Alterations of Mitochondrial DNA Copy Number and Telomere Length with Early Adversity and Psychopathology

Supplemental Information

Unadjusted Means (Raw values, prior to log transformation)

Mitochondrial DNA Copy Number. Unadjusted means were M=231.8, SD=121.9 for the No Adversity/No Disorder group; M=260.1, SD=117.7 for the Adversity/No Disorder group; M=286.3, SD=140.8 for the No Adversity/Disorder group; and M=281.3, SD=127.2 for the Adversity/Disorder group.

Unadjusted means for individual disorders were M = 269.1, SD = 113.9 for the group with lifetime MDD; M = 274.3, SD = 129.7 for the group with any lifetime depressive disorder; M = 283.8, SD = 128.3 for the group with lifetime PTSD; M = 294.3, SD = 122.7 for the group with any lifetime anxiety disorder; and M = 280.4, SD = 108.4 for the group with past alcohol or substance disorders.

Unadjusted means for individual types of adversity were M=296.3, SD=128.0 for the group who experienced parental loss, and M=270.6, SD=126.3 for the group with any form of maltreatment on the CTQ.

Telomere Length. Unadjusted means for the Adversity/Disorder grouping variable were M = 7181.00, SD = 4362.80 for the No Adversity/No Disorder group; M = 6432.9, SD = 3372.7 for the Adversity/No Disorder group; M = 5854.1, SD = 4250.9 for the No Adversity/Disorder group; and M = 5642.7, SD = 4138.2 for the Adversity/Disorder group.

Unadjusted means for the individual disorders were M = 4933.0, SD = 3189.2 for the group with lifetime MDD; M = 4855.5, SD = 3525.5 for the group with any lifetime depressive disorder; M = 5064.7, SD = 2616.6 for the group with lifetime PTSD, M = 4802.8, SD = 2960.1 for the group with any lifetime anxiety disorder; and M = 6893.3, SD = 4216.0 for the group with past alcohol or substance disorders.

Unadjusted means for the early adversity types were M=6482.9, SD=3932.3 for the group who experienced parental loss and M=6034.1, SD=3784.0 for the group with any form of maltreatment on the CTQ.

General Linear Models (GLMs)

The following tables show the results of the full GLMs. Standardized regression coefficients, B, are shown to display the relative contribution of variables in the model.

Table S1. General linear model predicting mitochondrial copy number

Source	df	F	В	p
Adversity/Disorder Grouping	3	8.40	.28	.00
Telomere Length	1	42.59	.36	.00
Age	1	2.23	09	.14
Gender	1	.01	.00	.95
Childhood Socioeconomic Adversity	1	.16	02	.69
Education	1	1.25	.06	.26
BMI	1	.02	01	.90
Error	280			

Table S2. General linear model predicting telomere length

Source	df	$\boldsymbol{\mathit{F}}$	В	p
Adversity/Disorder Grouping	3	6.39	25	.00
Mtdna Copy Number	1	42.59	.36	.00
Age	1	.62	04	.43
Gender	1	.22	.03	.64
Childhood Socioeconomic Adversity	1	1.26	07	.26
Education	1	.00	.00	.99
BMI	1	2.34	08	.13
Error	280			

Sensitivity Analyses

The following analyses did not control for the significant positive association of telomere length and mtDNA copy number.

Mitochondrial DNA Copy Number. When predicting mtDNA copy number without controlling for telomere length, the grouping variable was still a significant predictor of mtDNA copy number, but the Adversity/No Disorder group differed from the controls only at trend level (p = .069), and the Adversity/No Disorder group was not different from the Adversity/Disorder group (p = .193) or No Adversity/Disorder group (p = .245). The findings for individual disorders and adversity types all remained significant.

Telomere Length. When predicting telomere length without controlling for mtDNA copy number, the findings remained significant except that the effect for the No Adversity/Disorder group was reduced to trend level (p = .057), as was the difference between the Adversity/No Disorder group and the Adversity/Disorder group (p = .083). Findings for individual disorders remained significant except for lifetime PTSD (p = .100). Parental loss was associated with telomere length at trend level (p = .056) and the effect of any form of maltreatment on the CTQ was no longer significant (p = .127).