Hepatic Transcriptome Responses in Mice (Mus musculus) Exposed to the Nafion Membrane and Its Combustion Products

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Abstract

Nafion 117 membrane (N117), an important polymer electrolyte membrane (PEM), has been widely used for numerous chemical technologies. Despite its increasing production and use, the toxicity data for N117 and its combustion products remain lacking. Toxicity studies are necessary to avoid problems related to waste disposal in landfills and incineration that may arise. In this study, we investigated the histopathological alterations, oxidative stress biomarker responses, and transcriptome profiles in the liver of male mice exposed to N117 and its combustion products for 24 days. An ion-chromatography system and liquid chromatography system coupled to a hybrid quadrupole time-of-flight mass spectrometry were used to analyze the chemical compositions of these combustion products. The transcriptomics analysis identified several significantly altered molecular pathways, including the metabolism of xenobiotics, carbohydrates and lipids; signal transduction; cellular processes; immune system; and signaling molecules and interaction. These studies provide preliminary data for the potential toxicity of N117 and its combustion products on living organisms and may fill the information gaps in the toxicity databases for the currently used PEMs.

Introduction

During the last few decades, the polymer electrolyte membrane (PEM) has gained worldwide attention because of its wide use in a variety of chemical technologies, including for fuel cells, electrodialyzers, and sensors [1,2]. In particular, its application in fuel cells has attracted considerable interest because this technology has recently been used to supply energy to automobiles, for stationary power generation and for portable electronic devices [3,4]. Nafion, a perfluorosulfonic acid (PFSA) membrane developed by DuPont, is an important PEM because of its high proton conductivity, good chemical stability and high mechanical strength [5–7]. It consists of a carbon-fluorine backbone with perfluroro side chains containing sulfonic
acid groups. In light of the rapid development of the related technologies, the production and use of Nafion is expected to increase rapidly in the future. The disposal of Nafion wastes will therefore become more of a challenge, particularly because landfills and incineration are two common methods used for Nafion waste disposal [8].

Despite the widespread applications of Nafion in various technologies, recent studies focused primarily on its chemical degradation in fuel cells [9,10] and utilization in modified electrodes [11,12] as well as the removal of some pollutants [13,14], whereas the related toxicological data are lacking. The toxicity data provided by DuPont indicated an LD50 value greater than 20,000 mg/kg body weight in rats and no skin irritation to rabbits and humans [8]. It can be concluded that the potential toxic effects of Nafion are low. However, when Nafion is disposed via incineration, the toxicity of its combustion products may be increased because of the generation of fluorine ions (F−) and/or low-molecular-weight organofluorine compounds (S1 Table) [8]. To date, there are no toxicity data on these combustion products; therefore, additional studies are necessary to determine the better utilization and management of Nafion.

Recently, microarray-based toxicogenomics analysis has been used to evaluate the impacts of environmental toxicants on multiple cellular pathways [15–17]. This approach allows the identification of broad-scale gene expression patterns, which allows for the understanding of specific molecular responses and provides useful information on the mechanism of action of the toxicants [15]. However, this approach has not been used to assess the toxic effects of Nafion and its combustion products.

The objective of this study was to evaluate the potential toxicity of the Nafion 117 membrane (N117), a widely used Nafion membrane, and its combustion products on the mouse liver. Mice were exposed to three differentially treated N117 membranes for 24 days, and the histopathological alterations, oxidative stress responses, and transcriptome profiles were determined. Systematic analyses were conducted to identify the molecular pathways related to their potential toxicity. To the best of our knowledge, this is the first detailed assessment of the biotoxicity of N117 and its combustion products.

**Materials and Methods**

**Ethics statement**

All experiments involving animals were conducted in accordance with the National Institutes of Health (NIH) Guide for the Care and Use of Laboratory Animals (NIH publication number 85–23, revised 1996). All procedures were approved by the Institutional Animal Care and Use Committee of Model Animal Research Center of Nanjing University. Remarkable efforts were made to minimize the number of animals used in the study and their sufferings. All animals were housed in stainless-steel cages and acclimated for one week with regulated temperature (22 ± 1°C) and humidity (55 ± 5%), a 12/12 h light/dark cycle, and free access to food and water. At the end of the treatments, mice were fasted over night, anaesthetized with diethyl ether, and killed by cervical dislocation.

**N117 treatment**

To better mimic the actual exposure conditions of landfills or incineration sites, N117 (Shanghai Hesen Electric Co., Ltd, Shanghai, China) was cut into small pieces (1.0 mm × 1.0 mm) and then subjected to the following three different treatments: 1) directly mixed with mouse food; 2) combusted in the absence of oxygen with the products absorbed into the water to feed the mice; and 3) combusted in the presence of oxygen with the products absorbed into water to feed the mice. The details of the experimental treatments are described below.
1. Mixing of N117 with mouse food (1/100 wt% for N117/food): Mouse food, purchased from Qinglongshan Animal Breeding Center (Nanjing, China), was ground into a powder with mortar and pestle and then mixed with N117 pieces. After adequate mixing, a small amount of drinking water (30% v/w) was added to shape the mixture, which was then dried in an oven overnight at 103°C for the subsequent feeding test.

2. Combustion lacking oxygen supplementation (CLOS; 100 mg N117/L): Small pieces of the N117 (1 g) were heated in specialized pyrolysis equipment (AZ-HC-06a, Tianjin Aozhan Technology Co., Ltd, Tianjin, China). The operating parameters, including the atmosphere, heating temperature and time, were set according to the literature [8]. Specifically, the atmosphere was air, and the flow rate was 13 mL/min. The N117 sample was heated in a stainless steel tube at 10°C/min to 200°C and then for 5°C/min to 400°C; the temperature was then held for an additional 20 minutes for a total run time of approximately 75 minutes. The combustion products were absorbed with a 0.05 M sodium hydroxide (NaOH, 1 L) solution. The solution was then adjusted to pH 7.0 using 0.1 M hydrochloric acid, diluted 10-fold with drinking water and stored for the subsequent drinking test.

3. Oxygen-enriched combustion (OEC; 100 mg N117/L): Small pieces of the N117 (0.05 g) were ignited in a conical flask using the oxygen flask combustion (OFC) method (S1 File) [18,19], and the combustion products were absorbed with a 0.05 M NaOH solution (0.05 L). The solution was then adjusted to pH 7.0 using 0.1 M hydrochloric acid, diluted 10-fold with drinking water and stored for the subsequent drinking test.

Animal exposure
Twenty-four male mice (Mus musculus) (18–20 g), five-weeks of age, were purchased from Qinglongshan Animal Breeding Center (Nanjing, China). The mice were housed in stainless-steel cages and acclimated for one week with regulated temperature (22 ± 1°C) and humidity (55 ± 5%), a 12/12 h light/dark cycle, and free access to food and water. The mice were randomly divided into one control and three N117-treated groups, with six mice in each group. For the control group, a normal diet was provided for the mice daily. For the three experimental groups, one group received the N117-treated food and normal drinking water, and the other two groups received the normal food and treated water. They were exposed to these treatments for 24 days and named the Food group, CLOS group, and OEC group, respectively. After the exposure, all mice after diethyl ether anaesthesia were sacrificed by cervical dislocation for tissue collection.

F⁻ determination and LC/MS analysis
The tissue samples (liver, kidney, and muscle) were dissected from the mice, freeze-dried for 48 h with a Labconco Freeze Dry System (Labconco, Kansas City, MO, USA) and manually ground into powders, which were further passed through 0.4 × 0.4 mm screen. To quantify the fluorine concentration in the tissue samples, 0.05 g of each sample was first combusted using the OFC method with the gases absorbed in 50 mL 0.05 M NaOH solution. The solution was then adjusted to pH 7.0 using hydrochloric acid, followed by F⁻ analysis using a fluoride ion selective electrode (Shanghai Precision Scientific Instrument Co. Ltd., Shanghai, China) [18]. The F⁻ concentrations in the mouse urine were also measured with this electrode. For the drinking water samples, the F⁻ concentrations were determined using an ion-chromatograph system (ICS-1000, Dionex, USA) (S2 File).

The product composition of the initial absorption solutions of the different N117 combustion treatments was analyzed using liquid chromatography coupled with a high-resolution
hybrid quadrupole time-of-flight mass spectrometry LCMS-Q-TOF (LCMS-Triple TOF 5600, AB SCIEX, Foster City, CA) (S3 File).

**Histopathological analysis**
The liver tissue was harvested and fixed in 10% formalin, washed and dehydrated with a graded series of ethanol, and embedded in paraffin blocks. Sections of 5 μm thickness were stained with hematoxylin-eosin (H&E) and observed under a light microscope.

**Oxidative stress analysis and integrated biomarker response**
The liver samples were homogenized and centrifuged, and the supernatants were collected for biomarker determination. The biochemical parameters for oxidative stress, including the activities of superoxide dismutase (SOD) [20] and catalase (CAT) [21], reduced glutathione (GSH) content [22], and level of the lipid peroxidation product malondialdehyde (MDA) [23], were measured using commercial kits (Nanjing Jiancheng Bioengineering Institute, Nanjing, China) and normalized by protein content, which was determined by Bradford method [24]. Each experiment was performed in triplicate.

A method for combining all of the measured biomarker signals into one integral "stress index", termed the "Integrated Biomarker Response" (IBR) (S4 File) [25,26] was used to evaluate and compare the integrated impacts of N117 with different treatments on the antioxidant status of the mouse liver.

**RNA extraction and microarray analysis**
Total RNA from each mouse liver was extracted using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) and purified with a RNeasy mini kit (QIAGEN GmbH, Germany). The RNA concentrations were quantified using a NanoDrop ND-1000 spectrophotometer (Thermo Scientific, Wilmington, DE, USA), with the RNA integrity determined as the RNA Integrity Number (RIN) using an Agilent Bioanalyzer 2100 (Agilent Technologies, Santa Clara, CA, USA). RNA samples with a 2100 RIN value ≥ 7 and 28S/18S ≥ 0.7 were used for the gene expression analysis.

Microarray analysis was conducted by the Shanghai Biotech Corp (SBC, Shanghai, China). RNA extracted from one mouse liver, which was randomly selected in each group, was one sample. Four biological samples for all of the treatments were individually applied to the Agilent Whole Mouse Genome Oligo Microarray (4 × 44 K) platform (Agilent Technologies, Santa Clara, CA, USA) containing 41,174 unique probes. The subsequent RNA linear amplification and microarray hybridization were performed following the manufacturer’s instructions. After the hybridization, the microarray slides were scanned with an Agilent Microarray Scanner (Agilent Technologies, Santa Clara, CA, USA) at a 5 μm resolution for each slide with a photomultiplier tube setting of 100% and 10%. The data were extracted with the Feature Extraction software 10.7 (Agilent Technologies, Santa Clara, CA, USA). The raw data were normalized with the Quantile algorithm, Gene Spring Software 11.0 (Agilent Technologies, Santa Clara, CA, USA). Differentially expressed genes (DEGs) between the treated groups and control were identified as the genes with a greater than ± 2.0-fold-change and p-value < 0.05 (t-test). Gene ontology (GO) analysis (http://www.geneontology.org/) and Kyoto encyclopedia of genes and genomes (KEGG) pathway analysis (http://www.genome.ad.jp/kegg/pathway.html) were performed using the SBC Analysis System of the Shanghai Biotech Corp (http://www.ebioservice.com/).
Quantitative real-time polymerase chain reaction (QRT-PCR) analysis

Validation of the transcriptomic profiles was performed using QRT-PCR analyses with four target genes (Ugt1a2, Map3k6, Ccnb1 and Ccl5), which were differentially expressed in liver of mice exposed to N117 and its combustion products for 24 days. The primer sequences and PCR product sizes were shown in S2 Table. QRT-PCR analyses were conducted on all liver samples in each group (S5 File).

Statistical analysis

The data are expressed as the mean ± standard deviation (SD) and analyzed with the SPSS 16.0 software (SPSS Inc., Chicago, IL, USA). Tests for normal distribution (Kolmogorov-Smirnov) and homogeneity of variances (Levene) were applied. One-way ANOVA followed by Dunnett’s t test was performed to determine the significant differences of the biological parameters between the N117-treated groups and control. The results were considered significant at $p < 0.05$ and $p < 0.01$.

Results

Body and relative organ weight

Compared to the control, no significant differences ($p > 0.05$) were observed in the body and relative organ weights (liver and kidney) of the mice after a 24-day exposure to three different N117 treatments (S3 Table).

F⁻ analysis

The F⁻ concentrations in the mouse tissues, urine and drinking water samples in the control and N117-treated groups are listed in Table 1. Significant increases ($p < 0.01$) in the F⁻ concentrations were observed in the CLOS and OEC groups compared to the control. The highest value was found in the liver of the OEC group (1946.24 mg/kg dry weight), which was approximately 5.76 times that of the control. The F⁻ concentrations in the kidney, muscle and urine were approximately 2.45, 3.56, and 6.38 times greater in the OEC group than the control group, whereas a much higher F⁻ concentration was found in the drinking water for the OEC group (41.6 mg/L). The increases in the F⁻ concentrations in the CLOS group were less significant than the OEC group, and the F⁻ concentration in the drinking water for the CLOS group

Table 1. F⁻ concentrations in drinking water samples, mice tissues and urine in the control and N117-treated groups after 24-day exposure.

|                      | Control          | Food             | CLOS             | OEC             |
|----------------------|------------------|------------------|------------------|------------------|
| Drinking water (mg/L)| 0.68±0.05        | 0.65±0.08        | 8.45±0.59        | 41.6±1.51       |
| Liver (mg/kg dry weight) | 337.69±27.42    | 363.79±54.34    | 643.21±51.77**  | 1946.24±55.60**|
| Kidney (mg/kg dry weight) | 283.23±49.69    | 332.88±62.06    | 424.59±67.68    | 693.23±65.45**  |
| Muscle (mg/kg dry weight) | 282.97±23.61   | 378.48±61.52    | 771.19±39.52**  | 1008.80±38.66** |
| Urine (ng/L)         | 34.97±5.95       | 56.67±6.03      | 100.67±14.30**  | 223.00±15.62**  |

Values were obtained using an ion-chromatograph system (for drinking water) or a fluoride ion selective electrode (for mice tissues and urine). Twenty-four male mice (five-weeks of age) were individually exposed to normal diet (Control), 1/100 wt% N117-treated food (Food), 100 mg N117/L treated by combustion lacking oxygen supplementation (CLOS), and 100 mg N117/L treated by oxygen-enriched combustion (OEC) for 24 days, with six mice in each group. After the exposure, the drinking water samples, mice tissues (liver, kidney and muscle) and urine were assayed for F⁻ concentrations. Data are the mean ± standard deviation (SD), $n = 6$ for each data point.

** $p < 0.01$ indicates a significant difference compared with control using one-way ANOVA with Dunnett’s t test.

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was also lower than the OEC group was more elevated than the control (Table 1). In the Food group, only slight increases ($p > 0.05$) were found in the mouse tissues and urine compared to the control.

**LC/MS analysis**

As shown in S1 Fig, many significant $m/z$ peaks were observed in the absorption solution of the CLOS compared to the pure water, whereas only a few $m/z$ peaks were observed in the absorption solution of the OEC compared to the filter paper sample. Notably, many more $m/z$ peaks were recorded in the absorption solution of the CLOS than OEC. Furthermore, a more complete combustion of N117 from the OEC treatment led to less products but more $F^-$ than that from the CLOS treatment.

**Hepatic histopathology**

Histopathological alterations in the mouse liver after a 24-day exposure to the three N117 treatments are shown in Fig 1. No obvious effects were observed in the Food group, whereas
hepatocellular necrosis and inflammatory infiltration were obvious in the mouse livers of the CLOS and OEC groups.

**Oxidative stress induced by N117 and its combustion products**

The SOD and CAT activities, GSH content and MDA level in the control and N117-treated groups are shown in Fig 2. Compared to the control, the SOD activity was significantly inhibited \((p < 0.01)\) in the CLOS group. Significantly reduced CAT activity \((p < 0.01)\) was observed in the Food and CLOS groups, whereas a significant decrease \((p < 0.05)\) in the GSH content was detected in the OEC group. No significant increases \((p > 0.05)\) were detected for the MDA level in all of the treatment groups. For an overall evaluation of the antioxidant status, these signals were combined into one integral index, termed “IBR”. After these treatments, the IBR values ranged from 0.00 in the control to 5.94 in the CLOS group (Fig 3). The IBR of the treated groups was ranked CLOS > OEC > Food > Control.

Fig 2. Oxidative stress biomarkers in mice liver after a 24-day exposure to N117 and its combustion products. Antioxidant status was evaluated in mice liver by determining some oxidative stress biomarkers, and these assays were conducted using commercial kits (Nanjing Jiancheng Bioengineering Institute, Nanjing, China). Twenty-four male mice (five-weeks of age) were individually exposed to normal diet (Control), 1/100 wt% N117-treated food (Food), 100 mg N117/L treated by combustion lacking oxygen supplementation (CLOS), and 100 mg N117/L treated by oxygen-enriched combustion (OEC) for 24 days, with six mice in each group. The liver samples were dissected from the mice, homogenized and centrifuged, and the supernatants were collected for biomarker determination. The biochemical parameters for oxidative stress included superoxide dismutase (SOD; A), catalase (CAT; B), reduced glutathione (GSH; C), and the lipid peroxidation product malondialdehyde (MDA; D). Each experiment was performed in triplicate. Data are the mean ± standard deviation (SD), \(n = 6\) for each data point. The asterisks indicate significance of differences \(* p < 0.05; ** p < 0.01\) compared with the control using one-way ANOVA with Dunnett’s t test.

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Transcriptome profiles induced by N117 and its combustion products

The transcriptome responses of the mouse livers in the control and three N117-treated groups were analyzed using the Agilent Whole Mouse Genome Oligo Microarray. A total of 1370, 3013, and 2080 genes were identified as DEGs for the Food, CLOS, and OEC groups, respectively. The gene ontology analysis showed that the majority of these DEGs were associated with multiple biological processes, such as cellular process (GO: 0009987), biological regulation (GO: 0065007), and metabolic process (GO: 0006007) (Fig 4).

The biological significance of the DEGs for the different treatments was analyzed using the KEGG pathway database. A KEGG pathway was identified as significantly changed if there were more than three DEGs and the hypergeometric test yielded $p < 0.05$. The significantly altered KEGG pathways are listed in S4 Table and were used for further analysis. These differential pathways were divided into two primary categories, the metabolic pathway and the
Fig 4. Functional grouping of differentially expressed genes involved in biological processes. Gene expression profiling using Agilent Whole Mouse Genome Oligo Microarray platform containing 41,174 unique probes was performed in male mice (five-weeks of age) individually exposed to normal diet (Control), 1/100 wt% N117-treated food (Food), 100 mg N117/L treated by combustion lacking oxygen supplementation (CLOS), and 100 mg N117/L treated by oxygen-enriched combustion (OEC) for 24 days. RNA extracted from one mouse liver, which was randomly selected in each group, was one sample, and the obtained raw data were normalized with the Quantile algorithm, Gene Spring Software 11.0. Differentially expressed genes (DEGs) involved in biological processes between the treated groups (A, Food; B, CLOS; C, OEC) and control were identified as the genes with a greater than ±2.0-fold-change and p-value < 0.05 (t-test). Gene ontology (GO) analysis and Kyoto encyclopedia of genes and genomes (KEGG) pathway analysis were performed using the SBC Analysis System of the Shanghai Biotech Corp.

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biological process pathway. Different responses were observed after the different treatments (S4 Table), and the affected pathways involved 6 metabolic pathways and 6 biological processes, including carbohydrate metabolism, lipid metabolism, amino acid metabolism, xenobiotic biodegradation and metabolism, signal transduction, immune system and cellular process. For the Food group, 17 DEGs were involved in the metabolic pathway, including drug metabolism and retinol metabolism (Table 2). The CLOS group had 149 DEGs associated with the metabolic pathway, including drug metabolism, metabolism of xenobiotics by cytochrome P450, glutathione metabolism and retinol metabolism. For the OEC group, drug metabolism and glutathione and arachidonic acid metabolism were the significantly altered metabolic pathways, whereas the related cellular processes included focal adhesion, endocytosis, cell cycle and the p53 signaling pathway (Table 3). Several signal transduction pathways, such as the MARK and TGF-beta signaling pathways, were also significantly affected in the Food and OEC groups (Table 3). In addition, compared with the control, these N117 treatments significantly changed

### Table 2. Differentially expressed genes (DEGs) involved in metabolism pathways.

| Treatment | Pathway                                | Count | p-Value | Genes                                                                                   |
|-----------|----------------------------------------|-------|---------|-----------------------------------------------------------------------------------------|
| Food      | Retinol metabolism                     | 8     | 0.020   | Cyp26b1; Retsat; Cyp2a2; Cyp26a1; Cyp4a1; Ugt1a2; Cyp3a9; Cyp4a3                              |
|           | Drug metabolism                        | 9     | 0.019   | Cyp2a2; Gstm1; Gsta2; Fmo1; Ugt1a2; Maoa; Gstm3; Cyp3a9; Aox1                              |
| CLOS      | Pentose and glucuronate interconversions| 9     | 1.08×10^-4 | Ugt2b10; Akr1b1; Ugt2b17; Ugdh; Ugt2a3; Ugt2a1; Ugt1a1; Ugt1a2; Ugt2b37; Ugt2b15; Ugt2b   |
|           | Ascorbate and aldarate metabolism      | 10    | 4.22×10^-5 | Ugt2b10; Ugt2b17; Ugdh; Miox; Ugt2a3; Ugt2a1; Gulo; Ugt1a1; Ugt1a2; Ugt2b37; Ugt2b15; Ugt2b |
|           | Starch and sucrose metabolism          | 13    | 7.37×10^-4 | Ugt2b10; Hk1; Ugt2a3; Apl; Ugt2b15; G6pc; Ugt2b17; Gbe1; Ugdh; Ugt2a1; Gck; Ugt1a2; Ugt2b15; Ugt2b |
|           | Steroid hormone biosynthesis           | 16    | 6.38×10^-5 | Ugt2b10; Hsd3b6; Akr1d1; Hsd3b1; Ugt2a3; Cyp3a9; Ugt2b15; Cyp3a18; Srd5a1; Ugt2b17; Cyp11b3; Ugt2a1; Hsd11b2; Ugt1a1; Ugt2b15; Ugt2b |
|           | Glycine, serine and threonine metabolism| 13    | 1.57×10^-4 | Shmt2; Gmat; Gcat; Tdh; Pipox; Sardh; Phgdh; Alas2; Chdh; Gamt; Sds; Gldc                  |
|           | Glutathione metabolism                 | 18    | 3.06×10^-5 | Rm2; Gst1; Rm1; RGD1564906; Gpx2; Gsta5; Gsta4; Gclc; RGD1562107; Gstm2; Gpx3; Gstp1; Mgst3; Gstm1; Rm2b; RGD1558938; Gsta2; Gsto1; Sms; Gstm3 |
|           | Retinol metabolism                     | 17    | 9.63×10^-4 | Ugt2b10; Cyp1a2; Cyp2c7; Ugt2a3; Cyp4a1; Cyp3a9; Ugt2b15; Dhrs4; Cyp3a18; Ugt2b17; Cyp2a2; Ugt2a1; Rhdh5; Cyp2a5; Ugt1a2; Ugt2b37; Cyp4a3; Ugt2b |
|           | Drug metabolism                        | 29    | 2.75×10^-9 | Ugt2b10; Cyp1a2; Cyp2c7; Gst1; Gsta5; Gsta4; Fmo3; Cyp3a18; Ugt2b17; Mgst3; Ugt1a2; Ugt1a1; Cyp2a3; Ugt2b; RGD1564906; Ugt2a3; Cyp2d4; Cyp3a8; Ugt2b15; RGD1562107; Gstm2; Gsto1; Ugt2a2; Cyp2a1; Cyp2a5; Ugt2a1; Fmo1; Gstm3; Ugt2b37; Aox1 |
|           | Metabolism of xenobiotics by            | 24    | 1.01×10^-7 | Ugt2b10; Cyp1a2; Gst1; Cyp2c7; Gsta5; Gsta4; Cyp3a18; Ugt2b17; Mgst3; Ugt1a1; Ugt1a2; Ugt2b; Ephx1; Cyp2f4; RGD1564906; Ugt2a3; Ugt2b15; Cyp3a9; RGD1562107; Gstm2; Gsto1; Ugt2a1; Cyp2a1; Gsto1; Gstm2; Gsto3; Ugt2a2; Cyp2b3; Cyp3a9; Gpx3; Gsto1; Ugt2b15; Ugt2b37; Gstm3 |
|           | cytochrome P450                          |       |         |                                                                                          |
| OEC       | Arachidonic acid metabolism             | 12    | 0.030   | Cyp2b21; Gpx3; Tbxas1; Ggt6; Gpx2; Cyp2b3; Ggt5; Cyp4ff5; Gpx6; Cyp2b2; Cbr3; Ptgss1       |
|           | Glutathione metabolism                 | 13    | 6.55×10^-4 | G6pd; Rm2; Gpx2; Ggt5; Mgst2; RGD1562107; Gstm2; Gpx3; Ggt6; Mgst3; Gsta2; Gpx6; Gstm3 |
|           | Drug metabolism                        | 16    | 7.35×10^-4 | Aox2p; Cyp2b3; Fmo3; Cyp3a1; Mgst2; RGD1562107; Cyp2b21; Gstm2; Cyp2a2; Mgst3; Cyp3a1; Gsta2; Cyp2b2; Fmo1; Cyp2a3; Gstm3 |

Values were obtained using Agilent Whole Mouse Genome Oligo Microarray platform on mice liver. Twenty-four male mice (five-weeks of age) were individually exposed to normal diet (Control), 1/100 wt% N117-treated food (Food), 100 mg N117/L treated by combustion lacking oxygen supplementation (CLOS), and 100 mg N117/L treated by oxygen-enriched combustion (OEC) for 24 days, with six mice in each group. Differentially expressed genes (DEGs) involved in metabolism pathways between the treated groups and control were identified as the genes with a greater than ±2.0-fold-change and p-value < 0.05 (t-test). Gene ontology (GO) analysis and Kyoto encyclopedia of genes and genomes (KEGG) pathway analysis were performed using the SBC Analysis System of the Shanghai Biotech Corp.

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the pathways of immune system and signaling molecules and interaction in mice liver after a 24-day exposure, which mainly included cell adhesion molecules (CAMs), cytokine-cytokine receptor interaction, antigen processing and presentation, and chemokine signaling pathway (Table 4).

To validate the results of microarray analysis, four DEGs (Ugt1a2, Map3k6, Ccnb1 and Ccl5) were selected and verified by QRT-PCR. Taken together, the QRT-PCR data closely paralleled the gene expression pattern presented in the microarray data (S5 Table).

Discussion

The potential toxicity of N117 and its combustion products on mice following a 24-day exposure was assessed in this study using histopathological analysis, biomarker and microarray-based transcriptomics profile analyses. Notable, several significantly altered molecular pathways were identified, suggesting multiple toxicity mechanisms for N117 and its combustion products in mice.

Histopathological analysis

Histopathological changes indicate cellular damage when organisms are exposed to toxicants. The hepatocellular necrosis and inflammatory infiltration observed in the mouse liver in this study after exposure to the combustion products of N117 indicated their potentials to induce liver injury. As shown in S1 Table and S1 Fig, the possible byproducts under oxygen-lacking conditions included F⁻ and some low-molecular weight organofluorine chemicals, whereas F⁻ was the primary product under the oxygen-enriched conditions. Previous studies reported that F⁻ triggered histopathological alterations in the livers of mice [27] and rats [28] after in vivo exposure to different concentrations of sodium fluoride (NaF) in drinking water, which is consistent with the current study of the combustion products of N117.

Table 3. Differentially expressed genes (DEGs) involved in biological processes (signal transduction and cellular processes).

| Treatment | Pathway                          | Count | p-Value | Genes                                                                 |
|-----------|----------------------------------|-------|---------|----------------------------------------------------------------------|
| Food      | MARK signaling pathway           | 20    | 0.044   | Il1r1; Flnb; Fgfl21; Gadd45a; Ddit3; Chp2; Nf1; Dusp2; Nlk2; Map3k14; Map3k6; Mapk8; Jun; Nf1; Rela; Il1a; Map3k13; Cd14; Dusp7; Ntrk2 |
| OEC       | TGF-beta signaling pathway       | 15    | 0.011   | Tgfb1; Id3; Comp; Zfyve9; Id1; Acrvl1; Lefty2; MGC112830; Amhr2; Amh; Tnf; Inhba; LOC681309; Fst; Tgfb3 |
|           | Endocytosis                       | 25    | 0.047   | Ret; Adrb2; RT1-EC2; RT1-CE10; RT1-CE16; Amb2; RT1-CE7; Cxcr4; RT1-CE5; Pdgfra; Pdcd4; Col2a1; RT1-A3; RT1-CE2; RT1-M3-1; Rab31; RT1-CE4; RT1-CI; Hspa1a; Ldrlr; RT1-S3; RT1-A2; Pdcd2; Grk5; Mdm2; Sh3kbp1; Met; Csf1r |
|           | Cell cycle                        | 18    | 0.032   | Ccnb1; Tgfb1; Gadd45g; Pkmyt1; LOC298795; Cdk1; MGC112830; Mcm5; Esp1; Cdc25c; Ccnb2; Bub1b; Cdk1a1; Mdm2; Mcm2; Wee1; Tgfb3; Cdc20 |
| p53 signaling pathway | 11 | 0.046 | Ccnb1; Igfbp3; Gadd45g; LOC298795; Cdk1; Serpine1; Ccnb2; Shisa5; Cdk1a1; Mdm2 |
| Focal adhesion | 30 | 0.002 | Comp; Col5a2; Lama1; Mapk8; Acrn3; Jun; Col5a1; Vav1; Pdgfra; Col1a1; Fyn; Ppp1cb; Pik3cd; Itga7; Lama2; Pdgfrb; Lamb2; Rac2; Birc3; Vegfa; Itgb7; Prkcb; Vwfl; Itgα9; Mylk3; Col1a2; Flna; Met; Pak1; LOC681309 |

Values were obtained using Agilent Whole Mouse Genome Oligo Microarray platform on mice liver. Twenty-four male mice (five-weeks of age) were individually exposed to normal diet (Control), 1/100 wt% N117-treated food (Food), 100 mg N117/L treated by combustion lacking oxygen supplementation (CLOS), and 100 mg N117/L treated by oxygen-enriched combustion (OEC) for 24 days, with six mice in each group. Differentially expressed genes (DEGs) involved in signal transduction and cellular processes between the treated groups and control were identified as the genes with a greater than ±2.0-fold-change and p-value < 0.05 (t-test). Gene ontology (GO) analysis and Kyoto encyclopedia of genes and genomes (KEGG) pathway analysis were performed using the SBC Analysis System of the Shanghai Biotech Corp.

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Biomarker responses

Under normal conditions, reactive oxygen species (ROS) triggered by xenobiotic exposure are continuously decomposed in cells by antioxidant enzyme defenses, such as SOD and CAT, and low-molecular-mass scavengers, such as GSH [29,30]. Oxidative stress occurs in organisms when the steady-state ROS concentration is transiently or chronically enhanced during toxicant metabolism, leading to the increased damage to different cellular components [31]. In this study, cellular antioxidant defenses were significantly inhibited after the different treatments (SOD for CLOS group, CAT for CLOS and Food groups, and GSH for OEC group; Fig 2). These results suggested that N117 and its combustion products potentially induce oxidative stress in mouse livers. Although the toxicity data for the low-molecular-weight organofluorine compounds in the CLOS group are currently lacking, F\textsuperscript{−}-induced oxidative stress in rats or mice has been previously reported [32-34]. However, our data did not show any significant alteration in the MDA content after these various treatments, indicating the absence of lipid peroxidation during the 24-day exposure.

The IBR index was used to reflect the overall stress of the different treatments. This approach provides a simple but powerful tool for comparing the biological effects by combining different biomarker signals [25] and has been commonly used in recent environmental risk assessment studies [26,35]. The IBR index showed that the CLOS group exhibited a higher stress than the OEC group. This is most likely because the CLOS products contained both F\textsuperscript{−} and low-molecular-weight organofluorine compounds, whereas the OEC products were primarily composed of F\textsuperscript{−}. In addition, the IBR value for the Food group was higher than the control, suggesting the possible toxic effects of N117 via direct ingestion.

Transcriptomics analysis

After the 24-day exposure to three different treatments of N117, the altered KEGG pathways in the mouse liver were divided into two main categories, the metabolic pathway and biological process pathway. In this study, three major metabolic pathways, xenobiotic biodegradation and metabolism, carbohydrate metabolism, and lipid metabolism, and four biological processes, signal transduction, cellular processes, immune system, and signaling molecules and interaction, were selected for further analysis.

Xenobiotic biodegradation and metabolism was the most affected metabolic pathway for all three treatments, with 9, 53, and 16 DEGs in the Food, CLOS, and OEC groups, respectively. The majority of the DEGs involved in the detoxification of xenobiotics were associated with the aberrant regulation of cytochrome P450 enzymes (CYPs) and glutathione S-transferases (GSTs) [36], which is consistent with earlier studies of exposure of other toxicants [37,38]. As biomarkers for the toxicity of drugs and xenobiotics, CYPs and GSTs have been widely used to assess the potential toxicity of environmental pollutants [39,40]. Carbohydrate metabolism related pathways were also altered in the livers of the CLOS group, including pentose and glucuronate interconversion (9 DEGs), ascorbate and aldarate metabolism (10 DEGs), and starch and sucrose metabolism (13 DEGs). Of these DEGs, the down-regulated genes, including ugt1a1, ugt1a2, ugt2a1, ugt2a3, ugt2b, ugt2b10, ugt2b15, ugt2b17, and ugt2b37, were related to the regulation of UDP-glucuronosyltransferases (UGTs). Microsomal UGTs, primarily expressed in mammalian liver, are a family of isoenzymes that catalyze the transfer of UDP-glucuronic acid to endogenous and exogenous chemicals and/or their metabolites, rendering these substances more polar and facilitating their excretion into the bile and urine [41-43]. The alterations of the expression of these genes suggested that the combustion products of N117 under oxygen-lacking conditions might change the carbohydrate metabolism in the mouse liver at transcriptome level. In addition, for lipid metabolism, variations in the steroid hormone biosynthesis
(16 DEGs) and arachidonic acid metabolism (12 DEGs) were identified in the CLOS and OEC groups. These observations indicate that the N117 combustion products can disrupt lipid metabolism in mouse livers.

The microarray analysis also showed some significant alterations in signal transduction pathways, including the MARK signaling pathway (20 DEGs) in the Food group and the TGF-beta signaling pathway (15 DEGs) in the OEC group. The MARK signaling pathway is involved in various cellular functions, including cell proliferation, differentiation and migration, and couples intracellular responses to growth factor binding to cell surface receptors [44,45]. Aberrance of this pathway has been implicated in several human diseases, such as inflammation and cancers [36]. By contrast, the TGF-beta system functions via protein kinase receptors and Smad mediators to regulate many biological processes, including morphogenesis, embryonic development, immune regulation, wound healing and inflammation [46,47]. Alterations of the TGF-beta signaling pathway are involved in a broad range of pathologies, such as cancer, cardiovascular pathology, fibrosis and congenital diseases [46,48]. The DEGs in these two pathways were primarily down-regulated in this study, and the disruption of these DEGs

### Table 4. Differentially expressed genes (DEGs) involved in biological processes (immune system and signaling molecules and interaction).

| Treatment | Pathway                                      | Count | p-Value | Genes                                                                 |
|-----------|----------------------------------------------|-------|---------|----------------------------------------------------------------------|
| Food      | Cell adhesion molecules (CAMs)               | 15    | 0.009   | Jam2; RTI-T24-1; RTI-A3; Cd8a; Cd8b; RTI-M3-1; RTI-EC2; Cdln4; RTI-CI; RTI-M2; RTI-A2; RTI-CE10; RTI-CE16; RTI-DOa; RTI-Da |
|           | Antigen processing and presentation          | 14    | 2.08×10⁻⁷| Nfyα; Hspa5; RTI-T24-1; RTI-A3; Cd8a; Cd8b; RTI-M3-1; RTI-EC2; RTI-CI; RTI-M2; RTI-A2; RTI-CE10; RTI-CE16; RTI-DOa |
| CLOS      | Cell adhesion molecules (CAMs)               | 30    | 0.004   | Cdh4; Jam2; Cldn14; Mag; Igal; RTI-EC2; Cd40; Snp; Cd22; RTI-DMa; RTI-CE10; RTI-CE7; RTI-CE16; RTI-DOa; RTI-CE5; Cd226; Cadm3; RTI-DOb; RTI-Bb; RTI-A3; Cd8a; Cd8b; Slc3; RTI-Db1; RTI-CE4; RTI-CI; Igtb7; RTI-S3; RTI-A2 |
|           | Cytokine-cytokine receptor interaction       | 37    | 0.004   | Bmpr1a; Cxcr1; Tnfrsf18; Ltb; Csf1; Ili10; Ili13rb1; Cxcr3; Cd40; Ccl2; Acvrb; Iilr2; Ccl3; Tnfrsf12a; Lepr; Tnfrsf12; Il2rb; Ccl4; Il2ra; Il15; Cxcl2; Il1r1; Retl; Ghr; Cxcl1; Ccl5; Cxcl14; Amh; Tnf; Ili1a; Fli3; Tnfrsf14; Ili18; Kitig; Ccr2; Csf1r |
| OEC       | Antigen processing and presentation          | 24    | 1.49×10⁻⁷| Psme1; Hspa4; Klr3; Ili30; RTI-EC2; RTI-DMa; RTI-CE10; RTI-CE16; RTI-CE7; RTI-DOa; RTI-CE5; Cts; RTI-DOb; RTI-Bb; Tap1; RTI-A3; Cd8a; RTI-CE2; Cd8b; B2m; RTI-Db1; RTI-CE4; RTI-CI; RTI-S3 |
|           | Chemokine signaling pathway                  | 33    | 0.005   | Ccl24; Cxcr1; Gng13; Ptkacb; Gng5; Cxcr3; Stat4; Cx2; Arb2; Cxil7; Adcy5; Gng12; Cxcl; Rhoc; Vav1; Tiam1; Cxcl4; Stat1; Cx2; Pik3cd; Cxil; Rac2; Cxil5; Cxcl14; Prkcb; Cxil9; Rock2; Adcy4; Cxil6; Hck; Pak1; Cxir2; Cxil1 |
|           | Cytokine-cytokine receptor interaction       | 29    | 2.86×10⁻⁵| Mag; Igal; RTI-EC2; Cd40; Snp; Cd22; RTI-DMa; RTI-CE10; RTI-CE7; RTI-CE16; RTI-DOa; RTI-CE5; Cd226; RTI-DOb; RTI-Bb; Cdm11; RTI-A3; Cd8a; RTI-CE2; Cdbb; Sdc3; Cd34; RTI-M3-1; RTI-Db1; RTI-CE4; RTI-CI; Igtb7; RTI-S3; RTI-A2 |
|           | Cytokine-cytokine receptor interaction       | 29    | 0.004   | Tnfrsf25; Tnfrsf21; Tnfrsf18; Il6st; Ltb; Csf1; Ili10; Cxcr3; Cd40; Ccl2; Cxcr4; Prt; Ccl3; Pdgfra; Cxcl2; Retl; Tnfrsf4; Acvrf1; Pdgfb; Ccl5; Ccr1; Amhr2; Amh; Tnf; Fli3; Tnfrsf14; Met; Ccr2; Csf1r |
|           | Antigen processing and presentation          | 25    | 1.16×10⁻⁷| Klr3; Ili30; RTI-EC2; RTI-DMa; RTI-CE10; RTI-CE7; RTI-CE16; RTI-DOa; RTI-CE5; Cts; RTI-DOb; Tap1; Tap2; RTI-A3; Cd8a; RTI-CE2; Cdbb; RTI-M3-1; RTI-Db1; RTI-CE4; RTI-CI; Hspal1a; RTI-S3; RTI-A2 |
|           | Chemokine signaling pathway                  | 26    | 0.004   | Ccl24; Fgr; Cxcr3; Stat4; Ccl2; Amb2; Gnb4; Cxcr4; Ccl3; Vav1; Stat1; Cxcl2; Pik3cd; Cxil; Rac2; Rasgrf2; Cxil5; Ccr1; Ptkcb; Ccl9; Grk3; Prex1; Adcy4; Hck; Pak1; Cxil2 |

Values were obtained using Agilent Whole Mouse Genome Oligo Microarray platform on mice liver. Twenty-four male mice (five-weeks of age) were individually exposed to normal diet (Control), 1/100 wt% N117-treated food (Food), 100 mg N117/L treated by combustion lacking oxygen supplementation (CLOS), and 100 mg N117/L treated by oxygen-enriched combustion (OEC) for 24 days, with six mice in each group. Differentially expressed genes (DEGs) involved in immune system and signaling molecules and interaction between the treated groups and control were identified as the genes with a greater than ±2.0-fold-change and p-value < 0.05 (t-test). Gene ontology (GO) analysis and Kyoto encyclopedia of genes and genomes (KEGG) pathway analysis were performed using the SBC Analysis System of the Shanghai Biotech C

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demonstrates that N117 and its combustion products potentially affect the signal transduction pathways in mouse livers during a 24-day exposure.

For the cellular processes, several altered pathways were identified in the OEC group, including focal adhesion (30 DEGs), endocytosis (25 DEGs), cell cycle (18 DEGs), and the p53 signaling pathway (11 DEGs). These results indicate that the combustion products of N117 under oxygen-enriched conditions induce cytotoxicity in mouse livers, which may be caused by F⁻ toxicity because F⁻ is the primary component in the OEC products. Previous studies have also suggested the F⁻-induced cytotoxicity occurs in mouse embryonic stem cells [49] and cultured mouse osteoblasts [50]. Furthermore, the hepatocellular necrosis and inflammatory infiltration detected in our histological examination also reflected the toxic effects on cellular processes.

Moreover, these N117 treatments induced several significantly changed pathways of immune system and signaling molecules and interaction, such as CAMs, cytokine-cytokine receptor interaction, antigen processing and presentation, and chemokine signaling pathway. Interestingly, in this work, a subset of the DEGs identified in GO analysis (Ccl2, Ccl3, Ccl5, Ccr2 and Cxcl2) comprises part of the cytokine-cytokine receptor interaction and chemokine signaling pathway, which play a role in health and are essential in the communication between cells in the immune system [51]. These genes have been previously indicated to modulate the immunological and inflammatory responses in disease [52,53]. Besides, CAMs, which are important in a variety of physiological and pathological conditions related to interactions between cells and in measuring the specificity of cell-cell binding [54], were also significantly altered in mice liver after N117 treatments, indicating their possible toxic effects on this pathway.

Conclusions
Our data indicated that N117 and its combustion products induce histopathological damage and oxidative stress in mouse livers during the experimental exposure. Microarray-based transcriptomics profiling showed that these N117 treatments mainly disrupted the metabolism of xenobiotics, carbohydrates and lipids as well as some biological processes, including signal transduction, cellular processes, immune system, and signaling molecules and interaction. This is the first toxicity study to simulate mouse exposure to N117 and its combustion products via waste disposal processes, and these alarming results necessitate additional studies to evaluate the long-term toxic effects and elucidate the underlying cellular and molecular mechanisms of N117 toxicity on living organisms.

Supporting Information
S1 ARRIVE Guidelines Checklist.

S1 Fig. The total ion current (TIC) chromatograph (A1-D1) obtained by liquid chromatography coupled with a high-resolution hybrid quadrupole time-of-flight mass spectrometry (LCMS-Q-TOF) and the corresponding mass spectrum (retention time: 0.714–11.302 min) (A2-D2) subtracted by the mass spectrum (retention time: 0.323–0.610 min). (A) the pure water, (B) the absorption solution of N117 treated by combustion lacking oxygen supplementation (CLOS), (C) the absorption solution of filter paper using the oxygen flask combustion (OFC) method, (D) the absorption solution of N117 treated by oxygen-enriched combustion (OEC). Chromatographic separation was performed at a flow rate of 250 µL/min using a Thermo BDS Hypersil C₁₈ column (2.1 mm × 100 mm, particle size 2.4 µm) maintained at 40°C. The mobile phase was 0.3% formic acid in water and acetonitrile with an isocratic elution of 40:60 (v/v). Injection volume was 10 µL and elution time was 20 min for all samples. Mass
spectrometric analysis was carried out with a Q-TOF MS operating in a negative ion mode using an electrospray ion source.

(DOC)

S1 File. The experimental procedures of the oxygen flask combustion (OFC) method.
(DOC)

S2 File. Ion chromatography analysis for drinking water samples.
(DOC)

S3 File. LCMS-Q-TOF analysis for different combustion products of N117.
(DOC)

S4 File. The calculation procedures of the integrated biomarker response (IBR) index.
(DOC)

S5 File. The experimental procedures of quantitative real-time polymerase chain reaction (QRT-PCR) analysis.
(DOC)

S1 Table. The thermal degradation products of perfluorosulfonic acid copolymer.
(DOC)

S2 Table. Gene-specific primer sequences used for QRT-PCR.
(DOC)

S3 Table. The body weight (BW) and relative organ weight of mice in control and N117-treated groups after 24 days.
(DOC)

S4 Table. Categories of significantly altered KEGG pathways affected by N117 and its combustion products.
(DOC)

S5 Table. QRT-PCR validation of changes in selected genes identified by microarrays.
(DOC)

Author Contributions

Conceived and designed the experiments: MBF JW QGH ZYW. Performed the experiments: MBF JW ZYW. Analyzed the data: MBF JW. Contributed reagents/materials/analysis tools: JW SGY PS ZYW. Wrote the paper: MBF. Greatly contributed to the English language editing of the current manuscript: RJQ MH.

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