

Dietary advanced glycation endproducts (AGEs) increase their concentration in plasma and tissues, result in inflammation and modulate gut microbial composition in mice; evidence for reversibility

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Dietary advanced glycation endproducts (AGEs) increase their concentration in plasma and tissues, result in inflammation and modulate gut microbial composition in mice; evidence for reversibility

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ABSTRACT

Scope: Dietary advanced glycation endproducts (AGEs) are associated with negative biological effects, possibly due to accumulation in plasma and tissues and through modulation of inflammation and gut microbiota. Whether these biological consequences are reversible by limiting dietary AGE intake is unknown.

Methods and results: Young healthy C57BL/6 mice were fed a standard chow (n = 10) or a baked chow high AGE-diet (n = 10) (~1.8–6.9 fold increased protein-bound N^ε-(carboxymethyl)lysine (CML), N^ε-(1-carboxyethyl)lysine (CEL), and N^δ-(5-hydro-5-methyl-4-imidazolone-2-yl)-ornithine (MG-H1)) for 10 weeks or a switch diet with baked chow for 5 weeks followed by 5 weeks of standard chow (n = 10). We assessed accumulation of AGEs in plasma, kidney, and liver and measured inflammatory markers and gut microbial composition. After 10 weeks of baked chow, a substantial panel of AGEs were increased in plasma, liver, and kidney. These increases were normalized after the switch diet. The inflammatory z-score increased after the baked chow diet. Gut microbial composition differed significantly between groups, with enriched *Dubosia* spp. dominating these alterations. **Conclusion:** A high AGE-diet led to an increase of AGEs in plasma, kidney, and liver and to more inflammation and modification of the gut microbiota. These effects were reversed or discontinued by a diet lower in AGEs.

Abbreviations: AGEs, Advanced glycation endproducts; ASV, amplicon sequence variants; CEL, N^ε-(1-carboxyethyl)lysine; CML, N^ε-(carboxymethyl)lysine; CR, c-reactive protein; FDR, false discovery rate; GO, Glyoxal; IQR, interquartile range; LDA, linear discriminant analysis; LEfSe, LDA effect size; LOD, limit of detection; MG-H1, N^δ-(5-hydro-5-methyl-4-imidazolone-2-yl)-ornithine; MGO, methylglyoxal; PERMANOVA, Permutational multivariate analysis of variance; RCT, randomized controlled trial; rRNA, ribosomal RNA; 3DG, 3-deoxyglucosone.

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1. Introduction

Advanced glycation endproducts (AGEs) are abundantly present in processed food products and may contribute to beneficial effects on flavor, smell, and shelf life (Uribarri et al., 2010). These dietary AGEs are formed during the non-enzymatic reaction between free amino acids and reducing sugars, via the so-called Maillard reaction (Maillard, 1912). There is increasing evidence in humans and animals that consumption of dietary AGEs contribute to AGEs measured in plasma (Scheijen et al., 2018; Uribarri et al., 2007) and organs (Tessier et al., 2016) and that a diet high in dietary AGEs in humans has negative biological effects, such as low-grade inflammation, endothelial dysfunction, and insulin resistance (Nowotny, Schroter, Schreiner, & Grune, 2018). However, it is currently unknown whether AGE accumulation in tissues and the negative biological effects associated with a high AGE diet are reversible. Furthermore, how dietary AGEs are involved in the aforementioned biological effects remains poorly understood.

Dietary AGEs may exert their biological effects, at least partly, through modulation of the gut microbiota composition, which in turn is increasingly recognized to play a fundamental role in the pathophysiology of obesity, diabetes, and cardiovascular disease (Tilg & Moschen, 2014). Although there are some discrepancies, several animal studies have shown that a heat-treated diet high in dietary AGEs can modulate gut microbiota composition (Mastrocola et al., 2020; Qu et al., 2017; Seiquer, Rubio, Peinado, Delgado-Andrade, & Navarro, 2014; Snelson et al., 2021; Yang et al., 2020). However, the effects on inflammatory markers or increases in protein-bound and free AGE levels in plasma and/or tissues were not addressed in all of these studies. Furthermore, it is currently unknown whether changes in gut microbiota composition following a high AGE diet are reversible.

In light of the above, we hypothesized that mice fed a high dietary AGE diet for 10 weeks show increased blood and tissue AGEs, increased inflammatory markers, and different microbiota composition as compared to mice fed a standard dietary AGE diet. In addition, we studied whether changes following the high dietary AGE diet were reversible by implementing a switch after 5-weeks of high dietary AGE diet to the standard dietary AGE diet for 5 subsequent weeks.

2. Material and methods

2.1. Animal studies

To obtain a high AGE diet, a standard rodent chow (ssniff, Soest, Germany) was baked at 160 °C for two hours (hereafter referred to as “baked chow”).

9-week-old female C57BL/6/OlaHsd mice were randomly divided in three cages in groups of 10 (Fig. 1). After a one-week acclimatization period, the mice were divided in three groups fed either standard chow (n = 10), baked chow high in dietary AGEs (n = 10), or a “switch diet” (n = 10) for 10 weeks. In the switch diet group, mice were first fed the baked chow diet for 5 weeks, and subsequently the standard chow diet for 5 weeks. All diets were provided ad-libitum. Fecal and blood samples were collected after the 1-week acclimatization period. Blood samples were collected after 5 weeks of dietary intervention, while fecal samples were collected every 2.5 weeks. At the end of the 10-week dietary intervention, all mice were sacrificed by anesthetic overdose and plasma, liver and kidneys, and feces (from the rectum) were collected.

To test whether accumulation of AGEs in organs after a high dietary AGE diet are reversible, two extra groups of 10 mice were allocated to either 5 weeks of the standard chow diet or 5 weeks of the baked chow diet under the same conditions as mentioned above. At the end of the 5-week dietary intervention, all mice were sacrificed. All experiments were approved by the local ethical committee for animal experiments of Hasselt University and performed according to institutional guidelines (matrix 201503).

2.2. Advanced glycation end products and oxoaldehyde measurements

AGEs were measured in chow diets, plasma, liver, and kidney. AGEs in plasma were measured as this reflects uptake of dietary AGEs from the gastrointestinal tract. We chose to measure AGEs in liver and kidney as these are major organs and are highly susceptible to AGE accumulation and AGE-induced damage (Bettiga et al., 2019; Fernando et al., 2019). Livers and kidneys were homogenized using a Mini-bead beater homogenizer (Biospec) and 250 µL sodium phosphate buffer (0.1 M) supplemented with protease inhibitor (Roche) and 0.02% Triton-x (Sigma-Aldrich).

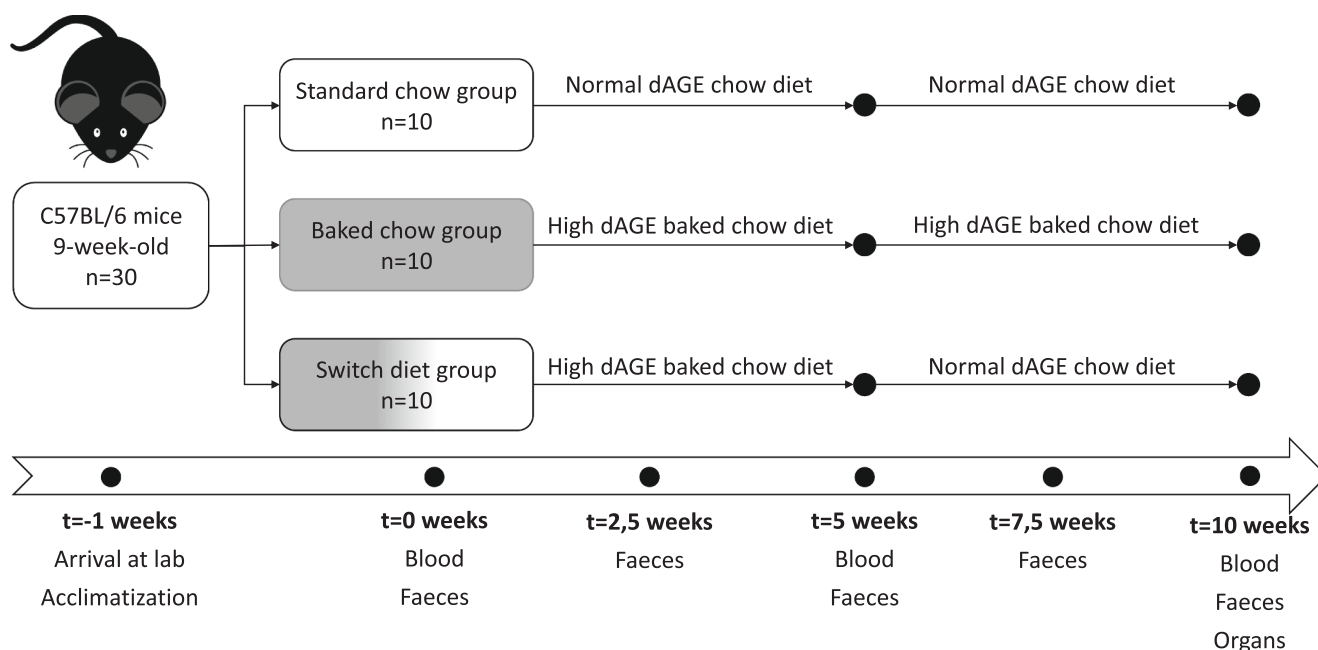


Fig. 1. Design of the main study. dAGE: dietary AGE.

Free- and protein-bound AGEs N ϵ -(carboxymethyl)lysine (CML), N ϵ -(1-carboxyethyl)lysine (CEL), and N δ -(5-hydro-5-methyl-4-imidazolone-2-yl)-ornithine (MG-H1) were analyzed in plasma, liver, and kidney homogenates by LC-MS/MS after extraction as described in detail before (Hanssen et al., 2014; Scheijen et al., 2016). Additionally, free-, protein-bound AGEs, and the oxoaldehydes methylglyoxal (MGO), glyoxal (GO) and 3-deoxyglucosone (3DG) were analysed in pulverized animal diets as described previously (Hanssen et al., 2014; Maassen et al., 2021).

2.3. Inflammation markers

IFN- γ , IL-10, IL-6, KC/GRO, and TNF- α were measured in plasma using MSD V-PLEX multiplex assay platforms (Meso Scale Diagnostics, Rockville, MD, U.S.A.). C-reactive protein (CRP) was measured using mouse CRP ELISA DuoSet kit (R&D system, Minneapolis, Minn, U.S.A.). For IFN- γ and IL-6, some samples were below the limit of detection (LOD) (<0.12 pg/ml for IFN- γ , n = 3, <3 pg/ml for IL-6, n = 14). These missing values were substituted by half the LOD, a commonly used method (Beal, 2001) (thus 0.06 pg/ml for IFN- γ and 1.5 pg/ml for IL-6).

2.4. Microbial 16S rRNA sequencing

DNA was isolated from the fecal pellets using a bead-beating procedure in combination with the customized MaxWell® 16 Tissue LEV Total RNA Purification Kit (XAS1220; Promega Biotech AB, Stockholm, Sweden) as described before (Fernández-Calleja et al., 2018). Triplicate PCR reactions of the 16S ribosomal RNA (rRNA) gene V4 region (515-F GTGYCAGCMGCCGCGGTAA, 806-R GGACTACNVGGGTWTCTAAT; 10 μ M each) were applied to the template DNA isolates (20 ng/ μ L) with a unique barcoded sequences library approach to identify individual fecal pellets with a total volume of 35 μ L per reaction. Formed PCR products were qualitatively confirmed and purified as described before (Fernández-Calleja et al., 2018). 200 ng of each sample was pooled after quantification with the Qubit® dsDNA BR Assay Kit and subsequently sequenced (Illumina NovaSeq 6000, paired-end, 150 bp; Eurofins Genomics Europe Sequencing GmbH, Konstanz, Germany).

The 16S rRNA gene sequencing raw data has been deposited in the European Nucleotide Archive (ENA) under accession number PRJEB41378.

2.5. Microbiota data processing and analysis

Sequences of the 16S rRNA gene were analyzed using the NG-Tax 2.0 pipeline (Poncheewin et al., 2020) with default settings, which generates *de novo* exact match sequence clusters: amplicon sequence variants (ASVs). ASVs with a relative abundance below 0.1% were removed and the threshold for taxonomic assignment was set at 80%. The SILVA 16S rRNA gene reference database release 132 (Quast et al., 2013) was used to assign taxonomy. The program R (version 3.6.1) was used for further data analysis. The Phyloseq package (McMurdie & Holmes, 2013) (version 1.30.0) was applied to combine the ASV table with the phylogenetic tree and the metadata. ASVs with a relative abundance > 0.1% in one of the individual samples were included for further data analysis. Bray-Curtis dissimilarities (beta-diversity) were assessed with the Phyloseq package. Relative abundance composition plots at phylum and genus level were created using the Microbiome package (Lahti & Shetty, 2017). The web-based tool Linear Discriminant Analysis (LDA) Effect Size (LEfSe) (Segata et al., 2011) was used to identify differential abundant taxa between the diet groups per sampling week. Spearman's rank correlations of microbial compositional data,glomerated at genus level, with relevant clinical data were performed with the associate function of the Microbiome package (Lahti & Shetty, 2017) (version 1.8.0). Therefore, subsets of each sampling week were created and taxa with a relative abundance > 1% in one of the samples were included.

2.6. Statistical analysis

Analyses were conducted using SPSS version 25 for Windows (IBM Corporation, Armonk, NY, USA). AGE and oxoaldehyde content of standard and baked chow are presented as mean \pm SD, and comparisons of AGE and oxoaldehyde content of standard and baked chow diets were tested using the two-tailed unpaired Students T-test.

Comparisons between free and protein-bound AGEs in plasma and tissues and inflammatory markers between groups were performed using the Mann-Whitney-U test. Free and protein-bound AGEs in plasma and tissues and inflammatory markers are presented as median [IQR] due to skewed distributions.

To increase statistical efficiency and reduce multiple testing when correlating inflammatory markers to the microbiota data, an inflammatory z-score was calculated by combining TNF- α , IFN- γ , KC/GRO, IL-6 and the inverse of IL-10 (1/IL-10) as described previously (Yudkin, Stehouwer, Emeis, & Coppack, 1999). To this end, first, z-scores for all individual parameters were calculated as follows: (individual value minus whole study population mean value)/ whole study population SD, thus resulting in a standardized variable ranging from approximately -2.5 to + 2.5 SD with a mean of 0. Second, as these individual z-scores share the same unit, they were averaged, resulting in one single inflammation score, which was subsequently standardized.

Permutational multivariate analysis of variance (PERMANOVA) of microbial beta diversity was assessed by application of the Adonis function (999 permutations) of Vegan package (Oksanen, Guillaume, Friendly, Kindt, Legendre, McGlenn, Minchin, O'Hara, Simpson, Solymos, Stevens, Szocs, & Wagner, 2019) (version 2.5-6). LEfSe analysis revealed differential abundant taxa if the P value of the non-parametric Kruskal-Wallis test between two diet groups was < 0.05 and the effect size of the logarithmic LDA score > 2.0. P values of correlations of microbial taxa with clinical data were adjusted for multiple testing using the Benjamini & Hochberg false discovery rate (FDR).

All analysis were considered statistically significant with P values < 0.05 unless otherwise stated.

3. Results

3.1. Increased levels of AGEs in chow after baking

To assess the effect of baking on chow diets, we compared free and protein-bound AGEs, and oxoaldehyde content between standard chow and baked chow. While levels of free CEL and MG-H1 increased after the baking procedure (34%, p < 0.05, and 266%, p < 0.01, respectively), free CML decreased after the baking procedure (-54%, p < 0.05) (Fig. 2). In contrast, levels of protein-bound AGEs in chow were uniformly and significantly increased following the baking procedure: 299% for protein-bound CML (p < 0.001), 691% for CEL (p < 0.001) and 182% for MG-H1 (p < 0.001) (Fig. 2). Likewise, levels of oxoaldehydes were also significantly increased following the baking procedure: 108% for MGO (p < 0.05), 53% for GO (p < 0.05) and 379% for 3DG (p < 0.001) (Fig. 2).

3.2. Mice fed baked chow ate less than mice fed standard chow, but weight gain was similar

To test whether weight gain over 10 weeks in young mice was different after the consumption of a baked chow diet, standard chow diet, or the switch diet, we determined food intake and weight at baseline and after 10 weeks. Mice fed the baked chow diet for 10 weeks ate less than mice fed the standard chow diet or switch diet: 2.96 g/day, 3.77 g/day, and 3.13 g/day, respectively (Table 1). In line with this, body weight was higher after 10 weeks of standard chow compared to baked chow or the switch diet, with median (g) [IQR] of 23.4 [22.2;24.1], 21.1 [20.5;21.2], and 21.8 [20.3;22.5], respectively. However, body weight of mice fed the baked chow diet was already

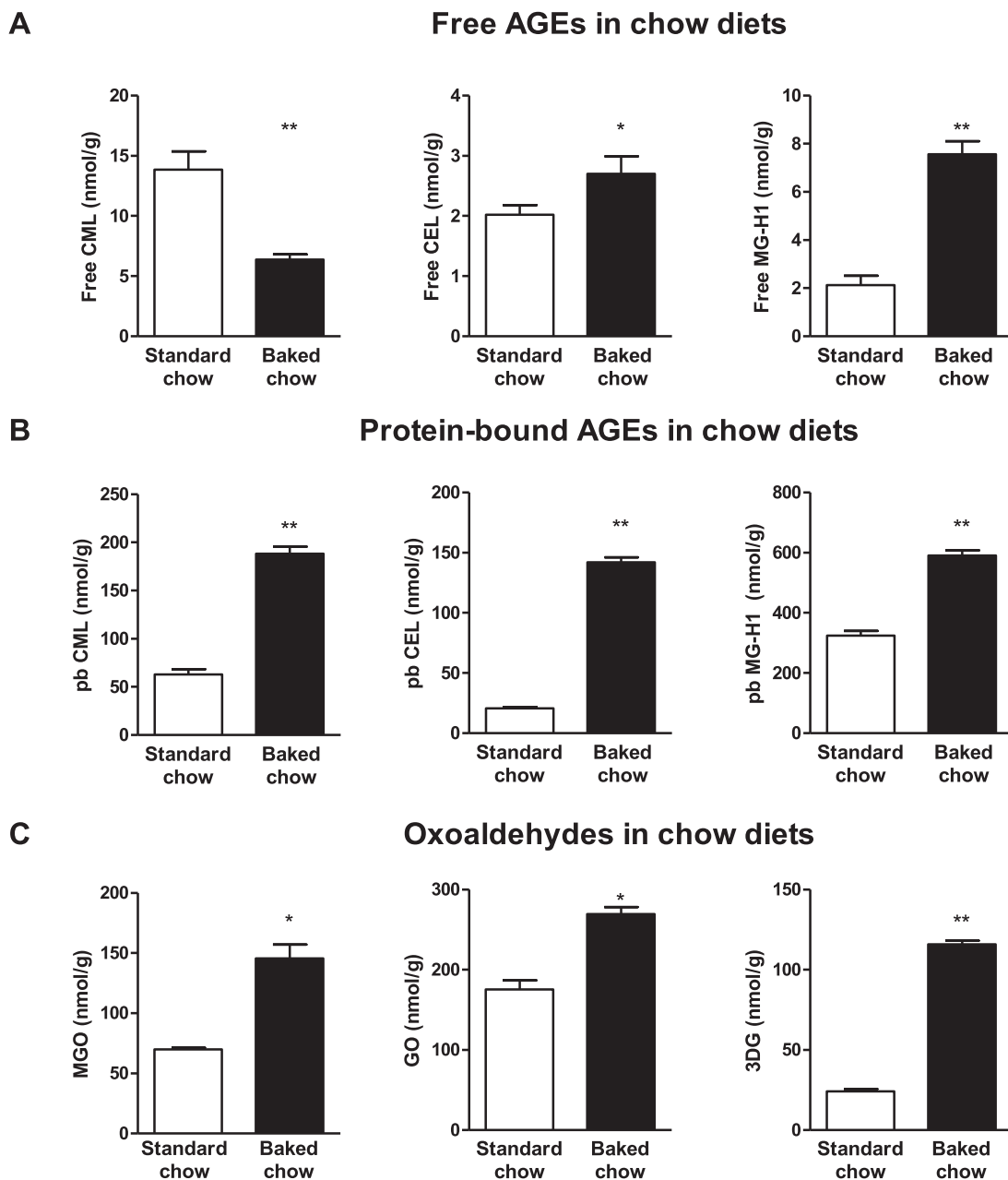


Fig. 2. Free AGEs (A), protein-bound AGEs (B), and oxoaldehydes (C) in standard chow and baked chow, as analyzed by UPLC-MS/MS. CML: N ϵ -(carboxymethyl) lysine. CEL: N ϵ -(1-carboxylethyl)lysine. MG-H1: N δ -(5-hydroxy-5-methyl-4-imidazolone-2-yl)-ornithine. MGO: methylglyoxal. GO: glyoxal. 3DG: 3-deoxyglucosone. pb: protein-bound. For free and protein-bound AGEs, bars and error bars indicate mean and SD of 5 samples per diet. For oxoaldehydes, bars and error bars indicate mean and SD of 2 samples per diet. * indicates $p < 0.05$ for the comparison between free and protein-bound AGEs and oxoaldehydes in baked chow and standard chow. ** indicates $p < 0.001$ for the same comparison.

Table 1
Mice characteristics.

	Standard Chow	Baked Chow	Switch Diet	<i>P</i> standard vs baked	<i>P</i> standard vs switch	<i>P</i> Baked vs switch
Weight baseline (g)	19.1 [17.9–19.5]	18.0 [17.1–18.2]	18.1 [16.4–19.3]	0.019	0.165	0.481
Weight sacrifice (g)	23.4 [22.2–24.1]	21.1 [20.5–21.2]	21.8 [20.3–22.5]	0.001	0.004	0.481
Weight gain (g)	4.2 [3.5–5.2]	3.3 [2.8–3.8]	3.1 [2.9–4.1]	0.043	0.035	0.796
Weight gain (%)	22 [19–27]	19 [15–22]	17 [16–23]	0.218	0.123	1.000
Food intake (g/day)	3.77	2.96	3.13			

Sample size $n = 10$ per group. Weight is presented as median [IQR]. Food intake was measured for each group as whole, thus statistical differences between groups cannot be computed. Significant differences ($p < 0.05$) are shown bold.

lower at baseline as compared to mice fed the standard chow diet, thus percentage weight gain was not different between mice fed standard chow, baked chow, or the switch diet after 10 weeks, with median (%) [IQR] of 22 [19;29], 19 [15;22], and 17 [16;23], respectively (Table 1).

3.3. A baked chow diet increases both free and protein-bound AGEs in plasma

To assess the impact of the chow diets on the level of AGEs after 10 weeks, we compared free and protein-bound AGEs in plasma between groups. All free AGEs in plasma were higher after baked chow compared to standard chow, with median (nmol/L) [IQR] of 536 [406;795] vs. 355 [332;430], $p = 0.004$ for CML, 228 [129;374] vs. 118 [86;135], $p = 0.011$, for CEL, and 242 [89;303] vs. 76 [61;102], $p = 0.019$ for MG-H1 (Fig. 3A).

Results for protein-bound AGEs in plasma were similar, except for MG-H1. After 10 weeks of the baked chow diet, protein-bound CML and CEL in plasma were higher compared to the standard chow diet, with median (nmol/L) [IQR] of 853 [805;919] vs. 561 [542;599], $p < 0.001$, and 311 [227;345] vs. 207 [183;235], $p = 0.011$, respectively. Protein-bound MG-H1 in plasma was lower after baked chow compared to standard chow: 1464 [1377;1506] vs. 1630 [1502;1911], $p = 0.011$ (Fig. 3B).

3.4. A baked chow diet increases AGEs in liver and kidneys, but mainly in the free form

We next determined the difference in AGE levels in organs after 10 weeks of the baked chow or standard chow diet. In liver, free CML and MG-H1, but not CEL, were higher after baked chow compared to standard chow, with median (nmol/g of protein) [IQR] of 8.0 [7.2;8.5] vs. 5.4 [4.9;5.7] for CML ($p < 0.001$), 2.3 [1.6;3.3] vs. 0.8 [0.8;0.9] for MG-H1 ($p < 0.001$), and 8.0 [7.4;8.7] vs. 8.1 [6.8;9.6] for CEL ($p = 0.730$) (Fig. 3C). Contrarily, protein-bound AGEs in liver were not different, although there was a trend for increased protein-bound CML in liver after baked chow compared to standard chow: 21 [20;22] vs. 18 [17;21] for CML ($p = 0.063$), 44 [39;46] vs. 41 [37;44] for CEL ($p = 0.393$), and 24 [21;28] vs. 25 [23;33] for MG-H1 ($p = 0.436$) (Fig. 3D).

In kidney, free AGEs were higher after baked chow compared to standard chow, although the difference in free CML did not reach statistical significance: 224 [143;284] vs. 161 [119;179] for CML ($p = 0.052$), 114 [67;147] vs. 48 [37;57] for CEL ($p < 0.001$), and 17 [8;26] vs. 5 [5;6] for MG-H1 ($p = 0.001$) (Fig. 3E). In contrast, protein-bound CML in kidney, but not CEL or MG-H1, was higher in kidney after baked chow compared to standard chow: 26 [24;29] vs. 22 [19;24] for CML ($p < 0.001$), 19 [18;22] vs. 21 [17;22] for CEL ($p = 0.796$), and 28 [22;44] vs. 31 [25;36] for MG-H1 ($p = 0.971$) (Fig. 3F).

3.5. The accumulation of AGEs in plasma and organs after a baked chow diet is reversible

We next studied whether the accumulation of AGEs after a baked chow diet can be reversed by a standard chow diet. We first assessed whether AGEs in plasma and organs were already increased after 5 weeks of the baked chow diet. In general, all free and protein-bound AGEs were increased in plasma, kidney, and liver of mice fed the baked chow diet for 5 weeks compared to mice fed the standard chow diet for 5 weeks (Supporting information Figure S1). However, protein-bound MG-H1 in plasma was not significantly different between mice fed the baked chow diet for 5 weeks compared to the standard chow diet, with median (nmol/g of protein) [IQR] of 1599 [1487;2128] vs. 1484 [1371;1704]. In addition, free CEL was lower and protein-bound MG-H1 was not different in liver of mice fed the baked chow diet compared to the standard chow diet: 8.3 [8.0;8.9] vs. 9.6 [8.8;10.4], $p = 0.007$ and 25.3 [21.5;28.9] vs. 26.5 [24.9;30.8], $p = 0.280$, respectively.

The accumulation of AGEs in plasma, kidney and liver after 5 weeks

of a baked chow diet were reversible, as most AGEs in plasma, liver, and kidney were significantly decreased after the switch to standard chow (Fig. 3A-F). Remarkably, free CML, free CEL, and protein-bound CEL in kidney were even lower after the switch diet as compared to 10 weeks of standard chow, with median (nmol/g of protein) [IQR] of 96 [85;102] vs. 161 [119;179], $p = 0.011$, 31 [29;33] vs. 48 [37;57], $p = 0.015$, and 16 [15;19] vs. 21 [17;22], $p = 0.009$, respectively (Fig. 3E and 3F).

3.6. A baked chow diet affects some markers of inflammation

To assess whether inflammation is increased after a baked chow diet compared to a standard chow diet, a panel of inflammatory markers were analyzed individually. TNF- α , IFN- γ , KC/GRO, CRP, and IL-6 were not different after 10 weeks of baked chow or standard chow (Fig. 4A). However, the anti-inflammatory cytokine IL-10 was significantly lower after baked chow compared to standard chow: 9.4 pg/ml [8.5;10.4] vs. 12.8 pg/ml [10.1;14.8], $p = 0.011$ (Fig. 4A).

We next combined the inflammatory markers into an overall inflammatory z-score that also included IL-10. The inflammatory z-score was higher after 10 weeks of baked chow than after 10 weeks of standard chow, with median (SD) [IQR] of 0.37 [-0.13;1.31] vs. -0.39 [-1.05;0.20], $p = 0.029$ (Fig. 4B).

Then we determined whether the increase in inflammation after a baked chow diet could be reversed by a standard chow diet. We first assessed whether inflammatory markers in plasma were already increased after 5 weeks of the baked chow diet. In contrast to AGEs in plasma and organs, inflammatory markers in plasma and the inflammatory z-score were not already increased after 5 weeks of baked chow compared to 5 weeks of standard chow (Supporting information Figure S2). In fact, plasma TNF α was lower after 5 weeks of baked chow compared to 5 weeks of standard chow, with median (pg/ml) [IQR] of 5.85 [5.16-6.39] vs. 7.50 [6.48;8.42], $p = 0.017$.

Although inflammatory markers in plasma were not already increased after 5 weeks of baked chow, the subsequent increase in inflammatory markers after 10 weeks of baked chow could be prevented by the switch to standard chow, with inflammatory z-score median [IQR] 0.37 [-0.13;1.31] vs. -0.41 [-0.81;0.26], $p = 0.034$ (Fig. 4B).

3.7. Gut microbial composition is altered after feeding a baked chow diet, and is reversible

Based on Bray-Curtis beta diversity dissimilarity distances, differences in overall microbial composition were shown in principal coordinate analysis (PCoA) plots from week 7.5 onwards (Fig. 5). Where in week 5 almost all samples were still distributed over the four quadrants (Axis 1 = 23.2%, Axis 2 = 13.5%), in week 7.5 the fecal samples from the mice fed the baked chow diet almost all clustered together in the bottom left quadrant, while almost all fecal samples from the standard chow diet clustered together in the upper right quadrant (Axis 1 = 23.6%, Axis 2 = 12%). The fecal samples from the mice fed the switch diet were distributed over the two upper quadrants and were located in between the samples of the baked chow and the standard chow diet. In week 10 the fecal samples of the mice fed the baked chow and the standard chow diet again were distributed over opposite quadrants, while the samples of the mice fed the switch diet were located in between these two diet groups, as expected (Axis 1 = 26.2%, Axis 2 = 23%). It should be noted that in week 10 not all fecal pellets were collected for each group ($n = 6$ for the baked chow group; $n = 4$ for the standard chow group; $n = 2$ for the switch group).

Further PERMANOVA analysis revealed that the provided diet explained a statistically significant ($p < 0.001$) variation in microbial composition from week 5 onwards with 19.5% variance explained by the diet in week 5, 22.9% in week 7.5 and 35.2% in week 10. Bray-Curtis dissimilarities of week 0 and week 2.5 were not statistically significant. Their corresponding PCoA plots are shown in the supplementary materials (Figure S3).

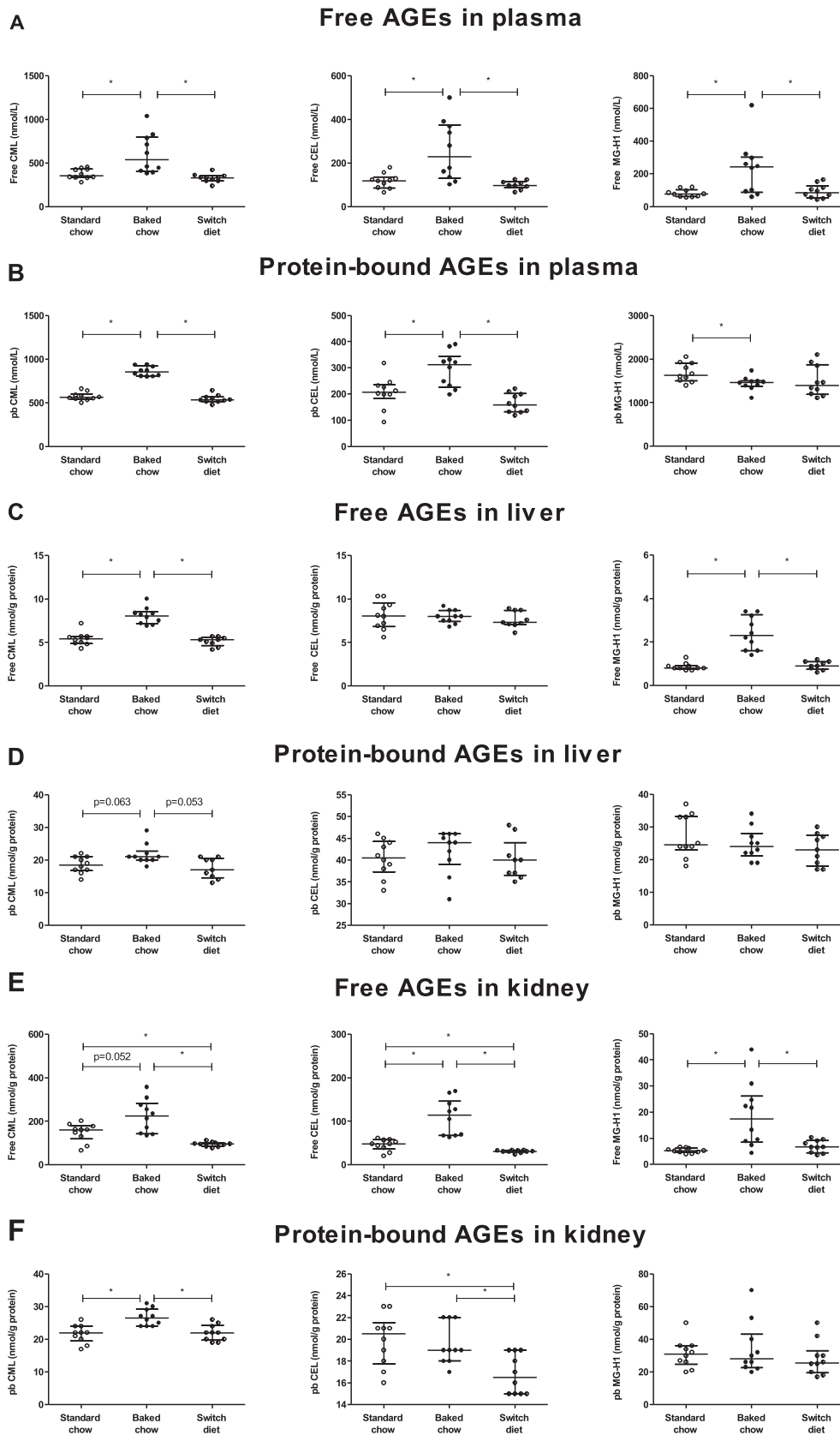


Fig. 3. Free and protein-bound AGEs in plasma (A,B), liver (C,D), and kidney (E,F) of mice after 10 weeks of standard chow, baked chow, and the switch diet, as analyzed by UPLC-MS/MS. CML: Nε-(carboxymethyl) lysine. CEL: Nε-(1-carboxyethyl)lysine. MG-H1: Nδ-(5-hydro-5-methyl-4-imidazolone-2-yl)-ornithine. pb: protein-bound. Center bars of scatter dots indicate median values while whiskers indicate inter quartile range. * indicates $p < 0.05$ for the difference between chow diets. $N = 10$ for all groups, except for free and protein-bound AGEs in liver for the switch diet group, where $n = 9$.

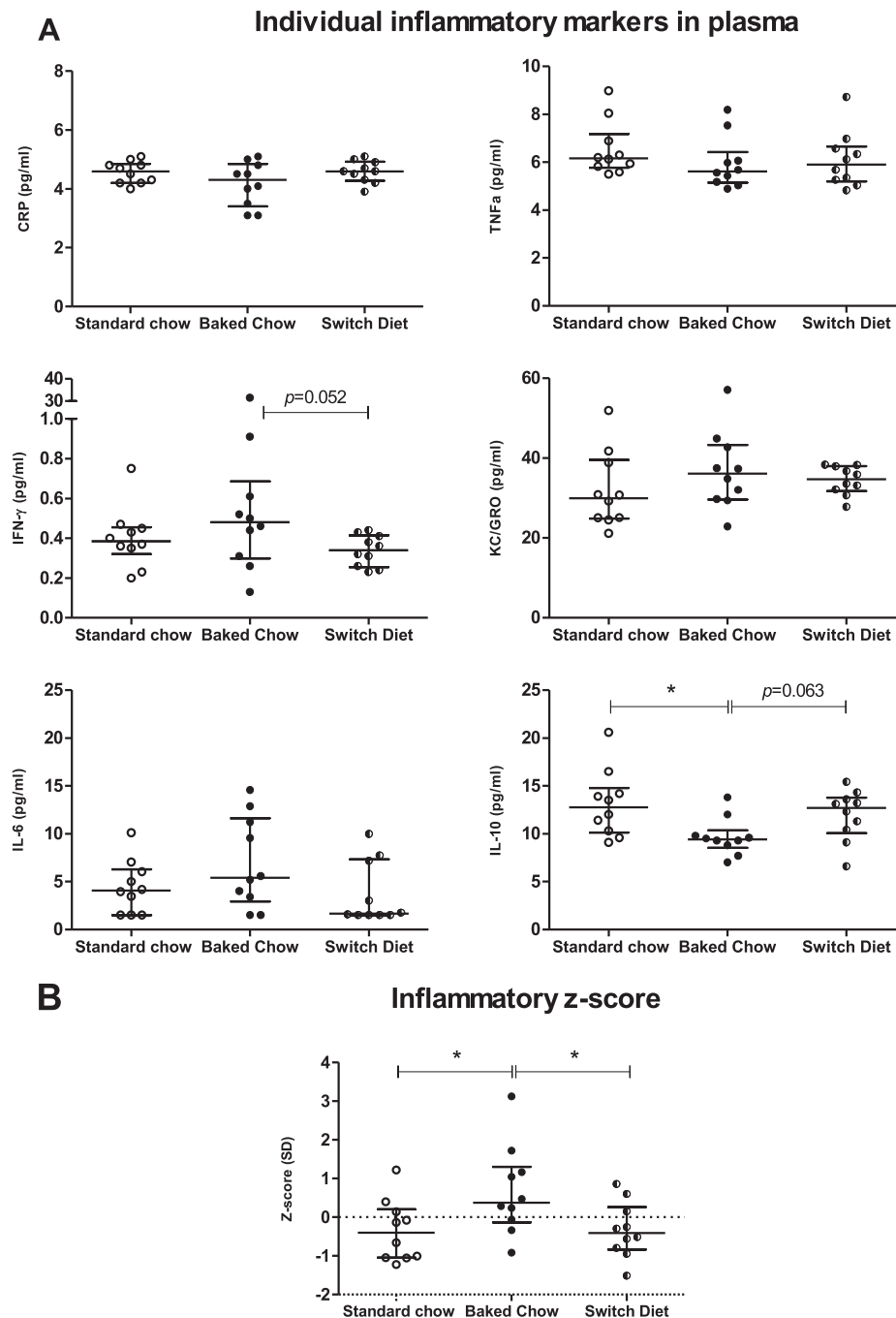


Fig. 4. Inflammatory mediators in plasma of mice after 10 weeks of standard chow, baked chow, and the switch diet. A: individual inflammatory mediators. B: Inflammatory z-score consisting of TNF- α , IFN- γ , KC/GRO, IL-6, and IL-10. Center bars of scatter dots indicate median values while whiskers indicate inter quartile range. * indicates $p < 0.05$ for the difference between chow diets. $N = 10$ for all groups.

Relative abundance composition plots both on phylum level and genus level per sampling week showed differences between the different diet groups (Figure S4), especially at genus level. In order to identify taxa which contributed to observed differences in microbial composition between the diet groups as assessed with Bray-Curtis dissimilarities, LefSe analysis of the microbiota composition was performed per sampling week from week 5 onwards. Differential abundant taxa were identified after comparing the microbial composition of the baked chow with the standard chow (Fig. 6) and the switch with the standard chow (Figure S5), and were visualized in cladograms (circular phylogenetic trees). Comparing the baked chow with the standard chow revealed that at week 5, six genera (i.e. *Olsenella*, *Ruminococcaceae_UCG_009*,

Dubosiella, *Turicibacter*, *Parasutterella* and *Akkermansia* with respective average relative abundances (%) of 0.7, 0.1, 3.6, 0.5 and 3.2) and four families (i.e. *Atopobiaceae*, *Erysipelotrichaceae*, *Burkholderiaceae*, *Akkermansiaceae*) were enriched after the baked chow diet and three genera (i.e. *Clostridiales_vadinBB60_group_uncultured*, *Roseburia* and *Faecalibaculum* with respective average relative abundances (%) of 0.1, 4.5 and 1.0) and one family (i.e. *Clostridiales_vadinBB60_group*) was enriched after consumption of standard chow. At week 7.5, twelve genera (i.e. *Bifidobacterium*, *Rikenella*, *Eubacterium_xylanophilum_group*, *Lachnoclostridium*, *Roseburia*, *Ruminiclostridium_5*, *Ruminococcaceae_uncultured*, *Ruminococcaceae_UCG_009*, *Ruminococcaceae_UCG_014*, *Dubosiella*, *Faecalibaculum*, *Desulfovibrio* with respective average relative abundances

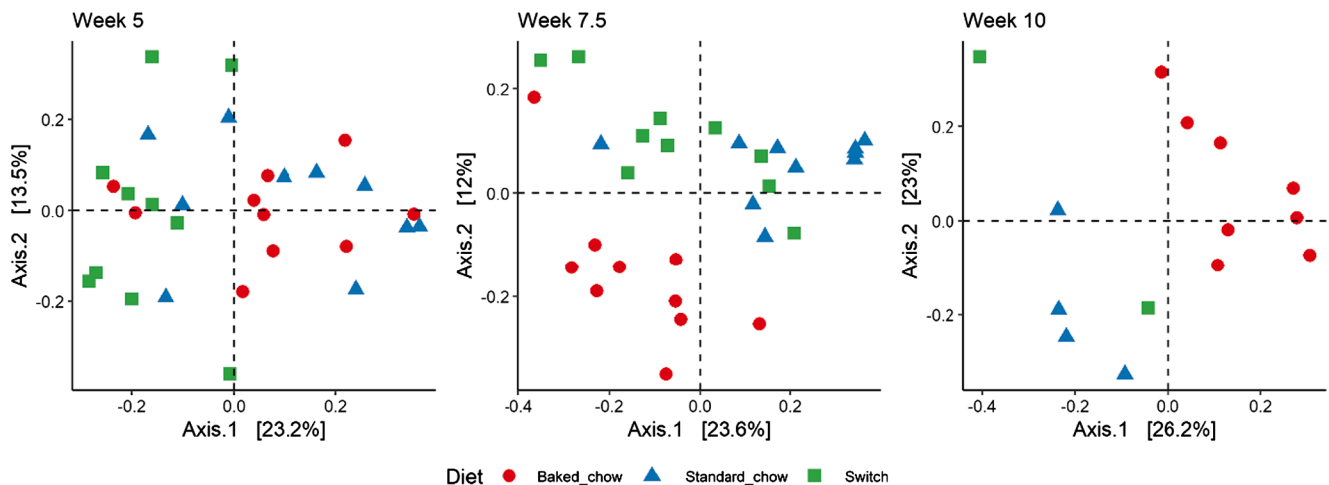


Fig. 5. Principal coordinate analysis plots (PCoA) of Bray-Curtis' beta-diversity dissimilarities of the fecal pellets per week. Each data point represents one fecal pellet of an individual mouse and labelled per diet (red circles for baked chow diet; blue triangles for the standard chow diet; green squares for the switch diet). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

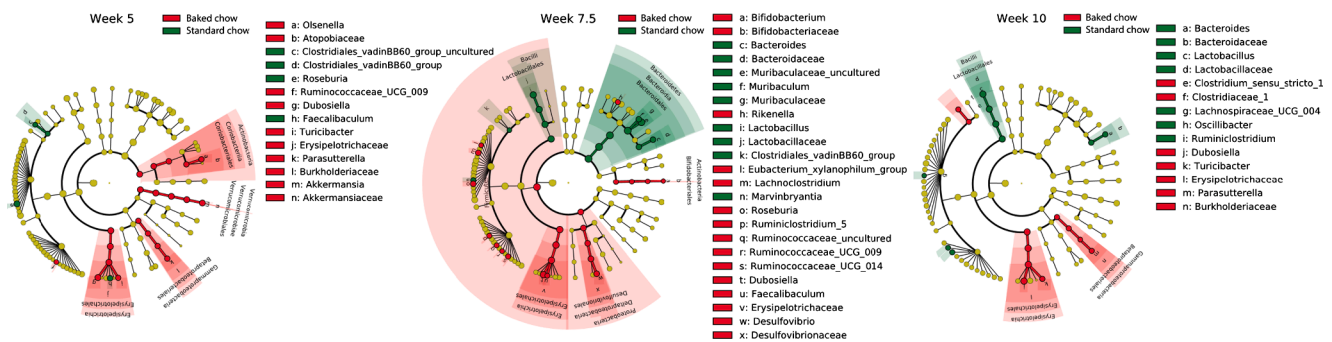


Fig. 6. LefSe results of the significant different taxa found by comparing the mice with the baked chow diet (red) to the standard chow diet (green), sampled in week 5 ($n = 10$ per group), 7.5 ($n = 10$ per group) and week 10 (baked chow: $n = 6$; standard chow: $n = 4$). Nomenclature was based on the highest achievable taxonomic resolution level. The alpha value was set to 0.05 and the \log_{10} LDA score threshold to 2.0. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

(%) of 0.2, 0.3, 0.1, 0.6, 1.5, 0.3, 1.1, 0.1, 0.3, 13.1, 0.2 and 1.7) and three families (i.e. *Bifidobacteriaceae*, *Erysipelotrichaceae* and *Desulfovibrionaceae*) were assigned as enriched taxa for the baked chow group, while for the standard chow diet five genera (i.e. *Bacteroides*, *Muribaculaceae_uncultured*, *Muribaculum*, *Lactobacillus* and *Marvinbryantia* with respective average relative abundances (%) of 1.2, 46.5, 1.2, 10.5 and 0.7) and four families (i.e. *Bacteroidaceae*, *Muribaculaceae*, *Lactobacillaea* and *Clostridiales_vadinBB60_group*) were enriched. The analysis performed at week 10 identified four genera (i.e. *Clostridium_sensu_stricto_1*, *Dubosiella*, *Turicibacter* and *Parasutterella* with respective average relative abundances (%) of 0.6, 26.2, 0.9 and 0.7) and three families (i.e. *Clostridiaceae_1*, *Erysipelotrichaceae* and *Burkholderiaceae*) to be enriched for the baked chow diet, while for the standard chow diet five genera (i.e. *Bacteroides*, *Lactobacillus*, *Lachnospiraceae_UCG_004*, *Oscillibacter* and *Ruminiclostridium* with respective average relative abundances (%) of 1.2, 22.5, 0.1, 0.4 and 0.7) and two families (i.e. *Bacteroidaceae*, *Lactobacillaceae*) were identified to be enriched.

The microbiota of baked chow-fed mice was found to be consistently enriched in the genus *Dubosiella*, while that of mice fed the standard chow diet were enriched in *Lactobacillus* and *Bacteroides* from week 7.5 onwards. The highest number of identified taxa after LefSe analysis were found in week 7.5 which could be explained by the fact that in week 10 not all fecal pellets were collected for each group ($n = 6$ for the baked chow group; $n = 4$ for the chow group; $n = 2$ for the switch group), while for week 7.5 fecal pellets from all mice ($n = 10$ per group) were collected

and subsequently analyzed for its microbial composition.

LefSe analysis of the switch diet with the standard chow diet (Figure S5) resulted in the highest number of enriched taxa in week 5 which decreased to week 7.5 and further to week 10. In week 10 only three genera (i.e. *Dubosiella*, *Turicibacter* and *Ileibacterium* with respective average relative abundances (%) of 2.9, 3.7 and 12.4) were shown to be higher abundant in the switch diet. This indicates that the microbial composition as affected by the baked chow diet is a diet-dependent reversible change.

3.8. Gut microbiota composition data are associated with clinical parameters

To assess whether gut microbiota composition was associated with other measured outcomes (i.e. AGE plasma and tissue levels, inflammatory markers), Spearman's rank correlations were determined after 5 and 10 weeks for mice fed the baked chow and the standard chow diet (Fig. 7). In week 5, the genera *Olsenella* and *Turicibacter* showed a statistically significant positive correlation with free plasma CML levels (*Olsenella*: $\rho = 0.769$, $p = 0.008$; *Turicibacter*: $\rho = 0.686$, $p = 0.030$) and CEL (*Olsenella*: $\rho = 0.711$, $p = 0.024$; *Turicibacter*: $\rho = 0.671$, $p = 0.031$). In week 10, no bacterial taxa were statistically significantly correlated with the other outcomes after correction for multiple testing, possibly due to less fecal pellets being collected at week 10. However, in week 10 several taxa showed a trend towards a high positive or negative

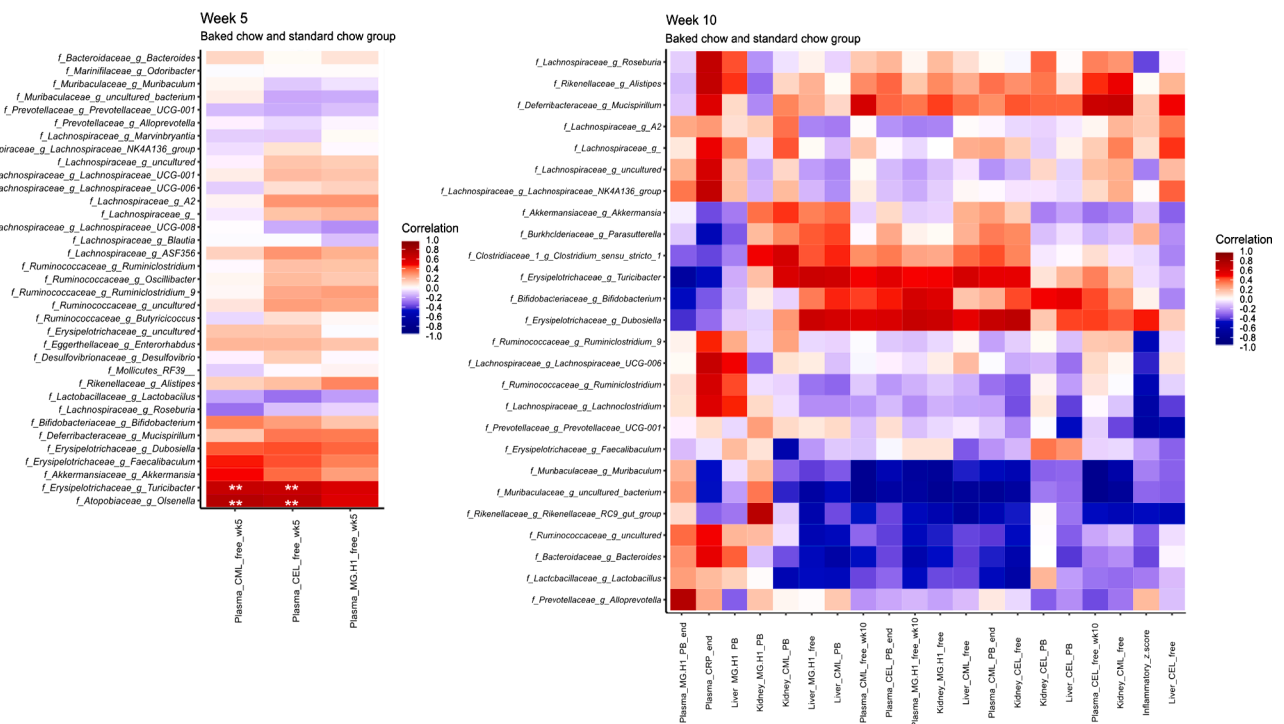


Fig. 7. Heatmaps of Spearman's rank correlation coefficient of relative abundance microbial composition data with relevant clinical parameters per sampling week, of mice in the baked chow and standard chow group. Bacterial taxa with a relative abundance > 1% in one of the samples were included. Statistically significant correlations after adjustment for multiple testing are marked with * ($p < 0.1$) or ** ($p < 0.05$). Nomenclature was based on the highest achievable taxonomic resolution level. pb: protein-bound.

correlation ($\rho > 0.7$ or $\rho < -0.7$) with these outcomes. As such, *Dubosiella* showed a high positive correlation with free kidney CEL levels ($\rho = 0.719$), *Lachnospiraceae* NK4A136 group with CRP levels in plasma ($\rho = 0.709$), *Alloprevotella* with protein-bound MG-H1 in plasma ($\rho = 0.755$), *Rikenellaceae* RC9 gut group with protein-bound MG-H1 levels in the kidney ($\rho = 0.745$) and *Alistipes* with plasma CRP levels ($\rho = 0.709$). On the other hand, *Rikenellaceae* RC9 gut group showed a high negative correlation with free CML in the liver ($\rho = -0.729$), *Muribaculaceae* uncultured bacterium with free CML and CEL in plasma ($\rho = -0.797$ and -0.825), protein-bound CML in plasma ($\rho = -0.755$), free CML in the liver ($\rho = -0.753$), and free CML and MG-H1 in the kidney ($\rho = -0.748$ and -0.783). *Muribaculum* showed high negative correlations with free CML and CEL in plasma, and free MG-H1 in plasma and kidney ($\rho = -0.797$, -0.825 , -0.718 , and -0.734 , respectively).

Since not all fecal pellets were collected at week 10, an additional Spearman's rank correlation analysis was performed using the microbial composition data of week 7.5 with other measured outcomes determined at week 10 (Figure S6). This revealed multiple statistically significant correlations. *Dubosiella* showed a positive correlation with free MG-H1 in liver ($\rho = 0.754$, $p = 0.018$), free CML in liver ($\rho = 0.683$, $p = 0.078$), protein-bound CML in plasma ($\rho = 0.796$, $p = 0.014$) and free CEL in kidney ($\rho = 0.771$, $p = 0.014$). *Desulfovibrio* showed a positive correlation with free MG-H1 levels in the liver ($\rho = 0.667$, $p = 0.098$). On the other hand, *Muribaculum* showed a negative correlation with free MG-H1 both in liver ($\rho = -0.777$, $p = 0.014$) and kidney ($\rho = -0.715$, $p = 0.040$). *Marvinbryantia* showed a negative correlation with protein-bound CEL in plasma ($\rho = -0.744$, $p = 0.021$). Although statistically significant results differed between Spearman's rank correlations using microbiome composition data from week 7.5 or week 10, the overall trend in found correlations was comparable.

Bacteroides and *Lactobacillus* showed a time-dependent increase from week 5 to week 10 towards negative correlations with free and protein-bound AGE levels in plasma and tissues, while *Dubosiella* showed a time-dependent increase towards positive correlations with free and protein-

bound AGE levels in plasma and tissues. These results are in line with the taxa identified in the LefSe analysis.

4. Discussion

Here, we show that consumption of a baked diet containing high levels of AGEs leads to an increase of AGEs in plasma, kidney, and liver and to a reduction of circulating IL-10 in addition to changes in gut microbiota composition in young mice. These effects were reversible or discontinued by switching the high AGE diet to a diet with lower levels of AGEs.

AGEs are generated in foods during various preparation methods involving dry heat, such as baking, grilling, frying, and toasting. Indeed, we saw a marked increase of the AGEs CML, CEL and MG-H1 after the baking of chow diet at 160 °C for two hours. Although food intake was lower in mice receiving the baked chow diet in comparison to the standard chow diet, total AGE intake was still increased in mice fed the baked chow diet. In line with this, free CML, CEL, and MG-H1 in plasma were higher in mice receiving the baked chow diet compared to mice receiving the standard chow diet. Although both free (CEL and MG-H1) and protein-bound (CML, CEL, and MG-H1) AGEs were increased in chow diets after baking, the uptake of dietary AGEs into the circulation is likely to depend mainly on digestion and absorption of protein-bound AGEs. It was previously shown that absorption of AGEs in vitro takes place mainly in the form of dipeptides, instead of free amino acid AGEs (Hellwig et al., 2011). In addition, levels of protein-bound AGEs in baked chow were approximately 30–80 fold higher than levels of free AGEs. In line with this, the increase in free plasma CML after the baked chow diet occurred in absence of an increase in free CML in baked chow.

Surprisingly, protein-bound CML and CEL in plasma were also higher, while protein-bound MG-H1 was lower in mice receiving the baked chow diet compared to mice receiving the standard chow diet. As such, we would not expect a direct increase in plasma protein-bound AGEs following the high AGE baked chow diet. This would be in line

with our previous data in humans showing that the free form of AGEs in plasma, but not the protein-bound form, is associated with dietary AGE intake in participants of the CODAM study (Scheijen et al., 2018). However, randomized controlled trials (RCTs) in humans are still inconclusive as there are reports of both an increase in protein-bound AGEs in plasma (Birlouez-Aragon et al., 2010) and no change (de Courten et al., 2016) after a high-AGE diet. This distinction is important, as the origin of both forms of AGEs is different, and it has implications for their biological effects, for example as free AGEs do not have affinity for the receptor for AGEs (RAGE) (Kislinger et al., 1999; Penfold et al., 2010; Xie et al., 2008).

We also observed some surprising findings in organs of mice fed the baked chow diet. While we observed higher levels of most free AGEs in liver and kidney in these mice, we also observed higher protein-bound CML in kidney. Direct evidence for dietary AGE accumulation in organs is already available in animal models, but free- and protein-bound AGEs have not been assessed simultaneously (Elmhiri et al., 2015; Roncero-Ramos et al., 2013; Somoza et al., 2006; Tessier et al., 2016). Tessier et al. used ¹³C-labeled CML to discriminate between dietary and endogenous CML and found that dietary CML accumulated in several organs of mice, but they could not discriminate between free and protein-bound CML due to low quantities of tissue being available (Tessier et al., 2016). The increase in protein-bound CML and CEL in plasma and protein-bound CML in kidney of mice fed the baked chow diet in the present study could be the result from indirect effects of dietary AGEs. It has previously been shown that dietary CML and MGO-modified albumin lead to significant increases of NF-κB and RAGE in kidneys of piglets and white adipose tissue respectively (Cai et al., 2012; Elmhiri et al., 2015). As such, protein-bound AGEs in plasma and tissues may reflect increased endogenous formation of AGEs, and not a direct uptake of dietary AGEs. In addition, we also measured reactive precursors of AGEs in the chow diet, which were increased upon heating. The increased levels of these reactive oxoaldehydes in baked chow diet may give rise to increased endogenous levels of protein-bound AGEs.

In line with the increase in AGEs in plasma and tissue, we observed reduced levels of circulating IL-10 in mice fed the baked chow diet as compared to mice fed the standard chow diet. Also driven by this, the inflammatory z-score was significantly higher after the baked chow diet compared to the standard chow diet. In line with our findings, Rajan et al. observed a significant increase in expression of several pro-inflammatory cytokines in 12-week old mice after a 6-month dietary intervention with baked chow diet (Sowndhar Rajan et al., 2018) and Mastrocola et al. also observed a decrease in IL-10 after feeding a high AGE diet for 22 weeks in 4-week old mice (Mastrocola et al., 2020). Although with the current data we cannot provide insights in the origin of increased inflammation following the baked chow diet, the increase in protein-bound CML in kidney is suggestive of involvement of the kidneys. Future studies could provide insight in a potential direct link between increased protein-bound levels in a specific tissue with the origin of observed inflammation. Nonetheless, the increase in inflammation after the baked chow diet may provide some insight into the mechanisms behind the biological effects of dietary AGEs. Circulating IL-10 levels are lower in obese insulin resistant individuals and treatment with IL-10 improves lipid-induced insulin resistance (Hong et al., 2009). As such, the decrease in circulating IL-10 after the baked chow diet could potentially contribute to the decrease in insulin sensitivity observed in humans after a high-AGE diet (de Courten et al., 2016). However, in humans, the increase in inflammation after a high-AGE diet is observed in some RCTs (Baye, Kiriakova, et al., 2017) but not all (Baye, De Courten, et al., 2017) and deserves further investigation.

To our knowledge, we are the first to show that the accumulation of AGEs in plasma and organs after a diet high in dietary AGEs is reversible. We showed that most AGEs in plasma, kidney, and liver were already increased after 5 weeks of the baked chow diet and that this accumulation of AGEs is fully reversible, by switching the diet to standard chow for 5 subsequent weeks. Although reversibility may be expected for

AGEs in plasma, as they are rapidly cleared by the kidneys (Förster & Henle, 2003), the latter does not necessarily apply to dietary AGEs accumulating in organs. We can currently only speculate on how dietary AGEs are transported into organs, if they end up in the intra- or extracellular matrix, and if glycated proteins (i.e. protein-bound AGEs) show different protein turnover rates. For example, the extracellular space is less subjected to protein turnover (Kerrigan, Mansell, & Sandy, 2000). Additionally, glycated proteins in food show resistance to enzymatic breakdown in the gastrointestinal tract (Pinto et al., 2014) and this could also potentially apply to enzymatic protein turnover. Inflammation, on the other hand, was not already increased after 5 weeks of the baked chow diet. Although mice fed the switch diet showed decreased inflammation compared to mice fed the baked chow diet for 10 weeks, we cannot speak of true reversibility. Nonetheless, dietary AGEs have been associated with negative biological effects in humans (Nowotny et al., 2018), but also several negative clinical outcomes, such as insulin resistance (de Courten et al., 2016), weight gain (Cordova et al., 2019), and vertebral fractures (Waqas et al., 2020). Therefore, the important finding of reversibility of AGE accumulation in plasma and organs may have beneficial implications for those whom ingest a diet high in AGEs to lower the dietary AGE-associated negative biological effects and clinical outcomes. Furthermore, the observation of reversible AGE accumulation after modulation of dietary AGEs also further strengthens their causal link.

Next to effects of baked chow diets on AGE levels in plasma and tissues and inflammatory markers, the baked chow diet also altered the gut microbiota composition of mice compared to the standard chow diet. From week 5 onwards significant effects of the baked chow diet on the gut microbiota composition were observed by assessing Bray-Curtis dissimilarities. Interestingly, statistically significant effects on gut microbiota composition were earlier observed (i.e. week 5) compared to effects on the inflammatory z-score and IL-10 levels in plasma (i.e. week 10). In the baked chow group, the genus *Dubosiella* was consistent enriched. *Dubosiella* was recently isolated from mice (Cox, Sohn, Tyrrell, Citron, Lawson, Pate, Iizumi, Perez-Perez, Goldstein, & Blaser, 2017), and has not been associated with specific functions before. However, *Dubosiella* belongs to the family *Erysipelotrichaceae* and was found to be related to *Allobaculum stercoricanis* (Cox et al., 2017). The genus *Allobaculum* spp. was increased in relative abundance in two animal studies after a glycated or high AGE diet (Han et al., 2018; Qu et al., 2017), which might imply a specific genera-cluster or family as a potential target for a high AGE diet. Interestingly, this does not corroborate with previous findings by Yang et al. (2020) who found an enrichment in the order *Erysipelotrichales* after the control diet instead of the high AGE diet. However, no specific genus was identified and it is relevant to note that old mice (15-month-old) were studied (Yang et al., 2020).

On the other hand, *Lactobacillus* and *Bacteroides* were consistent enriched in the standard chow group, corresponding to a relative decrease in the baked chow group, which corroborates partially or fully with earlier findings showing a contraction in *Lactobacillus* spp. or *Bacteroides* spp. (Delgado-Andrade et al., 2017; Mastrocola et al., 2020; Qu et al., 2017, 2018; Seiquer et al., 2014; Snelson et al., 2021). However, not all animal studies found this contraction in *Lactobacillus* spp. or *Bacteroides* spp. (Yang et al., 2020), indicating the need for multiple studies to derive clear and consistent conclusions regarding effects on gut microbiota composition.

In week 5, the genera *Olsenella* and *Turicibacter* showed a positive statistically significant correlation with free CML and CEL in plasma, while in week 10 no statistically significant correlations were found after correction for multiple testing. This could be explained by the higher number of parameters involved in the correlations of week 10 compared to week 5. Overall, *Dubosiella* showed high correlations with most parameters measured at week 10, and was included in statistical significant positive correlations when applying microbiome composition data from week 7.5 instead of week 10.

Our data suggests that *Dubosiella* spp. are associated with a diet high

in AGE levels. However, further work is required to identify the function or role of *Dubosiella* spp. in order to explain this link with AGEs. A hypothesized explanation of the enrichment of *Dubosiella* spp. in the baked chow diet group is the utilization of AGEs as a substrate by the bacterium. Multiple bacterial strains were reported in literature to degrade CML. For example *Cloacibacillus evryvenis* (Bui, Troise, Fogliano, & De Vos, 2019), isolated from human feces, and *Escherichia coli* strains (Hellwig, Auerbach, Mü, Samuel, Kammann, Beer, Gunzer, & Henle, 2019) were shown to degrade CML in vitro, which proves the ability of specific gut bacteria to utilize an AGE.

An attempt was made to investigate whether the effects of the baked chow diet on gut microbial composition were reversible by including the switch diet group. The results of both the Bray-Curtis dissimilarities and LEfSe analysis indicated that the trend of changing microbiota composition due to the baked chow diet is reversible. In week 10, LEfSe analysis identified 3 genera (i.e. *Dubosiella*, *Turicibacter* and *Ileibacterium*) to be enriched in the switch diet group. These findings suggest that minor differences in microbiota composition between the standard chow and the switch diet were still observed, which can be explained by the fact that the gut microbiota requires time to adapt towards a new diet composition. It could be that a 5-week dietary intervention of the standard chow diet after 5 weeks of the baked chow diet was too short to completely adapt and consequently forming a comparable microbiota composition as in the 10-week standard chow diet group.

Our study has several strengths. Protein-bound and free AGEs were measured both in the chow diets as well as in plasma and organs of mice with the highly specific gold standard UPLC-MS/MS technique. To our knowledge, we are the first to concurrently measure both forms of AGEs in both the experimental diet and tissue compartments. In addition, the validity of our findings is increased by inclusion of a control group and the switch group. Also, inclusion of the switch group enables us to show that changes in AGE accumulation following a high AGE diet are reversible in nature. Moreover, it is important to highlight that the chow baking conditions were not substantially different from the conditions used in other studies, and while AGE levels in the baked chow diet were increased, they still represented realistic dietary levels. Finally, we used very young, healthy mice without any underlying disease.

Our study also has several limitations. Primarily, we cannot rule out that the baking procedure has had effects on chow other than increasing AGEs, such as decreasing vitamin bioavailability or increasing acrylamide formation. Additionally, mice weighed less at the end of the study after the baked chow diet compared to the standard chow diet, and food intake was lower of mice in the baked chow diet group. However, percentual increase of weight did not differ between groups, and in addition bacterial taxa associated with weight loss, such as *Allobaculum* (Ravussin et al., 2012) or *Akkermansia muciniphila* (Everard et al., 2013), were not showing a similar trend towards lower weight in the baked chow diet group. If anything, lower food intake in the baked chow group reduced exposure to dietary AGEs and lead to underestimations of their effects. Finally, our observations of reversible AGE accumulation in kidney and liver may not be extrapolated to other organs. Likewise, our findings in mice cannot be directly extrapolated to humans, as species differences exist in for example metabolic rate and dietary habits, as also in gut microbiota composition (Hugenholtz & de Vos, 2018).

In summary, intake of dietary AGEs results in reversible elevated levels of AGEs in plasma and organs and alterations in the gut microbiota composition with a change in the inflammatory profile of healthy young mice. Randomized controlled trials on the effects of dietary AGEs on gut microbiota in humans are needed.

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CRediT authorship contribution statement

Katja C.W. Dongen: Formal analysis, Writing - original draft, Writing - review & editing, Visualization. **Armand M.A. Linkens:** Formal analysis, Writing - original draft, Writing - review & editing, Visualization. **Suzan M.W. Wetzels:** Investigation, Writing - review & editing, Project administration. **Kristiaan Wouters:** Conceptualization, Investigation. **Tim Vanmierlo:** Conceptualization, Investigation, Resources, Funding acquisition. **Marjo P.H. Waarenburg:** Methodology, Validation, Formal analysis. **Jean L.J.M. Scheijen:** Methodology, Validation, Formal analysis. **Willem M. Vos:** Resources, Writing - review & editing, Supervision, Funding acquisition. **Clara Belzer:** Resources, Writing - review & editing, Supervision, Funding acquisition. **Casper G. Schalkwijk:** Conceptualization, Resources, Writing - review & editing, Supervision, Funding acquisition.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary material

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.foodres.2021.110547>.

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