Mitochondrial Genetic Variants Identified to Be Associated with BMI in Adults

Antònia Flaquer¹,², Clemens Baumbach¹,², Jennifer Kriebel³, Thomas Meitinger⁴, Annette Peters³,⁵, Melanie Waldenberger³,⁵, Harald Grallert³,⁵, Konstantin Strauch¹,²

¹ Institute of Medical Informatics, Biometry and Epidemiology, Chair of Genetic Epidemiology, Ludwig-Maximilians-Universität Munich, Neuherberg, Germany, ² Institute of Genetic Epidemiology, Helmholtz Zentrum München - German Research Center for Environmental Health, Neuherberg, Germany, ³ Research Unit of Molecular Epidemiology, Helmholtz Zentrum München - German Research Center for Environmental Health, Neuherberg, Germany, ⁴ Institute of Human Genetics, Helmholtz Zentrum München-German Research Center for Environmental Health, Neuherberg, Germany, ⁵ Institute of Epidemiology II, Helmholtz Zentrum München-German Research Center for Environmental Health, Neuherberg, Germany

Abstract

It has been suggested that mitochondrial dysfunction plays a role in metabolic disorders including obesity, diabetes, and hypertension. The fact that mitochondrial defects can be accumulated over time as a normal part of aging may explain why some individuals can eat all sorts of foods and remain at normal weight while they are young. However, around the fourth decade of life there is a trend towards "middle-age spread" with weight gain and the body's decreasing ability to metabolize calories efficiently. To test the hypothesis that mitochondrial variants are associated with BMI in adults, we analyzed a total number of 984 mitochondrial single nucleotide polymorphisms (mtSNPs) in a sample of 6,528 individuals participating in the KORA studies. To assess mtSNP association while taking heteroplasmy into account we used the raw signal intensity values measured on the microarray and applied linear regression. Significant results were obtained for 2 mtSNPs located in the Cytochrome c oxidase subunit genes (MT-CO1: P adjusted = 0.0140 and MT-CO3: P adjusted = 0.0286) and 3 mtSNPs located in the NADH dehydrogenase subunit genes (MT-ND1, MT-ND2 and MT-ND4L: P adjusted = 0.0286). Polymorphisms located in the MT-CO3 and MT-ND4L genes have not been associated with BMI or related phenotypes in the past. Our results highlight the importance of the mitochondrial genome among the factors that contribute to the risk of high BMI. Focusing on mitochondrial variants may lead to further insights regarding effects of existing medications, or even to the development of innovative treatments.

Introduction

Excess of weight is a major risk factor for mortality and morbidity from cardiovascular disease [1], type 2 diabetes [2], and incident cancer [3]. Three million deaths per year worldwide are attributed to overweight and obesity in total [4]. Obesity is a medical condition in which excess body fat has accumulated to the extent that it may have an adverse effect on health [5]. Anyway, the conflation of lean and fat body mass becomes inconsistent in the elderly with both muscle mass and strength beginning to decline as part of aging [6]. Changes such as accumulation of intra- and extra-myocellular lipids, improper folding of structural and contractile proteins, and mitochondrial dysfunction are thought to occur with age [7–9]. Dysfunctional mitochondria are thought to play a key role in function decline with aging, as the mitochondria are the main producers of both cellular energy and free radicals.

Mitochondria are membrane-bound organelles found in the cytoplasm of almost all eukaryotic cells. Their primary function is to generate large quantities of energy in the form of adenosine triphosphate (ATP). In the early 1960s it was discovered that mitochondria contain their own DNA (mtDNA) of approximately 16.6 kb. It is known to code for 13 subunits of the mitochondrial respiratory chain complexes, 2 ribosomal (rRNA) genes, and 22 transfer RNA (tRNA) genes that are required for mitochondrial protein synthesis. Mitochondria consume oxygen and substrates to generate the vast majority of ATP while producing reactive oxygen species (ROS), also known as free radicals, in the process. An excess of ROS may damage DNA, proteins, and lipids if not rapidly quenched. This damage is termed oxidative stress. It has been recently recognized that oxidative stress is increased (ROS overproduction) in obesity [10,11] and in fat accumulation in adipocytes [12]. The 13 structural genes are essential for energy production through the process of oxidative phosphorylation.
(OXPHOS) and generation of ATP. The activity of the OXPHOS system that consists of five enzyme complexes (I–V) can be affected by a number of factors including age and physical training [13]. In addition to supplying ATP and involvement in oxidative stress, mitochondria also participate in other cellular processes, including signal transduction, cell cycle regulation, thermogenesis, and apoptosis. mtDNA is maternally inherited and lacks recombination [14]. One major difference between mtDNA and nuclear DNA (nDNA), which is relevant for the understanding of human diseases, is the heteroplasmy effect, which was originally believed to be a rare phenomenon. Because a cell carries many mitochondria, and also the mitochondrial genome has a higher mutation rate than the nuclear genome, there is heterogeneity of the relative frequency of a mtDNA variant within an individual tissue, cell, and even within the same mitochondrion. Therefore, mitochondria are considered heteroplasmic. The clinical expression of some phenotypes is determined by the relative proportion of normal and mutant mitochondrial genomes in different tissues. Mutations of mtDNA are under a growing scientific spotlight and there is increasing evidence that these mutations play a central role in many, if not most, human diseases.

In the past years, genome-wide association studies (GWASs) have identified over 90 common genetic variants associated with BMI in adult samples [15–19]. A growing body of research is demonstrating that altered mitochondrial energy production is a major anomaly capable of setting off a chain of metabolic events leading to obesity [20–23]. It has also been suggested that particular mtDNA haplogroups might be associated with inefficient energy expenditure [24] and obesity [25]. A mtDNA haplogroup is a collection of mtSNPs at certain genetic loci accumulated throughout human history that could be attributed to genetic drift and/or climate selection [26]. The fact that mitochondrial defects can be accumulated over time as a normal part of aging may explain why some individuals can eat all sorts of foods and remain at normal weight while they are young. However, around the fourth decade of life there is a trend towards “middle-age spread” with weight gain and the body’s decreasing ability to metabolize calories efficiently. In two newly published studies in animals, researchers show how the dynamics of the mitochondria play a key role in the body’s ability to control weight gain [27,28]. Hence, whether one becomes obese or remains lean can depend on the dynamics of the mitochondria, the body’s energy-producing “battery”.

The purpose of the current study was to conduct a mitochondrial GWAS to identify genetic variants influencing BMI. In particular, we tested 984 mtSNPs in a sample of 6,528 adults, aged 24–85 years.

Results

We performed mitochondrial genome wide association analysis separately for each genotyping chip and corresponding sample (see Table 1). After QC, a total number of 984 mtSNPs were included in the analysis. To assess mtSNP association while taking heteroplasmy into account, we used the raw signal intensity values that were measured on the microarray and applied the linear regression method as described in the Materials and Methods section. The resulting p-values after adjustment for multiple testing are plotted in Figure 1 for each genotyping chip. The x-axis represents the mitochondrial genome, showing the position and relative size of each of the 13 major mitochondrial genes.

Four and one mtSNPs were significant for Affymetrix 6.0 and Illumina Exome chip, respectively. No significant mtSNPs were observed when analysing the Axiom and Metabochip. The association

**Table 1. Summary of the quality control.**

| Chip     | mtSNPs | mtSNPs excluded UB | no_B38 | Study: Sample size | Intensity Ratio Outliers |
|----------|--------|--------------------|--------|-------------------|--------------------------|
| Affy. 6.0 | 411    | 0                  | 54     | F4:1,646          | 217 (0.05%)              |
| Affy. Axiom | 221    | 0                  | 31     | F4:1,550          | 62 (0.05%)               |
| Illum. Exome | 226    | 0                  | 0      | F4:2,721          | 123 (0.05%)              |
| Illum. Metabo | 126    | 0                  | 9      | F3:2,889          | 78 (0.05%)               |

The number of mtSNPs refers to the SNPs that passed QC and were included in the analysis. Several mtSNPs were excluded due to the upper bound cutoff "UB", or because the basepair position was not available in Build 38. The number of intensity measures per allele "rs" represents the total number of intensity measures per allele for the mtSNP.

The purpose of the current study was to conduct a mitochondrial GWAS to identify genetic variants influencing BMI. In particular, we tested 984 mtSNPs in a sample of 6,528 adults, aged 24–85 years.

Results

We performed mitochondrial genome wide association analysis separately for each genotyping chip and corresponding sample (see Table 1). After QC, a total number of 984 mtSNPs were included in the analysis. To assess mtSNP association while taking heteroplasmy into account, we used the raw signal intensity values that were measured on the microarray and applied the linear regression method as described in the Materials and Methods section. The resulting p-values after adjustment for multiple testing are plotted in Figure 1 for each genotyping chip. The x-axis represents the mitochondrial genome, showing the position and relative size of each of the 13 major mitochondrial genes.

Four and one mtSNPs were significant for Affymetrix 6.0 and Illumina Exome chip, respectively. No significant mtSNPs were observed when analysing the Axiom and Metabochip. The association

**Table 1. Summary of the quality control.**

| Chip     | mtSNPs | mtSNPs excluded UB | no_B38 | Study: Sample size | Intensity Ratio Outliers |
|----------|--------|--------------------|--------|-------------------|--------------------------|
| Affy. 6.0 | 411    | 0                  | 54     | F4:1,646          | 217 (0.05%)              |
| Affy. Axiom | 221    | 0                  | 31     | F4:1,550          | 62 (0.05%)               |
| Illum. Exome | 226    | 0                  | 0      | F4:2,721          | 123 (0.05%)              |
| Illum. Metabo | 126    | 0                  | 9      | F3:2,889          | 78 (0.05%)               |

The number of mtSNPs refers to the SNPs that passed QC and were included in the analysis. Several mtSNPs were excluded due to the upper bound cutoff "UB", or because the basepair position was not available in Build 38. The number of intensity measures per allele "rs" represents the total number of intensity measures per allele for the mtSNP.

The purpose of the current study was to conduct a mitochondrial GWAS to identify genetic variants influencing BMI. In particular, we tested 984 mtSNPs in a sample of 6,528 adults, aged 24–85 years.

Results

We performed mitochondrial genome wide association analysis separately for each genotyping chip and corresponding sample (see Table 1). After QC, a total number of 984 mtSNPs were included in the analysis. To assess mtSNP association while taking heteroplasmy into account, we used the raw signal intensity values that were measured on the microarray and applied the linear regression method as described in the Materials and Methods section. The resulting p-values after adjustment for multiple testing are plotted in Figure 1 for each genotyping chip. The x-axis represents the mitochondrial genome, showing the position and relative size of each of the 13 major mitochondrial genes.

Four and one mtSNPs were significant for Affymetrix 6.0 and Illumina Exome chip, respectively. No significant mtSNPs were observed when analysing the Axiom and Metabochip. The association

**Table 1. Summary of the quality control.**

| Chip     | mtSNPs | mtSNPs excluded UB | no_B38 | Study: Sample size | Intensity Ratio Outliers |
|----------|--------|--------------------|--------|-------------------|--------------------------|
| Affy. 6.0 | 411    | 0                  | 54     | F4:1,646          | 217 (0.05%)              |
| Affy. Axiom | 221    | 0                  | 31     | F4:1,550          | 62 (0.05%)               |
| Illum. Exome | 226    | 0                  | 0      | F4:2,721          | 123 (0.05%)              |
| Illum. Metabo | 126    | 0                  | 9      | F3:2,889          | 78 (0.05%)               |

The number of mtSNPs refers to the SNPs that passed QC and were included in the analysis. Several mtSNPs were excluded due to the upper bound cutoff "UB", or because the basepair position was not available in Build 38. The number of intensity measures per allele "rs" represents the total number of intensity measures per allele for the mtSNP.

The purpose of the current study was to conduct a mitochondrial GWAS to identify genetic variants influencing BMI. In particular, we tested 984 mtSNPs in a sample of 6,528 adults, aged 24–85 years.

Results

We performed mitochondrial genome wide association analysis separately for each genotyping chip and corresponding sample (see Table 1). After QC, a total number of 984 mtSNPs were included in the analysis. To assess mtSNP association while taking heteroplasmy into account, we used the raw signal intensity values that were measured on the microarray and applied the linear regression method as described in the Materials and Methods section. The resulting p-values after adjustment for multiple testing are plotted in Figure 1 for each genotyping chip. The x-axis represents the mitochondrial genome, showing the position and relative size of each of the 13 major mitochondrial genes.

Four and one mtSNPs were significant for Affymetrix 6.0 and Illumina Exome chip, respectively. No significant mtSNPs were observed when analysing the Axiom and Metabochip. The association
results that remained significant after adjustment for multiple testing (P_{adjusted} ≤ 0.05) are presented in Table 2. The two most significant mtSNPs associated with BMI are located in the Cytochrome c oxidase 1 and 3 (MT-CO1 and MT-CO3) genes of complex IV, with associated adjusted p-values of 0.014 and 0.029, respectively. Another three locations of NADH subunit dehydrogenase genes of complex I were also significant (MT-ND1: P_{adjusted} = 0.029; MT-ND2: P_{adjusted} = 0.029; MT-ND4L: P_{adjusted} = 0.029). In all significant mtSNPs the estimates of \( \beta_{sex} \) and \( \beta_{age} \) were significant and positive, indicating a significantly higher BMI in males and with older age. All five significant mtSNPs were present only on one chip without overlapping, i.e., they were not shared by the other chips.

A negative parameter estimate for the mtSNP \( (\beta_{SNP} < 0) \) indicates that the risk allele is the minor allele while a \( \beta_{SNP} > 0 \) indicates that the risk allele is the major allele. Taking the most strongly associated variant mt6663A→G in the MT-CO1 gene, based on our estimate \( (\beta_{mt6663A→G} = -0.55 \pm 0.1) \) BMI increases when the ratio of alleles A and G decreases, i.e., an increase in heteroplasmy with the G allele at this locus leads to an increased BMI. The same applies to the other mtSNPs; e.g. having mt10550A→G heteroplasmy with more G than A alleles results in a higher BMI than having only A alleles. The estimates of the model parameters for each significant mtSNP are provided in Table S1.

### Discussion

Excess of weight is a medical condition in which excess body fat has accumulated to the extent that it may have an adverse effect on health, leading to reduced life expectancy and/or increased health problems. The possible role of mitochondria in the development of obesity is mainly concerned with the efficacy of mitochondrial energy provision [29,30]. Apparently, the balance
Table 2. Summary of significant mtSNPs.

| Chip         | Gene                        | SNP          | Location | Type of mutation | rs_number       | Adj p-value | Minor allele | Major allele | Adj p-value | Minor allele | Major allele |
|--------------|-----------------------------|--------------|----------|------------------|-----------------|-------------|--------------|--------------|-------------|--------------|--------------|
| Affy6.0      | ND1: MT-ND1; subunit of NADH dehydrogenase (complex I) | mt3336T     | 2.6 × 10^-4 | synonymous       | 0.23            | 2.6 × 10^-4 | G            | T            | 2.6 × 10^-4 | G            | T            |
| Illum. Exome | ND1: MT-ND1; subunit of NADH dehydrogenase (complex I) | mt4851C     | 1.11     | synonymous       | 0.55            | 1.11        | C            | T            | 1.11        | C            | T            |
| Affy6.0      | ND1: MT-ND1; subunit of NADH dehydrogenase (complex I) | mt9698T     | 0.23     | synonymous       | 0.55            | 0.23        | T            | G            | 0.55        | T            | G            |
| Affy6.0      | ND1: MT-ND1; subunit of NADH dehydrogenase (complex I) | mt10550A    | 0.31     | synonymous       | 0.55            | 0.31        | A            | G            | 0.55        | A            | G            |
| Illum. Exome | ND1: MT-ND1; subunit of NADH dehydrogenase (complex I) | mt6663A     | 6.2 × 10^-4 | synonymous       | 0.55            | 6.2 × 10^-4 | G            | A            | 0.55        | G            | A            |

Genomic position in base pairs (bp), alleles, rs_number, and type of mutation are based on the NCBI dbSNP GRCh38 human genome assembly (rCRS, GeneBank minor allele. An estimated effect size (b), 0 indicates that the minor allele increases BMI. Nominal p-values and adjusted p-values are provided.

Table 2. Summary of significant mtSNPs.

Mitochondrially encoded cytochrome c oxidase subunits, complex IV, is a key oxidative enzyme regarded as one of the major regulation sites for the OXPHOS system, controlled by both nDNA and mtDNA. Its catalytic activity is primarily determined by 3 of the 13 subunits which are encoded by the mtDNA (MT-CO1, MT-CO2, and MT-CO3) [31]. The loss of function of this enzyme has been suggested to trigger ROS production, although the increase in radical accumulation rests not with the ETC but with non-mitochondrial sources [32]. However, the function of each subunit and the molecular mechanism behind the regulation on the activity of this important protein complex are largely unknown [33]. The interesting finding that suppression of cytochrome c oxidase activity also sensitizes cells to death signals [33] could shed some light on the role of defective cytochrome c oxidase in aging and age-related diseases as an increase of BMI with age.

MT-ND1mt3336T→G and MT-CO3mt9698T→C

Mitochondrially encoded NADH dehydrogenase subunit genes, complex I, is the first enzyme in the mitochondrial respiratory chain. It extracts energy from NADH, produced by the oxidation of sugars and fats, and traps the energy in a potential difference or voltage across the mitochondrial inner membrane. The potential difference is used to power the synthesis of ATP. Because NADH dehydrogenase is central to energy production in the cell, its malfunction may result in a wide range of metabolic disorders. Some of them are due to mutations in the mitochondrial genome, while others, which result from a decrease in the activity of complex I, or an increase in the production of ROS, are not yet well understood.

MT-ND1 and MT-ND2 genes were previously reported to be related with body fat mass and intramuscular fat in animals [23,34,35]. The significant findings in our study at mt3336T→G (MT-ND1) and mt4851C→T (MT-ND2) have associated effect sizes estimated to be −0.23 and −1.11, respectively. In other words, increasing the number of T alleles (or decreasing G alleles) at mt3336 in an individual will help to reduce the BMI, while increasing the number of T alleles (or decreasing C alleles) at mt4851 will contribute to increase BMI. In this study we identified for the first time a significant association between MT-ND1 (mt10550A→G) and BMI, with an estimated effect size of −0.31. Therefore, the G allele is the one to be related with high BMI.
values, i.e., people with heteroplasmy having a larger number of G alleles will have a higher BMI than people with a smaller number of G alleles. MT-ND1 is one of the 7 mtDNA-encoded subunits included among the approximately 41 polypeptides of respiratory complex I. MT-ND4L is probably a component of the hydrophobic protein fragment of the complex [36].

While associations of the MT-CO1 gene with intramuscular fat in animals have been reported [35], no significant findings have been reported so far for the MT-CO3 gene with BMI, nor with other BMI-related phenotypes. Liu et al. [37] suggested an effect of mtSNPs, including MT-CO2, MT-ND1, and MT-ND2 genes, on BMI but this effect failed to reach significance after adjusting for multiple testing. Similarly, Saxena et al. [38] detected nominal signals of association between BMI and mtSNPs, including MT-ND1, MT-ND2, and MT-ND3 genes; however after adjusting for multiple testing these associations disappeared.

Recently, Knoll et al., 2014 [39] did not find association between mtSNPs and obesity in children. However, this may be because heteroplasmy was not included in their study or different mtSNPs might be relevant for children than adults with regard to BMI.

Two of the significant variants MT-ND1™3336T→G and MT-CO1™9698T→G are missense mutations which lead to an amino acid change, thus being a non-synonymous variant. So, individuals with an excess of missense mutations may carry an appreciable fraction of an altered protein that is responsible for an elevated BMI. The other three significant variants MT-ND2™4851C→T, MT-CO3™9698T→C, and MT-ND4L™10550A→G are synonymous mutations, i.e., they code for the same amino acid. How an excess of synonymous mutations at these loci could impact BMI needs further investigation, since the single nucleotide change leads to an unchanged protein. However, different codons might lead to differential protein expression levels. Two of the synonymous variants, mt9698C and mt10550G, are present in the European mtDNA haplogroups U8 and K, respectively. However, to date only haplogroup T has been suggested as a risk factor for obesity [24].

These results together with previously published findings suggest that genetic variation in NADH dehydrogenase and cytochrome c is a risk factor for several metabolic disorders, including BMI and BMI-related phenotypes. Multiple lines of evidence suggest that mitochondrial dysfunction as well as altered mitochondrial genome expression, particularly of genes encoding complex I–IV may play a role in BMI. Searching for previous suggested candidate mtSNPs associated with BMI or related phenotypes has yielded a small number of positive findings, possibly due to low sample sizes or the lack of taking heteroplasmy into account in the analysis. In this study we have considered the heteroplasmic effect and found mtSNPs in candidate gene regions to be significantly associated with BMI that previously were suggested to be implicated in obesity, body fat mass, or intramuscular fat (MT-ND1, MT-ND2, and MT-CO1). In addition, two novel susceptibility mtSNPs in the MT-ND4L and in the MT-CO3 regions have been identified. In both polymorphisms heteroplasmic individuals towards a larger number of the respective minor allele increases the risk of developing a high BMI.

The regression models for all significant mtSNPs identified in this study also corroborate the generally acknowledged fact that males have slightly but significant higher BMI values than females and BMI also increases significantly with older age. With regard to sex, this effect is to be expected due to the fact that adult males and females with the same BMI are classified identically in regard to whether their weight is normal. However, even though men may have slightly higher BMI values than women, this does not necessarily mean that they are more overweight. For instance, women typically have more body fat than men, whereas men typically have more muscle than women, and muscle has a much greater density (it takes up less volume than an equal mass of fat).

It is also known that BMI is largely influenced by age in adults [40,41]. It has been suggested that BMI increases linearly with age in women, but in men it increases with age in two stages – a more intensive rise between age 20–40 and a much slower increase between age 40–60 [42]. High BMI can occur at any age, but with both muscle mass and strength beginning to decline around the age of 40, an increased BMI at this age may be related to mitochondrial dysfunction.

Conclusion

In summary, our study reports genetic variants located in the cytochrome c subunits 1, 3, and in the NADH dehydrogenase subunits 1, 2, and 4L of the mtDNA that are significantly associated with BMI in adults. Heteroplasmy in these variants towards the minor allele increases the risk of developing high BMI. Based on these findings we hypothesize that common metabolic disorders as obesity or high BMI are attributable at least in part to mitochondrial polymorphisms. Animal and human data consistently show that mitochondria are altered in aging, leading to increased mutations in mtDNA, decreased expression of some mitochondrial proteins, reduced enzyme activity, and altered respiration with reduced maximal capacity in sedentary adults. Since the primary role of mitochondria is to produce ATP to maintain the energy status of the cell, shifts in respiratory activity and capacity can lower the membrane potential, reduce cellular ATP concentration, and signal cellular apoptotic events. Increased apoptosis without correspondingly increased protein synthesis will eventually lead to net muscle fiber loss. All of these factors are likely to contribute to age-associated BMI, and mounting evidence suggests that most of these age-related changes can be either prevented or attenuated through increased physical activity.

There are some thoughts that the accumulation of oxidative damage caused by long-term ROS production is responsible for these changes with age. Cytochrome c and NADH dehydrogenase complexes are both involved in the regulation of mitochondrial ROS. Indeed, blocking ROS formation in animals by the targeted expression of antioxidant enzymes ameliorates age-associated dysfunction and returns mitochondrial parameters to those of young animals [43]. Whether this is also true for humans remains to be shown. However, behavioral factors that may improve mitochondrial function and BMI control, such as physical activity and caloric restriction, also reduce ROS production and increase antioxidant defenses. Although physical activity training in elderly adults probably does not completely reverse the primary effects of aging, it may clearly reduce the rate of mitochondrial decline and attenuate the aging processes [44].

These findings highlight the important role of the mtDNA among the factors that contribute to the risk of high BMI in adults and suggest that variants in the mitochondrial genome may be more important than has previously been suspected. Therefore, a challenge for possible novel therapeutic strategies might be the further elucidation of intracellular ROS signaling pathways for effective therapeutic gain.

Materials and Methods

Study design and population

The Cooperative Health Research in the Region of Augsburg (KORA) study is a series of independent population-based
were subjected to several medical examinations including BMI in the study.

Genotyping

DNA was extracted from full blood after the blood draw and then stored at −80°C. Genotyping was performed using different platforms such as the Affymetrix 6.0 GeneChip array (KORA F4), Illumina MetaboChip 200K (KORA F3 and F4), Illumina Human Exome Beadchip array (KORA F4), and Affymetrix Axiom chip array (KORA S4 and F4). A total number of 6,528 individuals were genotyped. Some individuals of KORA F4 were present on more than one chip, some of them (1,549) were included on all chips. Only single-nucleotide polymorphisms located in the mitochondrial genome (mtSNPs) were considered in this study. Genotyping data and samples are summarized in Table 3. Most of the covered mtSNPs have distinct positions identified by different chips. Although the Affymetrix 6.0 is the one containing the largest number of mtSNPs some regions are not well covered. The Illumina Metabochip contains the smallest number of mtSNPs and many regions are uncovered, especially the hypervariable regions (HVRI and HVRII) as well as the CO1 and CO2 genes. However, when all chips are considered together, good overall coverage of the mitochondrial genome is obtained (Figure 2).

Quality control

First a visual check was performed plotting the intensity measures of each mtSNP separately for each chip. In this step only the Axiom chip displayed anomalies in some intensity values. There seems to be an upper bound for the intensity values at an intensity of 4095. As a result some mtSNPs had a significant fraction of either their A or B intensities squeezed into the region just below that upper bound; apparently without the bound those intensities would have taken higher values. Since intensities close to the cut-off value of 4095 cannot be trusted, we proceeded as described in the Text S1.

Table 3. Distribution of characteristics of the study population.

| Chip          | mtSNP | KORA | Individuals (Males/Females) | Mean age Males/Females | Mean BMI Males/Females |
|---------------|-------|------|-----------------------------|------------------------|------------------------|
| Affy. 6.0     | 465   | F4   | 1666 (788/858)              | 60.4 ± 8.8/60.1 ± 8.7  | 28.2 ± 4.3/27.5 ± 4.9  |
| Affy. Axiom   | 252   | F4/F4| 3330 (1702/1828)            | 54.1 ± 14.2/53.6 ± 13.9| 27.7 ± 4.2/26.9 ± 5.1  |
| Illum. Exome  | 226   | F4   | 2721 (1294/1427)            | 55.5 ± 13.2/54.8 ± 13.1| 27.7 ± 4.0/26.9 ± 5.1  |
| Illum. Metabochip | 135 | F3   | 2889 (1393/1496)           | 56.6 ± 12.9/56.1 ± 12.5| 27.8 ± 3.7/27.0 ± 5.0  |
| F4            | 2817  | F4   | 1338 (1338/1479)           | 55.5 ± 13.2/54.9 ± 13.1| 27.7 ± 4.1/27.0 ± 5.1  |

The mtSNPs column shows the number of mitochondrial SNPs provided by each chip. Distributions are presented as means ± standard deviation.

doi:10.1371/journal.pone.0105116.003
described in Text S1. A summary of the quality control results is given in Table 1.

Statistical methods

To approach the heteroplasmy present in the mitochondria we used the raw signals of luminous intensity, where every measurement is associated with a specific mtSNP and represents one of its alleles [48,49]. The number of measures \( n \) per mtSNP depends on the vendor-specific technology employed on the genotyping chip. At the very least there have to be two signals, one for each of the two alleles. Often, however, the chip design includes more than one measurement per SNP and allele. That is, for every individual and SNP we have intensity measurements \((A_1, B_1), \ldots, (A_n, B_n)\) with \( n \geq 1 \) where \( A_i \) and \( B_i \) represent the intensities of the two alleles A and B. The best way to assess the association of BMI with the mtSNP intensities at a given age is to apply linear regression analysis using BMI as quantitative response variable. The mtSNP enters the model as a covariate via the log2-transformed intensity ratio, \( \log_2(A_i/B_i) \), where \( A \) and \( B \) denote the mean intensity, or single measure in case of \( n = 1 \), for the A allele and B allele, respectively. To improve the convergence properties of the model estimates we center this variable (\( z = \log_2(A_i/B_i) - \mu \)) as well as the additional quantitative covariate age at examination, i.e., the age at the time that the mtSNPs and BMI were measured. Sex is also introduced in the model as a covariate. The linear regression model for the \( j^{th} \)-mtSNP is:

\[
\text{BMI}_j = \beta_0 + \beta_1 z_j + \beta_2 \text{age}_j + \beta_3 \text{sex}_j + e_j \quad \text{for} \ j = 1 \ldots m
\]

where \( j \) denotes the individual and \( m \) is the sample size. The aim of this study is to evaluate how BMI is influenced by mitochondrial variants taking into account the age for a particular person on the phenotype BMI, and therefore the model adjusts for individual age differences in the study population. Using this model it is not possible to make any claims based on the effect of age on heteroplasmy at the particular mtSNP. However, due to the adjustment by age on BMI in the model, we can be sure that in case of positive findings the BMI change is in fact caused by variations of heteroplasmy at the particular mtSNP, and not by differences in age that jointly influence BMI and heteroplasmy, without an effect of heteroplasmy on BMI.

P-values were obtained from the Wald test, which is based on the asymptotic normality of the regression coefficient estimates, and corrected for multiple comparisons applying the Benjamini-Hochberg false discovery rate (FDR) using the R function p.adjust [50]. The resulting adjusted P-values are also known as “Q-values”. Each type of genotyping chip needs to be analyzed separately because different chips make use of different technologies, even between chips of the same manufacturer. The two Affymetrix chips, Affy6.0 and Axiom, share 170 positions and Illumina chips, Exome and Metabochip, share 44 positions. Only 9 positions are common to all four chips. Overlapping mtSNPs were analysed separately for each respective genotyping chip, giving in this way the opportunity of validation in the case of significant results. All the analyses were performed with the R statistical software [51].

Supporting Information

Figure S1 Examples of intensities affected by the cut-off.
(TIFF)

Figure S2 Resulting p-values from the 1st analysis (left side) and 2nd analysis (right side).
(TIFF)

Table S1 Estimates of the model parameters for mtSNPs with a \( P_{\text{nominal}} < 0.01 \). Genomic position in base pairs (bp), alleles, rs_number, and point mutation are based on the NCBI dbSNP GRCh38 human genome assembly (rCRS, GeneBank ID J01415.2). Alleles are given in terms of major→minor allele. An estimated effect size \( (\beta_{\text{SNP}}) < 0 \) indicates that the risk allele is the minor allele. Nominal p-values and adjusted p-values are provided. \( \mu \): mean of \( \log_2(A/B) \). Covariate sex baseline: male.
(DOC)

Text S1 Quality control.
(DOCX)

Acknowledgments

We thank all members of the field staff involved in the KORA study and we express our appreciation to all study participants.
Author Contributions
Conceived and designed the experiments: AF KS. Performed the experiments: HG MW JK. Analyzed the data: AF CB. Contributed reagents/materials/analysis tools: TM AP HG MW. Wrote the paper: AF. Revising the article critically for important intellectual content: KS HG MW.

References
1. Wormser D, Kaptoge S, Di Angelantonio E, Wood AM, Pencina L, et al. (2011) Separate and combined associations of body-mass index and abdominal adiposity with cardiovascular disease: collaborative analysis of 58 prospective studies. Lancet 377: 1083–1095.
2. Vazquez G, Duvall-Scherrer DR Jr, Silventoinen K (2007) Comparison of body mass index, waist circumference, and waist/hip ratio in predicting incident diabetes: a meta-analysis. Epidemiol Rev 29: 115–128.
3. Renihan AG, Tyson M, Egger M, Heller RF, Zwahlen M (2008) Body-mass index and incidence of cancer: a systematic review and meta-analysis of prospective observational studies. Lancet 371: 569–576.
4. World Health Organization (2010) International Statistical Classification of Diseases and Health Related Problems. Geneva: World Health Organization.
5. Haslam DW, James WP (2005) Obesity. Lancet 366: 1197–1209.
6. Frontera WR, Hughes VA, Lutz KJ, Evans WJ (1991) A cross-sectional study of muscle strength and mass in 45- to 78-yr-old men and women. J Appl Physiol (1985) 71: 643–650.
7. Cree MG, Newcomer BR, Katsumos CS, Sheffield-Moore M, Chinkes D, et al. (2000) Intramuscular and liver triglycerides are increased in the elderly. J Clin Endocrinol Metab 85: 3864–3871.
8. Hipkiss AR (2010) Mitochondrial Dysfunction, Proteostasis, and Aging: Causes or Effects, and the Possible Impact of NAD+-Controlled Protein Glycation. In: Makowski GS, editor. Advances in Clinical Chemistry. Burlington: Academic Press. pp. 125–150.
9. Johannsen DL, Conley KE, Bajpeyi S, Punyanitya M, Gallagher D, et al. (2012) Deficiency of electron transport chain in human skeletal muscle mitochondria in morbid obese population from southern Italy. Biomed Res Int 2013: 631082.
10. Mancini C, Di Marco A, Bottoni E, Galluzzi L, Mocchetti I (2007) Haplogroup T is an obesity risk factor: mitochondrial DNA haplotyping in a related BMI. Pediatr Res 65: 103–108.
11. Sankhla M, Sharma TK, Mathur K, Rathor JS, Butolia V, et al. (2012) Increased oxidative stress in obesity and its impact on metabolic syndrome. J Clin Endocrinol Metab 97: 281–285.
12. Lee H, Lee YJ, Choi H, Ko EH, Kim JW (2009) Reactive oxygen species and aging: a review. J Aging Res 2012: 194821.
13. R Core Team (2013) R: A Language and Environment for Statistical Computing. Vienna, Austria: R Foundation for Statistical Computing.
14. Maie P, Lombardi AM, Park JH, Puntis M, De Vos R, et al. (2009) Mitochondrial energy production correlates with the age-related BMI. Pediatr Res 65: 103–108.
15. Yang TL, Guo Y, Shen H, Lai SF, Liu YJ, et al. (2011) Genetic association study of common mitochondrial variants on body fat mass. PLoS One 6: e21595.
16. Bournat JC, Brown CW (2010) Mitochondrial dysfunction in obesity. Curr Opin Endocrinol Diabetes Obes 17: 446–452.
17. Speliotes EK, Willer CJ, Berndt SI, Monda KL, Thorleifsson G, et al. (2010) Separate and combined associations of body mass index. Nat Genet 42: 937–948.
18. Speliotes EK, Willer CJ (2010) Structure of NADH-ubiquinone reductase (complex I). J Biol Chem 285: 10601–10609.
19. Chretien D, Rustin P, Bourgeron T, Rotig A, Saudubray JM, et al. (1994) Reference charts for respiratory chain activities in human tissues. Clin Chim Acta 229: 53–70.
20. Elson JL, Andrews RM, Chinnery PF, Lightowlers RN, Turnbull DM, et al. (2005) Mitochondrial genome polymorphisms associated with type-2 diabetes or obesity. Mitochondrion 5: 15–33.
21. Lee SH, Kim SC, Choi BH, Lim D, Kim NK, et al. (2012) mt-COX1, mt-ND1 and CREBP are indicators of intramuscular fat content in Hanwoo (Korean cattle). Livestock Science 146: 160–167.
22. Ragan CI (1987) Structure of NADH-ubiquinone reductase (complex I). Curr Top Bioenerg Biomembr 15: 1–36.
23. R Core Team (2013) R: A Language and Environment for Statistical Computing. Vienna, Austria: R Foundation for Statistical Computing.