The widespread increase in inter-individual variability of gene expression in the human brain with age

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ABSTRACT

Aging is broadly defined as a time-dependent progressive decline in the functional and physiological integrity of organisms. Previous studies and evolutionary theories of aging suggest that aging is not a programmed process but reflects dynamic stochastic events. In this study, we test whether transcriptional noise shows an increase with age, which would be expected from stochastic theories. Using human brain transcriptome dataset, we analyzed the heterogeneity in the transcriptome for individual genes and functional pathways, employing different analysis methods and pre-processing steps. We show that unlike expression level changes, changes in heterogeneity are highly dependent on the methodology and the underlying assumptions. Although the particular set of genes that can be characterized as differentially variable is highly dependent on the methods, we observe a consistent increase in heterogeneity at every level, independent of the method. In particular, we demonstrate a weak but reproducible transcriptome-wide shift towards an increase in heterogeneity, with twice as many genes significantly increasing as opposed to decreasing their heterogeneity. Furthermore, this pattern of increasing heterogeneity is not specific but is associated with a wide range of pathways.

INTRODUCTION

Aging is commonly defined as a time-dependent decrease in the functional and structural integrity of an organism. Despite the ubiquity of aging in all living organisms, the molecular mechanisms responsible still require further elucidation. According to recent studies, aging differs phenotypically among individuals, including monozygotic twins [1,2] and within tissues from the same individuals [3]. Researchers have observed an age-related increase in variability in the epigenome [4,5] and transcriptome [6] of genetically identical samples, which may underlie the phenotypic differences. Age-related expression variability has been detected in many different cell and tissue types including mice stem cells, cardiomyocytes and immune cells [7–9], rat neural retina [10], fruit-fly, mice and human brain [6,11–14] as well as human pancreas, lung, blood, skin, fat and human fibroblasts in vitro [13,15–17]. Despite these reports, there is no agreement on the underlying mechanisms, extent and functional consequences. Suggested mechanisms include somatic [7,15] and germline mutations [11,17], changes in the DNA methylation [9,17,18] and chromatin modifications [5] and resulting chromatin compaction [12] as well as global dysregulation, caused by the change in transcription factor or miRNA expression [19].
Both genome-wide and hypothesis-driven approaches have been employed to explore the extent of expression variability with age. Among the former, some show a transcriptome-wide increase [6,9,12,13,15], while others focus only on those genes showing significant changes in their variability. Brinkmeyer-Langford et al. [11] report that an equal number of genes significantly increase or decrease their expression variability, whereas a recent study from Vinuela et al. [17] shows more genes decreasing rather than increasing their expression variability [17]. Hypothesis-driven studies mostly show an increase in variability for the genes measured [7,8,16], whereas Warren et al. [20] suggest this might be specific only to the non-renewing tissues. Similarly, Ximerakis et al. [14] show that change in transcription variability is in different directions in different cell types of mouse brain. The reports also vary in terms of the functional association of this variability. While some consider that increase in variability is widespread [6,12], others report that variability is concentrated in various cellular functions [10,11,18,21] – although these functions also differ between reports.

Age-dependent change in the expression variability is difficult to address due to the inherent noise in expression and the influence of other factors on variability. Thus, the data pre-processing steps to disentangle variability from the biological and technical confounders is of importance. Another technical aspect is the method to measure the change in the variability. Most studies tested for age-related change in the expression variability using either grouped (Bartlett’s test, Levene’s test, permutation test) [7,11,20] or regression-based tests (linear and loess regression) [6,10,17,18], with a few others using correlation-based approaches (gene co-expression, intra-class correlations) [21,22]. However, to our best knowledge, the effects of different batch-correction strategies and different methods to measure variability have not been explored on the same data.

In this study, we undertook a comprehensive investigation of the aging-related change in expression variability, using human brain expression dataset. We employed different pre-processing and variability measures and analyzed transcriptome-wide and gene-level changes in gene expression variability and the associated functions.

RESULTS

In order to study the change in gene expression variability during aging, we used one of the biggest published human brain transcriptome datasets, generated using microarray technology [23]. We limited the age range to between 20 and 80 years (Figure 1A), resulting in RNA expression data for 147 prefrontal cortex samples. We excluded prenatal, infant and childhood samples (up to 20 years old) because their expression levels are inherently coupled with developmental processes in the brain. We applied four batch correction strategies to account for technical and biological confounders (Supplemental Figure 1): i) only...
quantile normalization (QN), ii) QN followed by linear regression (regression), iii) QN followed by ComBat [24], and iv) QN followed by Surrogate Variable Analysis (SVA) [25]. Regression and ComBat are supervised approaches, i.e. known covariates should be supplied to the algorithm, whereas SVA estimates covariates from the data. We provide the results from Regression and SVA in the main text to include one supervised and one unsupervised approach. The results from other correction strategies are given in Supplemental Data in comparison with the Regression and SVA approaches (Supplemental Figures 2-8).

Analysis of the differentially expressed genes

First, we defined differentially expressed (DE) genes, based on the significance of the regression coefficients (FDR corrected p <= 0.05) for the linear model using the gene expression values as the dependent and age as the independent variable (see Methods). By applying two different pre-processing approaches on the data prior to DE estimation the number of DE genes was very different. SVA correction yielded 3499 DE genes, compared with 881 DE genes found by regression (Figure 1B, Supplemental Table 1). Nevertheless, 854 genes overlapped between the SVA and regression results, which constituted a quarter of genes found after SVA and 96% of the genes identified after regression correction. Quite a high overlap was consistent with the strong correlation between the expression level changes for the regression and SVA corrected data (Spearman ρ = 0.85, Supplemental Figure 3A).

To explore the biological processes affected by these changes in gene expression, gene set enrichment analysis was performed separately on the regression and SVA corrected data (Supplemental Table 2). It revealed 125 (out of 191 (regression) and 160 (SVA) categories) shared Gene Ontology Biological Process (GO BP) categories that were downregulated in the aging brain. Cognitive-function related GO terms, such as modulation of synaptic transmission, learning or memory, constituted a substantial fraction of these GO terms. In contrast, the number of the upregulated GO terms was much smaller and only 8 GO terms overlapped (out of 22 (regression) and 12 (SVA) categories) between the correction approaches, including detoxification, stress response to metal ions and cilium organization GO categories (see Supplemental Table 2).

Analysis of the differentially variable genes

Two different strategies were employed to measure change in the gene expression variability with age, namely continuous and grouped approaches. The continuous approach detects continuous monotonic change in variation from 20 to 80 years of age. The grouped approach compares the gene expression variation between two age groups: young (20 - 40 years old, N = 53) and old (60 - 80 years old, N = 22). Figure 2 illustrates the principles of these approaches and shows that the change in variability can be combined with any dynamics in the mean gene expression (upregulation, downregulation, no change). We checked if the changes in gene expression variability were confounded by the changes in gene expression level, but did not observe any relationship (Supplemental Figure 10, Fisher’s test p = 0.11, Odds ratio = 1.05).

In the continuous approach, we first fit a linear model to explain age-dependent change in expression (Figure 2, first column) and then used the residuals from this model to represent the variability. To measure change in the expression variability with age, we calculated the Spearman correlation coefficient (Δvar(ρ)) between the absolute value of residuals and age (Figure 2, middle column). The Δvar(ρ) measures ranged between -0.32 and 0.36 and were normally distributed (Shapiro-Wilk test, p > 0.05, see Methods) (Figure 3A). The distributions were significantly shifted towards positive values for both correction methods (median values range between 0.01 to 0.03, Wilcoxon test, p < 2.2e-16). Although the shift in the distribution was small, 57% to 63% percent of the genes showed increase in variability with age. However, we noted that the changes in variability calculated for each gene, using regression- and SVA-corrected data, were only weakly correlated, ρ(Δvar(reg), Δvar(SVA)) = 0.35 (Figure 3B).

In the grouped approach, we first generated a distribution of expected variability in gene expression for the young individuals and treated it as a null distribution to compare with the variability from the old individuals. We used interquartile range (IQR) as a measure of variability because it is robust to outliers. In order to calculate a distribution of expected variability in the young group, we randomly selected a subsample of 22 individuals (the number of samples in the old group) from the 53 individuals in the young group 10 000 times and calculated IQR. The change in variability, Δvar (IQR), was measured as a fractional change in the IQR between old and young groups (see Methods). The p-value was determined by calculating how many times we observed value as extreme as IQRold (see Methods). The distributions of change in variability, Δvar(IQR), were moderately skewed to the right and ranged from -0.70 up to 2.10 for the regression corrected data and from -0.78 up to 1.71 for the SVA corrected data (Figure 3C). The skew to the right was expected given that we calculate variability change as a fraction and, thus, it was more sensitive to increase in...
Figure 2. Changes in gene expression and its variability with age for some individual genes, using the different approaches. Example genes are chosen that increase (A) or decrease (B) expression variability with age, when the mean gene expression either increases, does not change or decreases. The types of change, for expression and variability respectively, is shown in the parenthesis following the gene name, for each row. Genes were selected to have the biggest absolute values of $\Delta \text{var}(\rho)$ and $\Delta \text{var}(\text{IQR})$ as well as demonstrate significant increase, decrease or no change in the expression level with age. The first column to the left illustrates mean expression level (regression-corrected) plotted against individual’s age on the $x^{0.25}$ transformed scale. The regression line is colored in blue, with the $\beta_1$ coefficient from the linear regression shown on the graph. The middle column illustrates the continuous approach to measure differential variability. Absolute values of the residuals (in red) from the regression line are plotted against age and the regression line between residuals and age (in blue) is drawn for illustrative purposes. The Spearman correlation estimates, $\Delta \text{var}(\rho)$ between the residuals and age are displayed on the graph and used in the subsequent analysis. The last column on the right illustrates the grouped approach to calculate differential variability. Gene expression levels (regression-corrected) of the individuals from the “young” (20 – 40 years old) and “old” (60 – 80 years old) groups are represented in the corresponding boxplots. A small random deviation (jitter) from the x-axis is applied for better visualization. $\Delta \text{var}(\text{IQR})$, the fractional change in the variability in the “old” group, as compared to the “young”, is displayed on the graph.
variability. In both cases, the distributions demonstrated a significant deviation from zero (Wilcoxon test, p-value < 2.2e-16 both for regression and SVA corrections). The data revealed that 6% and 2% more genes showed more variability in the old group, for regression and SVA approaches respectively. Similar to the continuous approach, the effect sizes calculated using regression and SVA corrected data correlated weakly \( \rho(\Delta \text{var}(\rho), \Delta \text{var}(\text{IQR})) = 0.24 \), Figure 3D).

### Gene-level differential variability

We then asked if we could detect any genes with a significant change in variability. Using the **continuous approach**, we did not detect any significant change in variability with age after the multiple testing correction (Supplemental Table 3). The **grouped approach** leads to 741 and 746 differentially variable (DV) genes (FDR corrected \( p \leq 0.05 \)) using the regression and SVA correction, respectively (Figure 4A, Supplemental Table 4). However, the two sets of DV genes identified only have 83 genes in common (Figure 4A), one of which shows an opposite direction of change in the two sets. The correlation between \( \Delta \text{var} \) (IQR) for regression and SVA corrected data is weak \( (p = 0.24) \), but correlation increases when we select only the common DV genes \( (p = 0.44) \) (Figure 4B). In agreement with our overview analysis above, we find twice as many DV genes with an increase in variability as those that decrease variability, using both correction methods: i) 533 genes increase and 241 decrease their variability in the regression correction, ii) 505 genes increase and 241 decrease their variability in the SVA correction (Figure 4A).

### Differential variability of functional groups

Following the individual gene analysis, we explored whether genes that tend to increase or decrease variability with age are localized in particular functional groups. We performed multiple gene set enrichment...
analyses (GSEA) using the change in the variability with age (Δvar) measures obtained in the continuous and grouped approaches on the gene sets from KEGG and Biological Process GO categories (Supplemental Tables 5, 6). We observed no genome-level significant enrichment in particular functional groups on the data either from the continuous (SVA correction), or the grouped approach (Regression and SVA corrections). However, we found that 4 pathways, namely beta-Alanine metabolism, Ras signaling pathway, Phosphatidylinositol signaling system, Bacterial invasion of epithelial cells (FDR corrected p ≤ 0.05) were enriched among the genes showing more variability of expression in the continuous approach (Regression correction). These pathways had positive normalized enrichment scores (NES) i.e. enrichment for the genes that increase variability with age. Moreover, these pathways also had positive NES for other approaches, even though they were not significant (Supplemental Table 5).

Distribution of the DV genes in the pathways

The gene set enrichment analysis shows if there are particular gene sets that include the genes with the highest increase or decrease. Failing to detect such functional categories, we asked how the variability measures for the genes were distributed in the different functional groups of genes. For each of 310 KEGG pathways, encompassing 5922 unique genes, we analyzed the distributions of Δvar measures (Supplemental Table 7), focusing on the median value for the change in variability (Figure 3A, C). In line with the overall tendencies we observed (Figure 3A, C), the majority of pathways contained a larger number of genes that become more variable with age, irrespective of the approach or correction method used. Although the increase in variability is ubiquitous and is observed across the majority of the pathways (74-94%), the increase is small (the mean value for the shift in distributions range between 0.021 and 0.033) – in accordance with the small, but significant increase observed in the distribution for all genes. Since the pathways are not mutually exclusive, we checked if there are particular genes that are present in many different pathways and cause the shift. However, no significant correlation between the pathway membership of gene and its variability measure (Δvar) was detected (Supplemental Figure 11). We repeated the analysis using GO Biological Process categories and observed a similar trend (see Supplemental Information, Supplemental Table 7, 8).
Using one of the largest publicly available human brain expression datasets, we have investigated the change in the variability of the gene expression with age. We applied and compared different approaches to identify differentially variable genes and correction strategies to adjust for the confounders. Our comparison showed that the correction strategy plays a pivotal role in identifying the specific set of differentially variable (DV) genes. However, irrespective of the approach and correction method used, we observed a transcriptome-wide increase in the gene expression variability, i.e. more genes showed a tendency to increase than to decrease expression variability with age. We also showed that most of the functional processes (as defined in KEGG and GO) were susceptible to the aging-related increase in the expression variability.

The difference between the continuous and grouped approaches can be explained by the power and initial assumptions of each method. While the continuous approach assumes a linear change in expression with age\(^{0.25}\) (see Supplemental Figure 4 for the results showing high concordance between models using age vs. age\(^{0.25}\) – correlation coefficient (\(\rho\)) ranges between 0.995 to 0.997), the grouped approach analyzes each age-group within itself and is not sensitive to different dynamics of gene expression change. However, the grouped approach requires expression levels to be similar within the young and old groups. The continuous approach is well suited to detect monotonic changes in variability, whereas the grouped approach can detect switch-like changes, e.g. when variability stays the same throughout the lifespan but changes abruptly at the age of 60. In contrast, the continuous approach focuses on the whole aging period, while the grouped approach overlooks the middle-age group (40-60). Finally, both methods are vulnerable to power issues as the continuous approach uses Spearman correlation, a non-parametric method, and the grouped approach analyzes only a subset of the data. Thus, we compared the variability measure of each gene, calculated using these two approaches. The variability measures are moderately correlated (\(\rho = 0.43\)) for the regression correction and strongly correlated (\(\rho = 0.71\)) for the SVA correction (Supplemental Figure 9). Overall, the differences in the results using these two approaches.
approaches create a challenge in interpretation, but they are not surprising given inherent differences in methodology and the small changes in variability we are investigating.

Another technical aspect we considered was the effect of pre-processing steps. While applying regression and SVA corrections, we showed that significantly DV genes hardly overlap between the corrections, with only 6% being in common (Jaccard similarity) (Figure 4A). Unfortunately, current approaches for handling transcriptome data are designed only to remove the confounding factors on the expression level and not on the expression variability. Thus, SVA and regression demonstrated much higher agreement in the differentially expressed (DE) genes (24% in common, Jaccard similarity) (Figure 1B). That raises a question: which set includes the genuine DV genes? The different correction strategies are quite distinct and might be accounting for different aspects, which is evident from the weak correlation between them (Spearman ρ between regression and SVA-corrected data for continuous approach – 0.35, grouped approach - 0.24, Supplemental Figure 6-7). In this case, the union may capture the full aspects of differential variability, whereas the overlap can provide the gene list in which we are most confident.

Independent of the correction strategy, two-thirds of the DV genes showed a significant increase in variability. These results agree with the reports of Li et al. [10] on mice neural retina, but disagree with findings of Brinkmeyer-Langford et al. [11] on the human brain and Vinuela et al. [17] on the multiple human tissues which show either equal amount of genes increase and decrease in variability or more decrease than increase. The small overlap of our DV gene set with Brinkmeyer-Langford et al. [11] (see Supplemental Information) could be explained by the technical aspects that we presented, i.e. variability measure and data pre-processing, as well as use of different experimental setups and different age-ranges.

We further asked if there is a shift towards an increase or decrease in variability (above or below zero) across the whole transcriptome, irrespective of the values and significance. In accordance with the previous findings on the human, rat and fruit fly [6,12,15], we found as many as 63% of genes showed increase in variability, whereas the value was lower for the grouped approach, i.e. 51%. Functional investigation of the differential variability showed that it is ubiquitous and was not concentrated in specific functional groups. That was further supported by the fact that as many as 74% to 94% of KEGG pathways included more genes with an increase in variability (Figure 5 A, B). We further asked if certain individuals with extreme ages cause an increase in variability with age. Analyzing the variability (absolute value of residuals) for each individual for the top 100 genes with the highest change in variability (ΔVar(ρ)) in either direction (Supplemental Figure 14, 15), we showed that the pattern is not caused by a limited number of individuals but is a general trend. Interestingly, the increase in variability was not restricted to samples with extreme ages but started as soon as the age of 40.

Most studies consider the accumulation of cellular damage, such as somatic mutations, with age as the main factor, causing increase in the gene expression variability with age. Indeed, Lodato et al [26] show increase in the number of single nucleotide variants in human brain with age, while Lee et al [27] documented somatic recombination of APP gene in human neurons and its increase with age. However, the causal link between the accumulation of mutations and increase in variability was not proven and Enge et al. [15] provide an evidence that somatic mutations are not enough to explain gene expression variability. Moreover, because brain is a post-mitotic tissue, it may demonstrate a different damage profile, as it is not as prone to replication-associated mutations as other tissues but associated with other types of damage, such as free radicals or loss of proteostasis. Notably, in this study we analyze different individuals; thus, the interpretation of the results could be different from the intra-individual variability which is observed across different cells from the same individual. If there are convergent mechanisms, more vulnerable cellular components, like hotspots, or transmission/propagation of signals that control the gene expression, interindividual heterogeneity could still reflect the changes at the cellular level. However, other explanations such as genotype differences, environment effects, and the difference in aging rates should be considered. A few studies have identified a small set of genetic variants that could change gene expression during aging (genotype-by-age interaction) [11,17,28]. However, these specific differences in genotype are not likely on their own to explain the transcriptome-wide shift that we observed. Still, we asked if the genes with a higher change in variability (ΔVar) are more or less tolerant to SNPs, using residual variation intolerance scores [29] which shows if a gene has more or fewer SNPs than expected (Supplemental Figure 13). There was no association between SNP tolerance and the change in variability, both when we considered and did not consider the direction of change (i.e. ΔVar and absolute value of ΔVar measures, respectively). The environmental factors influencing the epigenome, as well as stochastic effects driving an epigenetic drift [9,17,18,30] seems to be a likely explanation in this
case. Another explanation is that individuals age at
different rates and we see more variability in the rate at
older ages. However, in this case, the variability in
different genes would be driven by the same individuals
who age at different rates, which was not observed in
our analysis (Supplemental Figure 14-15). The change
in variability could also stem from the change in gene
expression levels. Although not replicated in mouse brain [14], Davie et al. [12] show that aging leads to an
overall decrease in the RNA content, which could also
be the reason for such a global increase in the
expression variability. However, we apply log2
transformation, which attempts to correct the mean-
variability dependence. Indeed, we do not observe any
significant association between the changes in
expression level and variability (Odds ratio = 1.05,
Fisher’s test p = 0.11, Supplemental Figure 10).

Although we used one of the largest, well-characterized
datasets, it is important to note that the sample size, the
unequal coverage of ages and the high technical and
biological variation all posed a challenge for the
analysis. Moreover, this data was generated using
microarray technology, which does not measure the
expression of all genes and is not as quantitative as
RNA-seq. Future studies addressing variability in gene
expression may consider the use of scRNA-seq data to
distinguish unique changes within a cell from the
coordinated changes within cell population or changes
in the cell composition. In order to gain insight into the
potential contribution of the changes in cell composition
to our results, we asked if there is an association
between cell-type specific and DV genes (Supplemental
Figure 12). We separated differentially variable genes
based on the direction of change in variability, approach
(grouped vs. continuous), and the correction strategy
(regression vs. SVA). There was a significant overlap
between genes with a decrease in variability and
myelinated oligodendrocyte-specific genes. Also,
oligodendrocyte progenitor- and neuron-specific genes
showed significant overlap with genes that increase in
variability, but this was not reproduced across different
approaches or correction strategies. This analysis,
however, cannot distinguish between if the change in
cell composition drives the change in variability or if
certain cell types become more variable with age; and
we need cell-type specific age-series data to answer this
question. Overall, the numbers of DV genes that
overlapped with cell type-specific genes were low, and
the changes in variability probably cannot be attributed
only to the changes in cell type composition.

Providing a systematic analysis of the same dataset at
multiple levels and considering multiple technical
challenges, we showed a slight but significant shift
towards an age-related increase in variability that was
not clustered in certain functions but distributed across
all pathways. It has been recently suggested that an
increase in expression variability is linked with the
 genetic risk for schizophrenia in males [31]. However,
future experiments are crucial to understanding whether
all genes, functions and organs are equally tolerant of
the variability we observed and whether this variability
has any causal relationship with the aging processes.

MATERIALS AND METHODS

Data processing steps

Dataset selection: We utilized one of the largest age-
series human brain expression datasets, featuring 269
prefrontal cortex samples from healthy individuals and
spanning the whole lifespan from development (prenatal
samples) through aging (80 years) [23]. These data were
collected using microarray technology from people of
both sexes and 4 races, namely African American (AA),
Caucasian (CAUC), Hispanic (HISP) and Asian (AS).
In the current analysis, we excluded fetal, childhood and
early adulthood samples before the age of 20, thus
limiting our sample size to 147. This was to exclude
developmental processes taking place in the brain until
the end of early adulthood, which exhibit discontinuous
expression changes between early adulthood and aging
[32]. Our main motivation was to study changes in gene
expression variability during aging, considering 20
years old as a starting point.

Data characterization: The pre-processed data (loess
normalization was applied on the background corrected
log2 intensity ratios (sample/reference) [23]); sample
and gene (probe set to Entrez gene mapping)
annotations were obtained from the NCBI Gene
Expression Omnibus (GEO) at accession number
GSE30272. Samples were processed in 19 batches, had
different quality measurements, namely pH and RNA
integrity number (RIN), and differed in the time of
collection after death (post-mortem interval (PMI)).
Using a PCA, we found no sample outliers as judged by
visual inspection of the first two principal components
(Supplemental Figure 2). However, the relationship
analysis between the above-mentioned factors (i.e.
batch, RIN, PMI and others) and age yielded significant
correlations for sex, post-mortem interval and RNA
integrity, pointing to potential confounders in the data
(Supplemental Figure 1). We further checked the
overlap between significantly differentially variable
genes in our analysis and previously reported genes that
are affected by PMI and detected only a limited overlap
(see Supplemental Information).

Probe set to Gene summarization: If one probe-set was
mapped to several genes, it was deleted to avoid
duplication. Conversely, when one gene had several
Batch correction: To compensate for technical variation between samples, quantile normalization (QN) was performed using the ‘normalize.quantiles’ function from the ‘preprocessCore’ R library. To differentiate between the age effect and the effect of the unwanted technical and biological variability, we have applied different expression correction strategies: linear regression of the known covariates, unsupervised estimation of covariates using surrogate variable analysis (SVA) [25,33], and ComBat, a parametric empirical Bayesian framework for covariate adjustment [24]. As a result, we analyzed the same data four times, corrected using QN, QN+regression, QN+SVA, QN+ComBat. Different corrections work by adjusting for the different covariates in the linear model that explains the gene expression, namely: i) QN – no covariates were added; ii) QN+regression – 25 covariates considered: technical batches (N = 19), sex (N=2), race (N=4), post-mortem interval, RNA integrity number, pH; iii) QN + SVA – 20 surrogate variables (SV) were inferred from the expression data using the ‘sva’ function from “SVA” R library; iv) QN+ComBat: the 6 confounding factors: batch (N = 19), sex (N=2), race(N=4), PMI, RIN and pH were adjusted for, one at the time, by repeatedly applying the ComBat function from the “SVA” R library to the expression data.

Differential expression

A least squares linear regression model was used to model gene expression level change with age. Age^{0.25} was used as an independent variable instead of age to account for the difference in the rate of gene expression changes between young (fast) and old (slow) as well as different density of the samples across ages. Nevertheless, the β_i coefficients from the linear model, that uses age^{0.25} correlate well with the one, that employs age (Supplemental Figure 4). Coefficients for the age covariate were used as a measure of the differential expression. P values for coefficients were adjusted using the FDR method with a threshold p ≤ 0.05 to account for multiple testing. Depending on the correction method applied, the linear model also accounted for different measured or unmeasured covariates (see Data processing steps) of the following general form:

\[ Y_i = \beta_{i0} + \beta_{i1} * \text{age}^{0.25} + \text{covariates} + \epsilon_i, \]

where \( Y_i \) is the normalized log-expression level of a gene with \( i = 1,…,n \). \( \beta_{i0} \) - intercept, \( \beta_{i1} \) - slope term and \( \epsilon_i \) - residual (or error) term.

Differential variability

The continuous approach: First, a linear model to fit gene expression during aging, using age^{0.25} and potential confounders, was constructed. Next, the Spearman correlation was calculated between the absolute values of the residuals, \(|\epsilon_i|\) from the linear model and age. Consequently, Spearman correlation estimates were used as a measure of the change in variability, referred as \( \Delta \text{var}(\rho) \). P values for the Spearman correlation estimates were corrected for multiple testing using FDR. FDR adjusted p ≤ 0.05 was used as a threshold to define significantly DV genes.

\[ \Delta \text{var}_i(\rho) = \text{Spearman} \rho(|\epsilon_i|, \text{age}) \]

The grouped approach: First, a corrected expression matrix was obtained by removing the effect of covariates (see data processing steps) from the data using the residuals from a linear regression model (\( Y_i = \beta_{i0} + \text{covariates} + \epsilon \)). The ‘grouped approach’ is a custom resampling-based test designed to compare gene expression variability between young (20 – 40 years old) and old (60-80 years old) groups using an interquartile range (IQR). IQR corresponds to the difference between the 75th and 25th percentiles of the distribution and is considered to be a robust measure of variability, meaning it is not susceptible to outliers and departure from normality in the data. In order to adjust for the unequal sample size of the young (N = 53) and old (N = 22) groups, we, first, calculated a null distribution of the IQR values for the young group by resampling it 10 000 times with the size of the old group. Next, we calculated significance as a percentage of samples where IQR_{old} was more extreme than IQR_{young} and corrected it for multiple testing using FDR correction, q ≤ 0.05. The ‘grouped’ measure of change in the variability, \( \Delta \text{var}(\text{IQR}) \), for the gene \( i \), corresponds to the difference between IQR value for the old, IQR_{i,old}, and IQR_{i,young} (i.e. mean IQR value from the young distribution), which is then divided by the latter, see formula:

\[ \Delta \text{var}_i(\text{IQR}) = \frac{(\text{IQR}_{i,old} - \text{IQR}_{i,young})}{\text{IQR}_{i,young}} \]

Gene Set Enrichment Analysis for KEGG pathways and GO categories

\( \beta_i \) coefficients from the differential expression and \( \Delta \text{var} \) measures from the differential variability analyses were used to perform gene set enrichment analysis, GSEA [34] using the “clusterProfiler” R library. KEGG pathways (N = 315) and BP GO terms (Biological processes Gene Ontology, N = 5822) with the size of
between 10 and 500 genes were considered as gene sets for the GSEA.

**Pathway distribution study**

KEGG pathway to gene mapping was obtained from “KEGGREST” R library and pathways were pre-filtered to contain between 5 and 500 genes. As a result, 310 KEGG pathways that comprise 5922 unique genes were used for the subsequent analysis. The boxplots illustrated distributions of the ∆var measure for genes in each pathway. Pathways were sorted according to their median ∆var measure in ascending order. The percentage of pathways that have their median ∆var above zero was calculated. The analysis was replicated using BP GO terms (N = 5919) of a size between 10 and 500 genes, which in total contained 12538 unique genes. Mapping of GO terms to genes was obtained from “org.Hs.eg.db” R library.

**Distribution tests**

Distributions of the ∆var - measures for all the genes were tested for normality using the Shapiro-Wilk test in R (‘Shapiro.test’ function) on the multiple subsamples, consisting of 5000 measures. Skewness of the distributions was calculated using the ‘fBasics’ function from “BasicStatistics” R library.

**Mean-variability relationship testing**

To visualize and test if the change in gene expression variability is associated with the change in gene expression level, we plotted the difference in the means between the young and old groups against difference in the interquartile range (IQR) between the young and old groups. Mean and IQR for the old group were calculated once, while mean and IQR for the young group were calculated 10,000 times for the subsamples (see Grouped approach) and then means of the distributions of the corresponding values (mean and IQR) were used in the analysis. Fisher’s exact test was performed on the values used for the plotting.

**Functional variation intolerance – variability relationship**

To test if there is a relationship between age-dependent change in variability and how tolerant a gene is to functional variations, we used residual variation intolerance (RVI) percentages based on ExAC v2 data [29]. RVI percentage shows whether genes have more or less common functional genetic variation relative to the genome wide expectation. A gene with a higher percentage has more common functional variation whereas a lower percentage shows the genes that are intolerant to functional variation. We tested the association using the Spearman’s correlation coefficient between the ∆var measures and RVI percentages, for different approaches and correction strategies, separately.

**Cell type-specific genes**

In order to test if the changes in cell type composition can drive the changes in heterogeneity, we tested the association between cell-type specific and DV genes. First, we analyzed a cell-type specific transcriptome dataset (GSE9566) [35] which has expression profiles for FACS purified cell types in the mouse brain. Data is downloaded from GEO database [36], RMA corrected using ‘affy’ package in R [37], log2 transformed, and quantile normalized using ‘preprocessCore’ package in R [38]. Only the genes with 1 to 1 human orthologs in Ensembl Compara [39] are used. We used ‘biomaRt’ package in R [40] to retrieve the mapping between probesets and genes. When there is more than one gene that map to a single probeset, we discarded those from the analysis. If multiple probesets represent one gene, we used the mean expression value of these probesets to calculate gene expression level. To define cell-type specific genes, for each human ortholog, we first standardized the gene expression values, and calculated the effect size of each cell type (oligodendrocytes, myelinated oligodendrocytes, oligodendrocyte progenitors, astrocytes and neurons) and identified genes with an effect size higher than 2 to a particular cell type. Using this cutoff, cell type-specific gene lists did not overlap. We next tested for the overlaps between these cell type-specific and differentially variable gene lists, using Fisher’s exact test.

**Software**

R version 3.5.0 and “data.table” were used to perform the analyses, while “ggplot2” and “ggpubr” R libraries were used to create visualizations of the data.

**Abbreviations**

BP GO: Biological Process Gene Ontology; DE: differentially expressed genes; DV: differentially variable genes; ∆var: the measure of change in the expression variability with age; GO: Gene Ontology; GSEA: Gene Set Enrichment Analysis; IQR: interquartile range; KEGG: Kyoto Encyclopedia of Genes and Genomes; PMI: post-mortem interval; QN: quantile normalization; RIN: RNA integrity number; Rho: Spearman correlation estimate; SVA: Surrogate Variable Analysis
AUTHOR CONTRIBUTIONS

H.M.D, V.R.K and J.M.T designed the study. V.R.K and H.M.D analyzed the data. V.R.K, H.M.D and J.M.T interpreted the results and wrote the manuscript. All authors read, revised and approved the final version of this manuscript.

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CONFLICTS OF INTEREST

The authors have no conflict of interest to declare.

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SUPPLEMENTARY MATERIAL

Please browse the links in Full Text version of this manuscript to see Supplementary Tables 1-7.

Supplemental Table 1. The results of the differential expression analysis performed on regression-corrected and SVA-corrected data.
Supplemental Table 2. The results of the GSEA Analysis for BP GO categories performed on the differentially expressed genes obtained on regression-corrected and SVA-corrected data.
Supplemental Table 3. The results of the differential variability analysis while applying “continuous approach” for the regression and SVA-corrected expression data.
Supplemental Table 4. The results of the differential variability analysis while applying “grouped approach” for the regression and SVA-corrected expression data.
Supplemental Table 5. The results of GSEA Analysis among the KEGG pathways on the differential variability measures from the "continuous" and "grouped" approaches.
Supplemental Table 6. The results of GSEA Analysis among the GO categories on the differential variability measures from the "continuous" and "grouped" approaches.
Supplemental Table 7. The median variability measures for the genes within each KEGG pathway and GO term, using different approaches.

Supplemental Information

Post-mortem interval associated differentially variable genes

Recent reports made on the GTEx data [1] suggest that extent of Post-mortem interval (PMI) – the time passed between death and sample collection, could be associated with the degradation of specific mRNA species and, consequently, change in both mean and variability of their gene expression. We have checked whether our set of differentially variable genes (DV) contains any of the reported 266 genes that are susceptible to the degradation dependent on PMI. We found 14, 14, 10 and 7 of those genes among the set of differentially variable genes identified in the grouped approach with quantile normalization, regression, ComBat and SVA correction, respectively.

Overlap of the DV genes with other publications

The total set of differentially variable genes identified by Brinkmeyer-Langford [2] comprises 848 distinct genes found across 13 brain regions. 171 of those genes are specific to the cortex and frontal cortex. Our set of DV genes found in grouped approach after regression correction (N total = 741) has an overlap of 21 genes with Brinkmeyer-Langford and 19 genes for the SVA-corrected expression (N total = 746). The percentage of overlap doesn’t increase, if we only consider cortex-specific DV genes from Brinkmeyer-Langford - it constitutes 4 and 3 genes for regression and SVA corrected sets, respectively.

Supplemental References

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Supplemental Table 8. Percentage of KEGG pathways and GO terms that have median expression variability above 0, according to different approaches.

|                     | Continuous approach |         |         | Grouped approach |         |         |
|---------------------|---------------------|---------|---------|------------------|---------|---------|
|                     | Regression          | SVA     | QN      | ComBat           | Regression| SVA     | QN      | ComBat |
| KEGG pathways       | 94%                 | 88%     | 99%     | 88%              | 77%      | 74%     | 94%     | 95%    |
| GO terms            | 92%                 | 84%     | 98%     | 79%              | 66%      | 61%     | 88%     | 88%    |
Supplemental Figures

Supplemental Figure 1. Dependency plots between the age and different biological and technical factors, such as batch, sex, race, post-mortem interval (PMI), pH and RNA integrity number (RIN). Box plots were constructed for categorical covariates (batch, sex and race), while scatter plots were made for the numerical covariates (PMI, pH, RIN). In order to test for the significance of the dependence between categorical covariates and age ANOVA-test was performed (p value is displayed on the graph), Spearman correlation estimate was calculated to estimate relationship between numerical covariates and age (Spearman rho and p value are displayed on the graph). * p ≤ 0.05
Supplemental Figure 2. PCA plots of the raw expression data and after different pre-processing steps. First row from left to right: probe expression level, gene expression level and quantile normalized gene expression level. Second row from left to right: quantile normalized data further corrected either with regression, ComBat or SVA. Color gradient depicts age range from pink – young to violet – old.
Supplemental Figure 3. (A) Illustration of the relationship between beta1 – coefficients from the linear regression models that explain gene expression corrected with different approaches (QN only, QN followed by regression, ComBat, SVA) using age 0.25. Scatter plots that illustrate relationship between the beta1 – coefficients from different corrections are displayed in the lower triangle of the matrix, distributions of the beta1 – coefficients are located on the diagonal and Spearman correlation estimates are displayed in the upper triangle of the matrix. (B) Venn diagram of the differentially expressed (DE) genes with age that were identified after applying different expression correction approaches (QN only, QN followed by regression, ComBat, SVA). The ComBat approach seem to greatly outnumber all the other approaches in the number of DE genes. We consider that consecutive application (6 times) of the linear model that contains age and particular confounding factor caused an overfitting of gene expression with age and lead to identification of so many DE genes, many of which are false positives.
Supplemental Figure 4. Scatter plots of the dependency between beta1 coefficients from the linear regression models that explain gene expression using age in comparison to age^{0.25}. Gene expression was corrected using ComBat (A), QN only (B), regression (C) and SVA (D). Linear regression line between the variables (in red) and Spearman correlation estimate is shown on the graph.
Supplemental Figure 5. Comparison of the distributions of the differential variability measures, \( \Delta \text{var}(\rho) \) and \( \Delta \text{var}(\text{IQR}) \), obtained in the expression data corrected with QN, Regression, SVA and ComBat. The black straight line depicts zero, the red dashed line – median of the distribution.
Supplemental Figure 6. Combination of comparisons of the variability measures (Δvar) obtained in the continuous approach between different ways of data pre-processing. Correction with quantile normalization only, linear regression, SVA and ComBat. Scatter plots between the differential variability measures obtained after different corrections are displayed in the lower triangle of the matrix, distributions of the differential variability measures are located on the diagonal and Spearman correlation estimates between the variability measures from different corrections are shown in the upper triangle of the matrix.
Supplemental Figure 7. Combination of comparisons of the variability measures (Δvar) obtained in the grouped approach between different ways of data pre-processing. Correction with quantile normalization only, linear regression, SVA and ComBat. Scatter plots between the differential variability measures obtained after different corrections are displayed in the lower triangle of the matrix, distributions of the differential variability measures are located on the diagonal and Spearman correlation estimates between the differential variability measures from different corrections are shown in the upper triangle of the matrix.
Supplemental Figure 8. Venn diagram of the differentially variable genes found in the grouped approach when expression data was corrected with QN, linear regression, SVA and ComBat.

Supplemental Figure 9. Scatter plots of the relationship between the change in the variability measures for the continuous and grouped approaches in data, corrected with QN, ComBat, regression and SVA corrections. Linear regression line between the variables (in red) and Spearman correlation estimate are shown on the graph.
Supplemental Figure 10. Relationship between the change in gene expression level and variability. x axis depicts difference between mean of the old group and mean of the young group distribution (generated by resampling young group with sample size of 22 for 10,000 times and calculating the mean), y axis illustrates difference between interquartile range (IQR) for the old and mean IQR from the young distribution (generated by resampling young group with sample size of 22 for 10,000 times and calculating IQR).

Supplemental Figure 11. Scatter plots that illustrate relationship between the pathway membership of a gene and its change in variability measure for the continuous and grouped approaches in data corrected with regression and SVA. Linear regression line between the variables is shown red on the graph.
Supplemental Figure 12. A heatmap showing the significant association between differentially variable and cell-type specific genes. Rows show different cell-types and columns show genes detected as differentially variable based on different approaches (grouped (gr_) or continuous (cont_)), correction methods (regression (regres_) or SVA) and direction of change (increase (u) or decrease (d)). The color shows log2(odds ratio) for the enrichment of a particular category in cell-type specific genes. All non-significant (p > 0.05) values are discarded and shown as white. Since there are no significantly differentially variable gene detected based on continuous approach, we took the first 1% of genes with the highest change in variability as the DV genes for this analysis. The number of genes tested for each cell type were as follows: oligodendrocyte progenitors 396, oligodendrocytes 64, astrocytes 21, myelinated oligodendrocytes 448, neurons 340.

*Non significant values are also shown in white
Supplemental Figure 13. Scatter plot showing the association between the change in variability with age (y-axis) and residual variation intolerance (RVI) percentage (x-axis), which shows whether genes have more or less common functional genetic variation relative to the genome wide expectation. A gene with a positive score has more common functional variation whereas a negative score shows the genes that are intolerant to functional variation. The scores are than assigned percentiles in decreasing order of scores, so that a higher score (genes having more common functional variation) is given a higher percentile. Each point shows a gene and the black line is the best fit line depicted for visualization purposes. The correlation between change in variability with age and RVI percentage is calculated using Spearman’s correlation and the coefficient is given for each plot. The same analysis was repeated for the absolute value of the change in variability and also only using the significantly DV genes and the results are comparable with Spearman’s
Supplemental Figure 14. A heatmap visualizing the variability for each individual (columns), and top 100 genes with the highest change in variability in either direction (rows) based on continuous approach and regression correction. The color shows the rank of individuals with respect to their variability for that particular gene. Darker colors show individuals with the highest variability for that particular gene. Genes are clustered using hierarchical clustering and individuals are ordered by age.
Supplemental Figure 15. Same as Supplemental Figure 14 but for continuous approach using SVA correction.