Gene expression in blood reflects smoking exposure among cancer-free women in the Norwegian Women and Cancer (NOWAC) postgenome cohort

Nikita Baiju1*, Torkjel M. Sandanger1, Pål Sætrom2,3,4,5 & Therese H. Nøst1,5

Active smoking has been linked to modulated gene expression in blood. However, there is a need for a more thorough understanding of how quantitative measures of smoking exposure relate to differentially expressed genes (DEGs) in whole-blood among ever smokers. This study analysed microarray-based gene expression profiles from whole-blood samples according to smoking status and quantitative measures of smoking exposure among cancer-free women (n = 1708) in the Norwegian Women and Cancer postgenome cohort. When compared with never smokers and former smokers, current smokers had 911 and 1082 DEGs, respectively and their biological functions could indicate systemic impacts of smoking. LRRN3 was associated with smoking status with the lowest FDR-adjusted p-value. When never smokers and all former smokers were compared, no DEGs were observed, but LRRN3 was differentially expressed when never smokers were compared with former smokers who quit smoking ≤ 10 years ago. Further, LRRN3 was positively associated with smoking intensity, pack-years, and comprehensive smoking index score among current smokers; and negatively associated with time since cessation among former smokers. Consequently, LRRN3 expression in whole-blood is a molecular signal of smoking exposure that could supplant self-reported smoking data in further research targeting blood-based markers related to the health effects of smoking.

Tobacco smoking is one of the major threats to public health, and it is currently responsible for more than 8 million deaths worldwide each year1. Exposure to tobacco smoke is a risk factor for many chronic diseases, such as cardiac and pulmonary diseases and several cancers. Further, smoking can suppress the immune system and modifies a range of immunological functions2. Subclinical outcomes, such as increased oxidative stress, reduced antioxidant defences, increased inflammation, impaired immune status, and altered lipid profiles, have been observed in smokers when compared to their counterparts who never smoked3. Notably, more respiratory symptoms caused by exposure to tobacco smoke have been observed in women than men4,5. Thus, tobacco smoking has several detrimental health effects, which might appear not long after smoking initiation or up to several decades after exposure1,6.

The toxic components of tobacco smoke are first absorbed in the lungs and then enter the blood stream before being distributed throughout the body, making blood an appropriate biological material to study the systemic influences of exposure to tobacco smoke7. In addition, the collection of whole-blood (or simply, 'blood
samples’) is easy and minimally invasive, and these samples can reveal features that are relevant for studies of human health effects. Current exposure to tobacco smoke has been linked with modulated expression of many genes in blood, for example LRRN3, CLDN1, GPR15, ATF4, SOD2, and CDKN1C. Altered gene expression in blood has also been linked to diseases for which smoking is a risk factor. However, there is a need for a more thorough understanding of the variability in gene expression profiles in whole-blood in relation to quantitative measures of smoking exposure among ever smokers. Therefore, this cross-sectional analysis used data from 1708 cancer-free women participating in the prospective, population-based Norwegian Women and Cancer (NOWAC) postgenome cohort. Microarray-based gene expression profiles from bio-banked whole-blood samples were assessed according to smoking status and quantitative measures of smoking exposure (hereafter referred to as ‘smoking metrics’), such as smoking intensity, smoking duration, time since smoking cessation (TSC), pack-years, and comprehensive smoking index (CSI) scores. Enriched pathways and gene ontology (GO) categories of significant genes associated with smoking were also assessed.

Results
General characteristics of the study population. The current study was based on microarray data from cancer-free women participating in the NOWAC postgenome cohort. The full cohort consists of approximately 50,000 women (mean age: 49.78 years; mean body mass index (BMI): 23.38 kg/m²), all of whom have given a blood sample. In total, 1708 of these women have been included as cancer-free controls in various studies and have gene expression profiles available for study, and only these women were included in the present analyses. All included women had completed up to three comprehensive questionnaires before blood collection (main questionnaires), and an additional questionnaire on lifestyle factors was completed at the time of blood collection. Thus, information was available for up to four time points in total. Smoking status and smoking metrics (smoking intensity, smoking duration, TSC, pack-years, and CSI scores) were based on information from all four questionnaires. Current smokers (CS) were defined as those who were currently smoking at the time of blood collection, former smokers (FS) were defined as those who reported smoking cessation prior to the time of blood collection, and never smokers (NS) were defined as those who reported they had never smoked either prior to or at the time of blood collection. CS and FS combined represented ever smokers. We defined passive smokers (PS) as those who were passively exposed to smoking at their homes as adults. Gene expression values were available for 7713 unique genes for all the women in this study.

We investigated associations between smoking status and potential covariates, such as age and BMI at blood collection, and white blood cell (WBC) proportions, using Chi-square or Kruskal–Wallis tests. We then performed a ‘global test’ to indicate any association between these variables and the overall gene expression data. We considered variables that were significant in both of these tests as potential confounders and adjusted for these in further models (Supplementary Table S1).

There were 473, 613, and 622 CS, FS, and NS, respectively, among the 1708 women. The distributions of age and BMI at blood collection did not deviate markedly from normality, whereas the distribution of alcohol consumption was skewed (Fig. 1). Each of these distributions were similar across different categories of smoking status (Fig. 1A–C), but FS had the highest mean BMI and alcohol consumption, and NS had the highest mean age (Supplementary Table S1). Further, the smoking metrics—smoking intensity, smoking duration, pack-years, and CSI score—had the highest means for CS as compared to FS (Fig. 1D–H). Finally, there were 192, 147, and 100 PS among CS, FS, and NS, respectively.

Estimated white blood cell proportions. We estimated proportions of 22 types of WBCs using an in silico gene expression deconvolution method. CD8 T cells, naive CD4 T cells, resting NK cells, M0 macrophages, resting mast cells, and neutrophils were significantly associated with both smoking status and overall gene expression (Supplementary Table S2 and Supplementary Fig. S1). Further, we used linear regression to assess the associations between WBC proportions and smoking metrics. We observed that CD8 T cells were negatively associated with pack-years and CSI score; naive CD4 T cells were positively associated with smoking intensity, smoking duration, pack-years, and CSI score; resting NK cells were negatively associated with smoking intensity, smoking duration, pack-years, and CSI score but positively associated with TSC; resting mast cells were negatively associated with smoking duration; and neutrophils were negatively associated with TSC (Supplementary Table S3).

Differentially expressed genes dependent on smoking status. We used two adjusted (minimally- and fully-adjusted) models to assess the relationships between smoking status and gene expression profiles, using the ‘limma’ package for gene-wise linear models. In minimally-adjusted models, we adjusted for technical variables such as laboratory batch (laboratory plates) and sample storage time, while in fully-adjusted models, in addition to the technical variables, we included the following variables that were associated with both the exposure and the outcome: selected WBC proportions, age, BMI, and use of hormone replacement therapy at the time of blood collection, as well as information on alcohol consumption and use of oral contraceptives, which was taken from the main questionnaires. The presence of differentially expressed genes (DEGs) was determined by three comparisons of smoking status groups: CS-vs-NS, CS-vs-FS, and FS-vs-NS. We considered Benjamini–Hochberg false discovery rates (FDR) with the significance threshold FDR ≤ 0.05.

In minimally-adjusted models, there were 1009 DEGs in the CS-vs-NS comparison; 427 up-regulated and 582 down-regulated genes. Correspondingly, in the CS-vs-FS comparison, there were 1371 DEGs (559 up-regulated, 812 down-regulated). In fully-adjusted models, there were 911 DEGs in the CS-vs-NS comparison (355 up-regulated, 556 down-regulated; Fig. 2A,D), and 1082 DEGs in the CS-vs-FS comparison (435 up-regulated, 647 down-regulated; Fig. 2B,E). The two adjusted models had 670 overlapping DEGs in the CS-vs-NS comparison.
(Supplementary Table S4) and 839 in the CS-vs-FS comparison (Supplementary Table S5). Similarly, the CS-vs-NS and CS-vs-FS comparisons had 776 and 652 overlapping DEGs in the minimally- and fully-adjusted models, respectively. In the fully-adjusted models, there were 230 up-regulated and 422 down-regulated genes that overlapped between the CS-vs-NS and CS-vs-FS comparison and displayed the same direction of effects. The top-ranked gene (i.e., the gene with the lowest FDR adjusted p-values) in all comparisons was **LRRN3** (Supplementary Fig. S2). Receiver operating characteristics (ROC) curve analyses showed that expression levels of **LRRN3**, as measured by the Illumina arrays, could strongly distinguish CS from NS and moderately distinguish FS (with ≤ 10 years TSC) from NS (Supplementary Fig. S3). Moreover, in a subset of our dataset, **LRRN3** expression showed similar discriminative power as DNA methylation at the **AHRR** CPG site (cg05575921), which is a known marker for smoking exposure19. There were no DEGs in the FS-vs-NS comparison in either

**Figure 1.** Descriptive statistics of study participants by smoking status for (A) age at blood collection, (B) body mass index (BMI) at blood collection, (C) alcohol consumption at baseline, (D) smoking intensity, (E) smoking duration, (F) time since smoking cessation (TSC), (G) pack-years, and (H) comprehensive smoking index (CSI) scores. Yellow, blue, and red coloured violin plots represent kernel density estimates for never, former, and current smokers, respectively. White boxes extend from the 25th to the 75th percentile, vertical bars inside the box represent the median, whiskers extend 1.5 times the length of the interquartile range right and left side of the 75th and 25th percentiles respectively, and outliers are represented as black dots. The green diamond shaped dot represents the respective mean.
model (Fig. 2C). However, LRRN3 was the only DEG that remained significant when we included only FS with TSC ≤ 10 years and compared it with NS in the minimally-adjusted model (with log2 fold-change (logFC) = 0.34 and FDR = 3.63E−04). The p-values were uniformly distributed only in the FS-vs-NS comparison, but not in the other comparisons, as presented in quantile–quantile plots (Supplementary Fig. S4). Further, we used the ‘limma’ package to analyse the effects of passive smoking among NS, by contrasting all NS who were PS in adulthood with the other NS using the minimally-adjusted model. There were no DEGs when testing differences between PS (n = 100) and non-PS (n = 428).

**Figure 2.** Volcano plots for the test statistics in fully-adjusted models from the tests of differentially expressed genes (DEGs) in comparisons of (A) current versus never smokers, (B) current versus former smokers, and (C) former versus never smokers; and forest plots for the 10 top-ranked DEGs in tests of DEGs in comparisons of (D) current versus never smokers and (E) current versus former smokers. In volcano plots (A–C), red dots display up-regulated genes, blue dots display down-regulated genes, while grey dots display genes with FDR > 0.05; the x-axis presents log2 fold-changes and the y-axis presents − log10 of FDR adjusted p-values; and gene names displayed are the 20 top-ranked DEGs in the respective tests. In forest plots (D and E), dots in the x-axis represent log2 fold-changes and the y-axis represents DEGs with the lowest FDR adjusted p-values ranked from the top; the horizontal line for each gene represents their confidence interval; and the vertical blue dotted line represents no difference.
Analyses of smoking metrics within ever smokers. To identify genes associated with magnitude of smoking exposure, we used the ‘limma’ framework to identify genes for which the expression level correlated with the given smoking metrics among ever smokers. Specifically, we extended the minimally-adjusted model to include the given smoking metrics and analysed CS and FS separately.

In analyses of CS, the top-ranked gene, LRRN3 (logFC = 0.60, FDR = 4.70E−05), was positively associated with CSI score (Fig. 3). Further, there were five genes positively associated and two genes negatively associated with smoking intensity (Supplementary Table S6), where LRRN3 was the top-ranked gene, with a positive association (Supplementary Fig. S5). Likewise, there were three genes positively associated and two genes negatively associated with pack-years (Supplementary Table S7), where LRRN3 was the top-ranked gene, with a positive association (Supplementary Fig. S6). There were no genes significantly associated with smoking duration among CS.

Figure 3. Distributions of expression values for the top-ranked gene (LRRN3) (A) among never (yellow) and former (blue) smokers and (B) among current smokers according to comprehensive smoking index (CSI) scores. In figure (A), boxes extend from the 25th to the 75th percentile, horizontal bars represent the median, whiskers extend 1.5 times the length of the interquartile range above and below the 75th and 25th percentiles, respectively, and outliers are represented as points. In figure (B), the red line represents the linear regression fit and the shaded grey area its standard error.
In analyses of FS, the top-ranked gene, \textit{LRRN3} (logFC = −0.014, FDR = 2.63E−03), was negatively associated with TSC (Fig. 4). Correspondingly, \textit{NMRAL1} (logFC = −0.008, FDR = 2.72E−02) was negatively associated with pack-years (Supplementary Fig. S7). No genes were significantly associated with smoking intensity, smoking duration, or CSI scores among FS.

**Functional enrichment analyses.** To investigate the potential common functions of the identified DEGs affected by smoking, we performed functional enrichment analyses to identify GO biological processes (BP), GO molecular functions (MF), GO cellular components (CC), Kyoto encyclopedia of genes and genomes (KEGG) pathways, and REACTOME pathways enriched for DEGs in the CS-vs-NS and CS-vs-FS comparisons (Supplementary Tables S8–12, Fig. 5, and Supplementary Fig. S8). Analyses were performed for DEGs in fully-
Figure 5. Summary of functional enrichment analyses for up- and down-regulated genes for the (A) GO(BP) and (B) REACTOME pathway databases. The colour of the dots indicates the adjusted p-value, where red dots represent the most enriched categories; the 'GeneRatio' indicates the proportion of genes overlapping between lists of differentially expressed genes (DEGs) and the genes in gene ontology categories. GO: gene ontology; BP: biological processes; CS-vs-NS: comparison of current smokers versus never smokers; CS-vs-FS: comparison of current smokers versus former smokers; U: Up-regulated genes; D: Down-regulated genes.
adjusted models and separately for up-regulated and down-regulated genes. The numbers of enriched terms in the respective categories are presented in Table 1.

For both up-regulated and down-regulated genes, enriched categories overlapped considerably for genes that were significant according to the FDR in the CS-vs-NS and CS-vs-FS comparisons. However, there were more enriched categories among genes in the CS-vs-NS comparison, except for GO(MF), where there were significant categories only in the CS-vs-FS comparison. Considering the terms themselves, up-regulated genes were mostly enriched for terms related to translation, such as ribosome (KEGG and GO(CC)), protein localisation to endoplasmic reticulum (GO(BP)), and translation (REACTOME). Terms were also related to immune responses, such as humoral immune response, inflammatory response, and B cell activation (GO(BP)). In contrast, down-regulated genes were enriched for many terms related to circulatory functions, including response to wounding and regulation of blood vessel size (GO(BP)), and extracellular signalling, such as G protein-coupled receptor ligand binding (REACTOME), and plasma membrane region (GO(CC)).

**Discussion**

This study presents DEGs across categories of smoking status, as well as genes associated with different smoking metrics within ever smokers in the whole-blood of cancer-free women from the NOWAC postgenome cohort. These assessments, which used quantitative and repetitive smoking metrics, bring novel knowledge about the systemic responses to smoking exposure within ever smokers.

The study participants had similar proportions of CS, FS, and NS. They had comparable mean age and BMI at the time of blood collection as that of the full cohort, and to that of participants in other studies targeting the relation between smoking exposure and gene expression. Among the 7713 genes assessed, 911 and 1082 genes were differentially expressed in CS-vs-NS and CS-vs-FS comparisons, respectively. When looking at the DEGs in the CS-vs-NS comparison and the significant genes indicated in corresponding tests in a large meta-analysis containing 10,233 participants (51% women), we found that among the 285 DEGs in our study that overlapped with the 1270 DEGs in that study, 282 genes had the same direction of effects. Moreover, the mean expression levels for the 285 DEGs between CS-vs-NS that overlapped with DEGs identified in corresponding tests in the meta-analysis were higher than those DEGs that did not overlap (6.83; t = 5.23, p-value = 2.63E–07). Still, the absolute logFC for the overlapped (0.12) and non-overlapped DEGs (0.09; W = 128,066, p-value = 1.29E–14) were similar. This implies that the relation to smoking was consistent for hundreds of genes between these studies and demonstrates the comprehensive effects of smoking on gene expression in blood.

Around 40% of the genes were over-expressed in CS as compared to both NS and FS (i.e., 60% were under-expressed). Although higher proportions of up-regulated genes have been observed more frequently in other studies, higher proportions of down-regulated genes have also been observed. Interestingly, there could be sex differences in the directionality of observed DEGs, as one study comparing smokers and non-smokers observed that 29% of DEGs in men were down-regulated, compared to 62% in women. However, only about 4% of the DEGs in our study were in X-chromosomes in both the CS-vs-NS and CS-vs-FS comparisons. Notably, differences in gene expression between adult men and women do not need to originate in genes on the X-chromosomes, but a meta-analysis of sex expression differences in blood found that 25% of DEGs do map to the sex chromosomes. Thus, it is unlikely that the higher proportion of down-regulated genes in our study was due to the inclusion of women only.

Among NS, there were no genes associated with self-reported passive smoking in their homes as adults when compared to individuals with no passive smoking exposure. This could indicate that gene expression was more influenced by tobacco smoking of the women themselves. However, this could also be due to lack of statistical power or an imprecise exposure measure (lack of detailed information on timing, duration, and intensity of exposure).

Among CS, there were one, five, and three significant genes that were positively associated with CSI scores, smoking intensity, and pack-years, respectively. Among these, the top-ranked gene, LRRN3, was up-regulated in CS, which demonstrated that even within CS, LRRN3 had a higher expression among those with a higher smoking exposure, as represented by increasing CSI scores, smoking intensity, and pack-years. Among FS, there was one significant gene that was negatively associated with TSC (LRRN3) and one that was negatively associated with pack-years (NMRAL1). This demonstrated that within FS, those who had quit smoking recently had a higher expression of LRRN3 than those who had quit long ago, and FS with more pack-years had a lower expression of NMRAL1 than those with fewer pack-years. Also, when restricting the FS-vs-NS comparison to recent

| Database | CS-vs-NS | CS-vs-FS |
|----------|----------|----------|
| GO(BP)   | 33       | 22       |
| GO(MF)   | 4        | 6        |
| GO(CC)   | 23       | 14       |
| KEGG     | 1        | 1        |
| REACTOME | 31       | 34       |

Table 1. Number of enriched terms in different categories of enrichment analyses in comparisons of current versus never smokers (CS-vs-NS) and current versus former smokers (CS-vs-FS).
quitters (with TSC ≤ 10 years), LRRN3 remained significant in minimally-adjusted models. This indicates that there are differences in gene expression related to ongoing smoking exposure in women that persist for LRRN3 in those who recently stopped smoking. Thus, LRRN3 expression increases during smoking exposure and years after smoking cessation, but it eventually reverts back to levels similar to those of NS. However, according to the results of our linear model (Fig. 4), it appeared to take approximately 20–30 years for LRRN3 expression in FS to reach the average expression among NS. The difference in results from the overall FS-vs-NS comparison and those restricted to FS with TSC up to 10 years emphasizes that TSC needs to be taken into account when analysing smoking effects in FS.

LRRN3 was the top-ranked DEG in most comparisons, and its expression differences were large compared to the other DEGs identified. LRRN3 has been consistently indicated to be over-expressed in the whole-blood of CS or FS in previous studies9-11,13,14,16,21,22. This gene is highly expressed in the adrenal glands, the brain, and the lungs, but also in 11 other tissues23, and LRRN3 codes for a membrane protein. The GO database has little information on LRRN3’s potential functions, except that electronic annotations indicate that it is involved in the positive regulation of synapse assembly23,24. Notably, LRRN3 has six known SNPs25, but genetic variants in participants were not available in this study. Top-ranked DEGs other than LRRN3 in the CS-vs-NS and CS-vs-FS comparisons were PID1, RGL1, and STAB1, and in the analyses of ever smokers was NMRAL1. These genes are expressed in various tissues that differed across genes. The main functions of the aforementioned genes are to increase the proliferation of pre-adipocytes (PID1)26; to be involved in probable guanine nucleotide exchange factor (RGL1)27; and to act as a scavenger receptor for acetylated low-density lipoprotein, bind to both gram-positive and gram-negative bacteria, and to play a role in the defence against bacterial infection (STAB1)28. However, the interpretation of the potential function of these genes in blood in relation to smoking is not clear.

We performed functional enrichment analyses for GO(BP), GO(MF), and GO(CC) categories; and for KEGG and REACTOME pathways. This gave insight into knowledge of pathways for the identified DEGs29. The overlap in the enriched categories of the up-regulated and down-regulated genes in the CS-vs-NS and CS-vs-FS comparisons indicated that similar GO categories and pathways were enriched when current smoking exposure was compared to both FS and NS. Still, the enrichment was clearer when CS were compared to NS than to FS. The latter might be because the effect of smoking was not completely absent or was being slowly reduced in FS. In addition, the overall lack of overlap for enriched categories of the up-regulated and down-regulated genes likely demonstrated that separate groups of genes are involved in different pathways.

The GO enrichment analysis indicated categories such as peptide metabolic and biosynthetic processes, protein formation and translation, humoral immune response, structural constituent of ribosome and molecule activity, ribosomal subunits, and adherens junction were up-regulated in CS. In contrast, processes such as response to wounding, circulatory system, regulation of blood vessels and tube size and diameter, neuron projection development, drug and hydrogen peroxide catabolic processes, heme binding, cell body, and hemoglobin complex were down-regulated. Categories indicated in the KEGG and REACTOME enrichment analyses were largely in line with those in GO analysis. In summary, these categories indicate that the DEGs we identified were enriched for functions related to the physiological effects of smoking on the human body, which are well documented in the literatures. This is particularly relevant for the physiological functions linked to the cardiovascular system, as DEGs measured in blood could be directly influenced by such altered functions. For example, carbon monoxide binds to haemoglobin, thereby reducing the blood’s oxygen-carrying capacity30. Accordingly, our system, as DEGs measured in blood could be directly influenced by such altered functions. For example, carbon monoxide binds to haemoglobin, thereby reducing the blood’s oxygen-carrying capacity30. Accordingly, our system, as DEGs measured in blood could be directly influenced by such altered functions. For example, carbon monoxide binds to haemoglobin, thereby reducing the blood’s oxygen-carrying capacity30.
between smoking exposure and gene expression in blood have been conducted in rather small samples, ranging of gene expression in blood. Lastly, although RNA-sequencing has become a routinely used technology, results reflect smoking exposures even decades after cessation. In a subset of our data, Lund et al. demonstrated similar ability to discriminate CS and FS (with ≤ 10 years TSC) from NS as compared to methylation at the CpG cite in the AHRR gene. Therefore, the abilities of LRRN3 expression as a quantitative marker for discrimination of smoking status should be investigated in other population samples and with the comparison to other markers.

This study comprised a large number of women (n = 1708), whereas most studies targeting associations between smoking exposure and gene expression in blood have been conducted in rather small samples, ranging from 9 to 219 participants. The two exceptions are one population-based cohort study in the Netherlands with 3319 participants (65% women) and a meta-analysis with 10,233 participants (51% women). As mentioned, our results are in line with those observed in these studies. The present study included only cancer-free women, although we cannot disregard influences of other common chronic diseases. Further, this study was based on whole-blood samples, which is a relevant tissue to investigate the effects of smoking, as it expresses a large proportion of the genes in the human genome. Still, the top-ranked gene was the AHRR gene. Therefore, the abilities of LRRN3 expression as a quantitative marker for discrimination of smoking status should be investigated in other population samples and with the comparison to other markers.

In conclusion, our results demonstrated associations between smoking exposure and gene expression profiles in whole-blood of cancer-free women in the NOWAC postgenome cohort. The use of quantitative, reliable, and repeated measurements of past and recent smoking exposures was the novelty of this study, as it contributes new knowledge on systemic responses of smoking exposure. Close to a thousand DEGs in comparisons between CS and NS or FS, LRRN3, was the top-ranked gene. LRRN3 was also associated with CSI score, smoking intensity, and pack-years among CS; and with TSC among FS. Consequently, LRRN3 expression in blood is a molecular signal of smoking exposure that could supplant self-reported smoking data in gene expression studies of the association between smoking exposure and specific phenotypes. The biological functionality of the DEGs identified were linked to circulatory functions, translation, and immune responses, and could indicate systemic impacts of smoking. Genes that are differentially expressed depending on smoking exposure could be of interest in studies that focus on the effects of smoking exposure on health. This study has provided knowledge on the relationship of genes and pathways with detailed information on smoking exposure among cancer-free women.

Methods
Study population. The NOWAC study is a nation-wide, population-based prospective cohort study initiated in 1991. Currently, it includes approximately 172,000 Norwegian women aged 30–70 years. Women were randomly selected from the Norwegian National Population Register and sent an invitation letter along with a first questionnaire, which included detailed questions related to smoking exposure, height, weight, reproductive history, hormone replacement therapy, alcohol consumption, family history of breast cancer, dietary patterns, use of medication, and others. Since then, each woman has answered between one and three follow-up questionnaires (main questionnaires). The NOWAC study database takes information from the Cancer Registry of Norway, as well as national death and emigration registries. Details about the NOWAC study are available in Lund et al.

The current study was based on data from the NOWAC postgenome cohort, a sub-cohort of the NOWAC study. This consists of approximately 50,000 women who, between 2003 and 2006, had blood samples collected in PreAnalytiX (PAX) gene-tubes for gene expression analysis and, at the same time, answered a less extensive questionnaire about their lifestyle. The current study incorporated microarray-based expression profiles in biobanked whole-blood samples from cancer-free women in the NOWAC postgenome cohort, who were originally enrolled as controls in several studies on breast, lung, ovarian, and endometrial cancers, and diabetes. We obtained relevant questionnaire and registry information from NOWAC databases and excluded those women that did not respond to any questions on smoking exposure, those who participated in more than one study, and those who were diagnosed with cancer before 2017. This resulted in a final analytical sample of 1708 women.

Smoking status and smoking metrics. The main questionnaires included detailed questions regarding past and current smoking exposures, including ages at smoking initiation and cessation, average number of cigarettes smoked per day across age intervals, and details about passive smoking. Smoking status and smoking metrics (smoking intensity, smoking duration, TSC, pack-years, and CSI scores) were based on information from all main questionnaires and the questionnaire completed at the time of blood collection. Smoking intensity was defined as the average number of cigarettes smoked per day during years of active smoking, smoking duration was the duration of active smoking in years, and TSC was the time since smoking cessation in years. Pack-years quantify individual, long-term exposure to tobacco smoking; this variable was calculated by the formula:
Number of pack-years = (smoking intensity/20) × smoking duration. We considered 20 cigarettes in 1 pack, which is standard in the Norwegian context. CSI score is a cumulative measure of smoking exposure that incorporates smoking intensity (int), smoking duration (dur), and TSC (tsc). CSI scores were calculated using the formula\(^{18}\):

\[
\text{CSI} = (1 - 0.5 \delta \tau)/0.5 \exp(-\delta), \quad \text{where } \tau \text{ is an estimated half-life parameter, and } \delta \text{ is an estimated lag time parameter describing TSC and total duration as follows:}
\]

\[
tsc^* = \max(tsc - \delta, 0) \quad \text{and} \quad \text{dur}^* = \max(dur + tsc - \delta) - tsc^*.
\]

**Laboratory analyses and pre-processing of the gene expression data.** Total RNA was extracted and purified from PAX gene-tube samples according to the PAX gene blood RNA kit protocol at the Genomics Core Facility, Norwegian University of Science and Technology (NTNU), Trondheim. A NanoDrop ND 8000 spectrophotometer (ThermoFisher Scientific, Wilmington, DE, USA) was used to assess RNA purity, and bioanalyser capillary electrophoresis (Agilent Technologies, Palo Alto, CA, USA) was used to assess RNA integrity. Complementary RNA (cRNA) was prepared using the Illumina TotalPrep-96 RNA amplification kit, and hybridised to Illumina human WG-3 or HT-12 expression bead chip microarrays. The raw microarray images were processed in Illumina genome studio. The laboratory analysis date varied from January 2011 to January 2015.

For each study sample set separately, potential outliers were evaluated based on plots such as principal component analysis (PCA) plots and boxplots of probe signals displaying variation along with the laboratory quality measures\(^{44}\). We performed background correction, removed bad quality probes, and filtered probes detected in less than 20% of samples. Further, we performed log\(_2\) transformation and quantile normalisation before all data were combined and inspected for batch effects using PCA plots. We performed gene annotation using the Bioconductor packages ‘lumi’, ‘lumiHumanIDMapping’, and ‘illuminaHumanv4.db’\(^{45-47}\). If there were more than one probe annotated to each gene, the probe with the largest inter-quartile range was kept, which resulted in 7713 unique genes in the data analysed. Estimates for the proportions of 22 populations of WBCs in samples were obtained using the CIBERSORT procedure\(^{64}\).

**Statistical analyses.** We considered covariates and WBC proportions as potential confounders if they were significantly associated with smoking status using Chi-square or Kruskal–Wallis tests, and with overall gene expression data according to the ‘global test’ from the Bioconductor package ‘global test’\(^{51}\) for the gene-wise linear models. The presence of DEGs was determined by three comparisons of smoking status groups: CS-vs-NS, CS-vs-FS, and FS-vs-NS, using a significance threshold of FDR ≤ 0.05\(^{52}\). We used two adjusted (minimally- and fully-adjusted) models to assess the association between smoking status and DNA methylation at the CpG site cg05575921, using ROC curves. Differences in average expression and log\(_2\)FC between groups of DEGs were tested using t-test and Wilcoxon rank sum test, respectively. To evaluate common biological functions of results from the gene-wises tests, we performed functional enrichment analyses of GO(BP), GO(MF), and GO(CC) categories, and KEGG\(^{55}\) and REACTOME pathways for DEGs from different smoking status groups.

**Ethical statement.** The Regional Ethical Committee of North Norway (REK) has approved the NOWAC study and the NOWAC postgenome cohort (Reference Numbers: 2010/2075/REK Nord and 2014/1605/REK Nord, respectively), and the collection and storage of human biological material, the individual case–control studies, and gene expression analyses that this project was constructed from. The women gave written informed consent for the blood collection and for gene expression analyses\(^{42}\). All methods were carried out in accordance with relevant guidelines and regulations in the manuscript for human.

**Data availability**

Data cannot be shared publicly because of local and national ethical and security policy. Data access for researchers will be conditional on adherence to both the data access procedures of the Norwegian Women and Cancer Cohort and the UiT – The Arctic University of Norway (contact via Tonje Braaten <tonje.braaten@uit.no> and Arne Bastian Wiik <arne.b.wiik@uit.no>) in addition to an approval from the local ethical committee.

Received: 14 September 2020; Accepted: 15 December 2020
Published online: 12 January 2021

**References**

1. World Health Organization. *Don’t Let Tobacco Take Your Breath Away: Choose Health, not Tobacco: 31 May, World tobacco day* (accessed 10 August 2019). [https://www.who.int/docs/default-source/world-no-tobacco-day/wntb-2019-brochure.pdf?sfvrsn=deac371c_22](https://www.who.int/docs/default-source/world-no-tobacco-day/wntb-2019-brochure.pdf?sfvrsn=deac371c_22) (2019).

2. Sopori, M. Effects of cigarette smoke on the immune system. *Nat. Rev. Immunol.* 2, 372–377 (2002).
49. Goeman, J. J., Van De Geer, S. A., De Kort, F. & Van Houwelingen, H. C. A global test for groups of genes: testing association with a clinical outcome. Bioinformatics 20, 93–99 (2004).

50. R Core Team. R: A Language and Environment for Statistical Computing. R Foundation for Statistical Computing, Vienna, Austria (2015 (ver 3.2.1) and 2019 (ver 3.6.2)). https://www.R-project.org.

51. Ritchie, M. E. et al. limma powers differential expression analyses for RNA-sequencing and microarray studies. Nucl. Acids Res. 43, e47-e (2015).

52. Benjamini, Y. & Hochberg, Y. Controlling the false discovery rate: a practical and powerful approach to multiple testing. J. R. Stat. Soc. Ser. B (Methodol.) 57, 289–300 (1995).

53. Yu, G., Wang, L.-G., Han, Y. & He, Q.-Y. clusterProfiler: an R package for comparing biological themes among gene clusters. Omics J. Integr. Biol. 16, 284–287 (2012).

54. Yu, G. & He, Q.-Y. ReactomePA: an R/Bioconductor package for reactome pathway analysis and visualization. Mol. BioSyst. 12, 477–479 (2016).

55. Kanehisa, M. & Goto, S. KEGG: kyoto encyclopedia of genes and genomes. Nucl. Acids Res. 28, 27–30 (2000).

Acknowledgements
The authors gratefully extend their gratitude to all the Norwegian women participating in the NOWAC study. We thank B.A. and all the other personnel, who have handled the administration of the data collection and the biobanks. We thank T.B., M.L. and N.S. for providing the data set. Special thanks to J.I. for helping with data wrangling and T.P.-T. for editing the language of the manuscript. This research received no external funding. The publication charges for this article have been funded by a grant from the publication fund of UiT —The Arctic University of Norway, Tromsø, Norway.

Author contributions
N.B. made the contributions as first authors. T.H.N. and T.M.S. conceptualized the main research idea. T.H.N. curated the data. T.H.N., P.S., T.M.S., and N.B. designed the research methodology. N.B. performed the formal data analysis, wrote the manuscript, and Supplementary Information with significant contribution from T.H.N. All the co-authors discussed the results and reviewed the manuscript and Supplementary Information.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-020-80158-8.

Correspondence and requests for materials should be addressed to N.B.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2021