory technicians were blind to the case-control status. Duplicate samples from 50 cases and 100 controls were interspersed throughout the plates for concordance analysis, which rate was 100% for all assays. Hardy–Weinberg Equilibrium (HWE) was tested in controls using the \( \chi^2 \) Goodness-of-fit test: no HWE deviation was detected.

Based on functional information [18,19], we categorized the NAT1 genotypes by the associated rapid (NAT1 *10/*10, *4/*4, *4/*11, *10/*11 or *11/*11 genotypes) or slow acetylator (NAT1 *4/*4 or *4/*10 genotype) phenotype, and the NAT2 genotypes by the rapid (NAT2 *4/*4 genotype) intermediate (NAT2 *4/*5, *4/*6, and *4/*7 genotypes) and slow acetylator phenotype (NAT2 *5/*5, *5/*6, *5/*7, *6/*6, *6/*7 and *7/*7 genotypes).

Using the questionnaire information, we categorized smoking as a binary variable (current smokers and ex smokers who quitted from <5 years vs never smokers and ex smokers who quitted smoking ≥ 5 years before interview). Coffee intake was highly prevalent in our study population, with 72% study subjects having coffee at least once a day; therefore, we categorized coffee intake as daily vs less than daily. We used the frequency of intake of well done and very well done red meat and cooking method to define a score of dietary intake of heterocyclic amines; we tested score values above the median against below median values. Dietary folate intake was derived from the frequency of intake of 20 food items reported as its major sources [20]; we summed up the intake frequency of each individual item weighted by its reported average content in folate to build a score, which was subsequently categorized as a binary variable (above the median value vs below median). Use of any permanent hair dye was considered to convey exposure to specific aromatic amines according to the type of colour [21], particularly before 1980. To allow acceptable statistical power to this analysis, we only used the binary ever/never use of permanent hair dyes.

Odds ratios (OR) and 95% confidence intervals (95% CI) for lymphoma (all subtypes combined) and B-cell lymphoma associated with the rapid NAT1 phenotype and with the intermediate and slow NAT2 phenotype, the lifestyle risk factors as described above, and the respective interaction terms were estimated using unconditional logistic regression analysis, adjusting by age (continuous), gender and education. We used the likelihood ratio test [22] to formally test the interaction between the NAT1, NAT2 phenotype and the dietary and lifestyle variables we considered. Adjustment for the frequency matching factors was deemed necessary because of the varying acceptance rate of the biological part of the study protocol by age and gender. Education was used as a surrogate for socio-economic status (SES), and it was included as a covariate in the regression models to account for SES-related variation in the acceptance rate of the full study protocol, and in lifestyle habits, possibly conveying exposure to known lymphoma risk factors. All the analyses were performed with SPSS version 16.0 (SPSS Inc., Chicago, IL, USA).

The study protocol was approved by the Ethical Committee of the University Hospital of Cagliari, and by the Ethical Committee of the Local Health Unit N. 3, Nuoro. All study participants signed an informed consent form according to the Helsinki declaration.

**Results**

Table 1 shows selected characteristics of the study population. As a result of the lower acceptance rate of blood withdrawal among male controls, the gender distribution differed by case/control status (\( \chi^2 = 3.71, p = 0.054 \)). All the dietary and lifestyle variables were approximately equally distributed by disease status, but ever use of hair dyes, which prevalence was approximately double among the cases in respect to the controls. The NAT1 haplotypes could not be determined for 49 cases and 20 controls, leaving 199 cases and 188 controls available for the analysis of the NAT1 interactions. The NAT2 haplotypes could not be determined for 54 cases and 23 controls, leaving 195 cases and 185 controls available for the analysis of the NAT2 interactions. Among study participants with available NAT1/NAT2 genotypes, the genotypes expressing the NAT1 rapid acetylator phenotype and those expressing the NAT2 slow acetylator phenotype were more represented among the cases.

Table 2 shows the results of the logistic regression model predicting risk of lymphoma (all subtypes combined) and B-cell lymphoma, associated with the dietary or lifestyle risk factor, the NAT1 phenotype, and the respective interaction term. Although, after adjusting for the NAT1 polymorphism, risk of lymphoma (all subtypes combined) and mature B-cell lymphoma associated with the estimated above median heterocyclic amines dietary intake was not elevated risk significantly increased among subjects carrying the rapid NAT1 acetyltransferase phenotype (OR = 4.2, 95% CI 1.2 – 14.5; p for interaction = 0.024; and OR = 4.2, 95% CI 1.2 – 14.8; p for interaction = 0.026, respectively). A dietary folate intake above the median was also associated with an elevated risk of lymphoma (all subtypes combined) (OR = 3.8, 95% CI 1.0 – 14.5), and B-cell lymphoma in particular (OR = 4.1, 95% CI 0.7 – 22.4), but no gene-environment interaction was detected (\( p = 0.094 \) and \( p = 0.076 \), respectively). Risk of lymphoma, and particularly B-cell lymphoma, was significantly elevated among hair dye users independent on the NAT1 acetyltransferase phenotype, while risk was null among hair dye users with the rapid NAT1 acetyltransferase phenotype, although no interaction was detected. Current smoking and daily coffee intake did not show an independent association with risk of lymphoma (all subtypes combined) nor with risk of B-cell lymphoma, nor we saw evidence of an interaction with the NAT1 rapid acetyltransferase phenotype.

Although with a very wide confidence interval, ever use of hair dyes was a strong independent predictor of risk of lymphoma (all subtypes combined) and of B-cell lymphoma, but risk was below unity among subjects with the intermediate and slow NAT2 acetyltransferase phenotype (\( p \) for interaction = 0.005, and \( p = 0.009 \), respectively) (Table 3). In the models including the NAT2 polymorphism, smoking, coffee, heterocyclic amines, and folate did not show an independent association with risk of lymphoma (all subtypes combined) and B-cell lymphoma, nor we detected an interaction between these lifestyle and dietary risk factors and the NAT2 acetyltransferase phenotype.

**Discussion**

We used functional data to infer the genotype-based NAT1/NAT2 acetyltransferase phenotype, and questionnaire information to infer exposure to putative carcinogen substrates from diet and lifestyle habits. Our
results suggest an interaction between above median dietary intake of heterocyclic amines and the NAT1 rapid acetylator phenotype in increasing risk of lymphoma and particularly B-cell lymphoma. We also observed that the NAT2 slow acetylator phenotype reduced to null the elevated risk of lymphoma, and B-cell lymphoma in particular, associated with use of hair dyes. Our findings are consistent with reports of an elevated non Hodgkin lymphoma (NHL) risk associated with intake of very well done red meat [7] and with use of permanent hair dyes [1,2], as well as with the hypothesis that polymorphisms in the genes involved in the metabolism of heterocyclic amines and aromatic amines, the putative carcinogens responsible for the associations we detected, modulate risk. On the other hand, we did not observe an interaction between the NAT1/NAT2 polymorphism with dietary folate intake, smoking and coffee. However, whether positive or null, our findings are purely tentative because of the small size and the limited statistical power of our study; therefore, the extreme variations in risk we observed should be interpreted cautiously.

As a consequence of the insufficient statistical power, we were unable to test the interactions by histological subtype. This is a further important limitation in our study, as substantial heterogeneity in risk factors has been described by lymphoma subtypes [23]. Nonetheless, our results prompt the need of further more powerful analyses, possibly using biomarkers of dose of the exposures of interest to reduce misclassification.

In conclusion, while our results seem intriguing in confirming the hypothesis of gene-environment interactions in the etiology of lymphoma and particularly B-cell lymphoma, caution is recommended in their interpretation, as further carefully planned, more powerful and more detailed studies are warranted.

Acknowledgments

Funding for this study was provided to PC by the Italian Ministry for Education, University and Research (PRIN 2007 prot.2007WEJLB, PRIN 2009 prot. 20092ZELR2), and the Italian Association for Cancer Research (AIRC, Investigator Grant 11855). Support to MGE was provided by Fondazione Banco di Sardegna 2010-2012, and Regione Autonoma della Sardegna (LR7 CRP-59812/2012). This study would have been impossible without the voluntary and disinterested contribution of all the patients and the population controls who accepted donating blood samples for the purposes of our study.

Author contributions

PC, MGE, and AS designed the study. M.Z, SS, GM, and MGE conducted the laboratory analyses. GS, TN, MC, and PC managed the data and conducted the

Table 1: Selected characteristics of the study population.

| Covariate | Cases | Controls | Total |
|-----------|-------|----------|-------|
| Age, mean (sd) | | | |
| men | 55.7 (13.8) | 58.1 (13.0) | 56.8 (13.5) |
| women | 57.9 (12.6) | 57.4 (12.4) | 57.6 (12.5) |
| all | 56.5 (13.4) | 57.8 (12.7) | 57.1 (13.0) |
| Gender: n (%) | | | |
| men | 121 (60.8) | 96 (51.1) | 217 (56.1) |
| women | 78 (39.2) | 92 (48.9) | 170 (43.9) |
| Education: n (%) | | | |
| <= middle school | 154 (66.1) | 134 (68.6) | 303 (66.4) |
| high school | 32 (15.7) | 43 (21.2) | 85 (18.2) |
| university | 12 (5.6) | 11 (5.8) | 26 (5.7) |
| missing | 0 | 1 | 1 |
| NAT1 phenotype | | | |
| slow | 155 (77.9) | 170 (90.4) | 325 (84.0) |
| rapid | 44 (22.1) | 18 (9.6) | 62 (16.0) |
| NAT2 phenotype | | | |
| rapid | 11 (5.7) | 46 (24.9) | 57 (15.0) |
| intermediate | 79 (40.7) | 79 (42.7) | 158 (41.7) |
| slow | 104 (53.6) | 60 (32.4) | 164 (43.3) |
| missing | 5 | 3 | 8 |
| Smoking | | | |
| Never, ex smokers (5 years) | 140 (70.4) | 138 (73.4) | 278 (71.8) |
| Current, ex smokers (<5 years) | 59 (29.6) | 50 (26.6) | 109 (28.2) |
| Heterocyclic amines | | | |
| Below median | 79 (39.7) | 75 (39.9) | 154 (39.8) |
| Above median | 120 (60.3) | 113 (60.1) | 233 (60.2) |
| Coffee | | | |
| Less than daily | 55 (27.6) | 53 (28.2) | 108 (27.9) |
| Once a day or more | 144 (72.4) | 135 (71.8) | 279 (72.1) |
| Folate | | | |
| Below median | 124 (50.0) | 104 (50.0) | 228 (50.0) |
| Above median | 124 (50.0) | 104 (50.0) | 228 (50.0) |
| Hair dyes | | | |
| Never | 174 (81.4) | 176 (93.6) | 350 (90.4) |
| Ever | 25 (12.6) | 12 (6.4) | 37 (9.6) |

Table 2: Risk of lymphoma (all subtypes combined) and B-cell lymphoma associated with dietary and lifestyle risk factors, the NAT1 rapid acetylator phenotype, and the respective interactions in Sardinia, Italy. ORs are adjusted by age (continuous), gender, and education.
| External risk factor | NAT2 acetylator phenotype intermediate vs rapid | NAT2 acetylator phenotype intermediate vs rapid | Likelihood ratio test, OR (95% CI) |
|----------------------|---------------------------------------------|---------------------------------------------|---------------------------------|
| Categories | OR (95% CI) | OR (95% CI) | df (p value) |
| All lymphomas combined |  |  |  |
| Model 1. Smoking | Current smokers vs never + ex smokers | 1.9 (0.4 – 8.1) | 5.0 (2.0 – 12.2) | 9.3 (3.8–22.9) | 1.52, 2, 0.468 |
| Model 1: interaction terms |  |  |  |
| Model 2. Coffee | Daily vs less than daily | 0.4 (0.1 – 1.4) | 2.8 (0.9 – 8.8) | 2.9 (0.9 – 9.1) | 3.82, 2, 0.148 |
| Model 1: interaction terms |  |  |  |
| Model 3. Heterocyclic amines | >median vs <median | 0.8 (0.2 – 3.1) | 4.4 (1.5 – 13.2) | 6.9 (2.2–21.8) | 0.06, 2, 0.970 |
| Model 1: interaction terms |  |  |  |
| Model 4. Folate | >median vs <median | 0.4 (0.1 – 2.0) | 3.1 (1.4– 7.1) | 5.1 (2.2–11.7) | 2.48, 2, 0.289 |
| Model 1: interaction terms |  |  |  |
| Model 5. Hair dyes | Ever use vs never | 25.5 (2.2 –293) | 5.1 (2.2 –11.7) | 10.2 (4.4–23.6) | 4.1 (1.4 – 12.2) |
| Model 1: interaction terms |  |  |  |
| Model 1. Smoking | Current smokers vs never + ex smokers | 1.6 (0.3 – 7.5) | 4.7 (1.9 – 11.5) | 8.3 (3.4–20.7) | 1.59, 2, 0.452 |
| Model 1: interaction terms |  |  |  |
| Model 2. Coffee | Daily vs less than daily | 0.4 (0.1 – 1.8) | 3.3 (1.0 – 10.9) | 3.2 (1.0–10.5) | 2.71, 2, 0.258 |
| Model 1: interaction terms |  |  |  |
| Model 3. Heterocyclic amines | >median vs <median | 0.7 (0.2 –2.7) | 4.1 (1.4 –12.2) | 6.0 (1.9–19.1) | 0.09, 2, 0.956 |
| Model 1: interaction terms |  |  |  |
| Model 4. Folate | >median vs <median | 0.2 (0.02–1.7) | 2.9 (1.3–6.7) | 4.5 (2.0–10.5) | 4.25, 2, 0.119 |
| Model 1: interaction terms |  |  |  |
| Model 5. Hair dyes | Ever use vs never | 29.9 (2.6 –349) | 5.5 (2.3 –13.2) | 9.8 (4.0–23.6) | 9.23, 2, 0.010 |
| Model 1: interaction terms |  |  |  |

Table 3: Risk of lymphoma (all subtypes combined) and B-cell lymphoma associated with NAT2 acetylator phenotypes, as inferred from the respective haplotype-based genotypes, in Sardinia, Italy. ORs are adjusted by age (continuous), gender, and education.
19. Zang Y, Doll MA, Zhao S, States JC, Hein DW (2007) Functional characterization of single-nucleotide polymorphisms and haplotypes of human N-acetyltransferase 2. Carcinogenesis 28: 1665-1671.
20. European Oncology Institute. Data base of food nutrients for epidemiological studies in Italy.
21. Munshi S (2004) Hair coloring products: colloid and surface phenomena.
22. Breslow NE, Day NE (1980) Statistical methods in cancer research. Volume I - The analysis of case-control studies. IARC Sci Publ : 5-338.
23. Morton LM, Slager SL, Cerhan JR, Wang SS, Vajdic CM, et al. (2014) Etiologic heterogeneity among non-Hodgkin lymphoma subtypes: the InterLymph Non-Hodgkin Lymphoma Subtypes Project. J Natl Cancer Inst Monogr 2014: 130-144.