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## Evaluating mitochondrial DNA variation in autism spectrum disorders

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### SUMMARY

Despite the increasing speculation that oxidative stress and abnormal energy metabolism may play a role in Autism Spectrum Disorders (ASD), and the observation that patients with mitochondrial defects have symptoms consistent with ASD, there are no comprehensive published studies examining the role of mitochondrial variation in autism. Therefore, we have sought to comprehensively examine the role of mitochondrial DNA (mtDNA) variation with regard to ASD risk, employing a multi-phase approach.

In phase 1 of our experiment, we examined 132 mtDNA single-nucleotide polymorphisms (SNPs) genotyped as part of our genome-wide association studies of ASD. In phase 2 we genotyped the major European mitochondrial haplogroup-defining variants within an expanded set of autism probands and controls. Finally in phase 3, we resequenced the entire mtDNA in a subset of our Caucasian samples (~400 proband-father pairs). In each phase we tested whether mitochondrial variation showed evidence of association to ASD. Despite a thorough interrogation of mtDNA variation, we found no evidence to suggest a major role for mtDNA variation in ASD

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#### Author contributions

JLM, JLH, and MPV designed the study. JLM, AH, ERM led the writing and revising of the manuscript. PLW, IK, DJH, JG performed and directed all molecular work including the genotyping and mtDNA resequencing efforts. AH, JLM, MS, and ERM performed the QC and statistical analysis. HHW, RKA, MLC, JLH, and MPV were involved in recruiting autism families and controls for this study. RKM and SMW provided control sample DNA. All authors have read and contributed to the manuscript.

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susceptibility. Accordingly, while there may be attractive biological hints suggesting the role of mitochondria in ASD our data indicate that mtDNA variation is not a major contributing factor to the development of ASD.

## Keywords

mitochondrial DNA; autism; autism spectrum disorders; association studies; genetic

## INTRODUCTION

Autism Spectrum Disorders (ASDs) are neurobehavioral disorders characterized by deficits in social abilities, problems with language and communication, and the presence of patterns of repetitive behaviors, restricted interests, and resistance to change. ASD has an estimated population prevalence of one in every 1000 individuals within the general population with a male:female ratio of 4:1 (Fombonne, 2002; Fombonne, 2009). Autism, along with Asperger's syndrome, Rett syndrome, and other pervasive developmental disorders, are more generally classified as autism spectrum disorders (ASDs) and may affect as many as one in 88 children in the United States (Autism and Developmental Disabilities Monitoring Network Surveillance Year 2008 Principal Investigators, 2012). Little is known about the etiology of ASD; however overwhelming evidence from numerous studies has indicated that idiopathic autism has a complex genetic etiology. Twin and sibling studies overwhelmingly suggest a strong genetic component and high heritability for ASD. Studies show a concordance rate of ~60% for classic autism and up to ~90% for ASD among monozygotic (MZ) twins and <10% among dizygotic (DZ) twins (Folstein & Piven 1991; Ritvo et al., 1985; Bailey et al., 1995). Numerous linkage and association studies, including genome-wide association studies (GWAS), and candidate gene studies have failed to characterize an appreciable amount of the genetic variation believed to be involved in this devastating disease. These approaches have identified a multitude of possible locations and genes for susceptibility, but few consensus regions or genome-wide significant associations have resulted (Anney et al., 2010; Autism Genome Project Consortium et al., 2007; Ma et al., 2009; Wang et al., 2009; Weiss et al., 2009). Recently, Copy Number Variants have been revealed to explain some of the variation in ASD, strengthening the hypothesis that multiple sources contribute to ASD etiology (Salyakina et al., 2011; Griswold et al., 2012)

The mitochondrial genome is small and circular (16,569 base pairs), possesses a distinct code from the nuclear genome (Wallace et al., 1999) and has a unique maternal inheritance pattern. Multiple copies of the mitochondrial DNA (mtDNA) are contained in each mitochondrion; some differ in sequence, this phenomenon is called 'heteroplasmy'. It encodes 13 protein subunits of the mitochondrial electron transport chain and a distinct set of rRNAs and tRNAs all of which are critical for life-sustaining oxidative phosphorylation and energy generation (Wallace, 1994). Relative to the nuclear genome, the mitochondrial genome has been understudied in the search for common genetic variation associated to human disease, despite the fact that mitochondria play a vital role in cellular energy production (Wallace et al., 1999; Papa, 1996; Wallace, 1997).

Variation in mitochondrial DNA has been examined in numerous neurological, age-related common genetic diseases (Howell et al., 2005; Raule et al., 2007; Canter et al., 2008; van der Walt et al., 2003; Rollins et al., 2009). Increased production of reactive oxygen species (ROS) due to mitochondrial respiratory activity and the resultant damage to both mtDNA and nuclear DNA have long been implicated in disease (Wallace et al., 1999; Kang & Hamasaki 2003; Scheffler, 2001; Feig et al., 1994; de Zwart et al., 1999; Penta et al., 2001). There has been increasing speculation that oxidative stress and abnormal energy metabolism

may play a role in ASD, consistent with some level of mitochondrial dysfunction (Lombard, 1998; Chugani et al., 1999; Chauhan & Chauhan 2006; Clark-Taylor & Clark-Taylor 2004; Oliveira et al., 2005; Rossignol et al., 2007), although this remains somewhat controversial (Lerman-Sagie et al., 2004). In addition, numerous clinical reports have described patients with mitochondrial disorders or mutations who have symptoms consistent with ASD (Oliveira et al., 2005; Poling et al., 2006; Fillano et al., 2002; Tsao & Mendell 2007; Filipek et al., 2003; Graf et al., 2000; Pons et al., 2004). Further, mitochondrial inheritance is consistent with the observed increased neuropsychological abnormalities in the mothers of ASD children (Baron-Cohen et al., 1997; Bishop et al., 2004; Constantino & Todd 2005). To date there has been one investigation of the mitochondrial haplogroups in ASD (Kent et al., 2008). The small sample size (n=162) generated borderline significant results, but the study was underpowered to detect anything but large main effects with odds ratios >2.0.

Additional evidence for the role of mitochondria in autism stems from data regarding the *SLC25A12* gene, a nuclear encoded protein for the mitochondrial aspartate/glutamate transporter, ARALAR (De Zwart et al., 1999; Satrustegui et al., 2007)). Mice with a homozygous deletion of the *Aralar* gene develop severe birth defects soon after birth and die approximately 20 days postnatal. In addition, aralar deficiency causes a large drop in aspartate and its derivative n-acetylaspartate (NAA) in the brain and in primary neuronal cultures. Interestingly, NAA is commonly used in <sup>1</sup>H-NMR spectroscopy as a potential diagnostic marker for neuronal function or loss (Pan & Takahashi 2005; Tsai & Coyle 1995), and is reduced in certain brain regions of autistic patients (Otsuka et al., 1999). In humans, two polymorphisms within *SLC25A12* were associated with an increased risk of autism in a dataset of 411 families (Ramos et al., 2004). A positive replication was found in a relatively homogeneous dataset of Irish autism families (Segurado et al., 2005), but negative reports in more heterogeneous datasets (Rabionet et al., 2006; Blasi et al., 2006; Correia et al., 2006) have also been published. There are several possible explanations for the inconsistency of results, including underpowered datasets and genetic heterogeneity, which could arise from having differing mitochondrial haplogroup backgrounds that may affect *SLC25A12* function.

In summary, the potential role of mitochondrial variation in ASDs remains intriguing and warrants thorough investigation. While the nuclear genome has been the focus of countless studies over the past couple of decades in search for autism susceptibility genes, our study marks the largest examination of the mitochondrial genome in ASD, and the first report of mtDNA resequencing in ASD.

## MATERIALS AND METHODS

### Patient Ascertainment and Description

Individual patient samples included in this study (n=1298) (Table 1) consist of samples ascertained at the John P. Hussman Institute for Human Genomics (HIHG) at the University of Miami, Miller School of Medicine (Miami, Florida) (n=668), the University of South Carolina (Columbia, South Carolina) (n=317), the Center for Human Genetics Research at the Vanderbilt University (Nashville, Tennessee) (n=108), and samples obtained from the Autism Genetic Resource Exchange (AGRE) (n=205) (Autism Genetics Resource Exchange, 2008). Families were enrolled through a multi-site genetics study of autism and recruited via family support groups, advertisements, and clinical and educational settings. All participants were ascertained and sampled according to approved Institutional Review Board (IRB) protocols. Participants with ASD met the following minimum criteria for inclusion: (1) chronological age between 3 and 21 years of age; (2) presumptive clinical diagnosis of ASD; (3) expert clinical determination of ASD diagnosis using DSM-IV criteria supported by the Autism Diagnostic Interview (ADI-R) (Rutter et al., 2003b).

Diagnostic determination was based on review by clinical psychologists with extensive experience in autism and related disorders. In those instances where an ADI-R was not available, a best-estimate diagnosis was assigned using all available clinical information including clinician summaries, caregiver report, and medical records; (4) minimal developmental level of 18 months as determined by the Vineland Adaptive Behavior Scale (VABS) (Sparrow et al., 1984) or the VABS-II (Sparrow et al., 2005) or an IQ equivalent > 35. These minimal developmental levels assure that ADI-R results are valid and reduce the likelihood of including individuals with severe mental retardation. We excluded participants with severe sensory problems (e.g., visual impairment or hearing loss), significant motor impairments (e.g., failure to sit by 12 months or walk by 24 months), or identified metabolic, genetic, or progressive neurological disorders. Family history and pedigree information (including any known health and psychiatric history of family members) was collected in a standard semi-structured interview with a biological parent of the proband, frequently the mother. Phenotypic data regarding the family also was collected through a review of available medical and psychiatric records of the proband and/or affected sibling, as well as review of available photographs of the proband, siblings and parents in the patient charts. Confounding by race and ethnicity was addressed using both a stratified analysis and a principal components analysis (PCA) in phase 1, haplogroup definition in phase 2, and a homogeneous sample of self-reported Caucasian non-Hispanic individuals in phase 3.

### Control Ascertainment and Description

Control samples utilized in this study (n=2,646) were obtained from multiple sources (Table 1). Healthy children (n=513) between the ages of 4 and 21 years were recruited by the HHG. Participants were screened for eligibility using a series of preliminary questions to determine whether the child, his or her parent, or sibling has been diagnosed a developmental, behavioral, neurological or other disability or condition. If none of those conditions were present, parents of minor children or participants reviewed and signed the informed consent and completed the Social Communication Questionnaire (Rutter et al., 2003a) to screen for potential ASDs. These control participants provided a saliva sample for these and other ongoing genetic studies. A second set of control individuals (n=327) were part of ongoing studies of preterm birth. These samples were collected via the Centennial Medical Center in Nashville from the cord blood of term pregnancies (> 37 weeks gestation). Mothers between the ages of 18 and 40 were recruited for this collection, with cord blood being collected from live singleton births. Our third set of control individuals (n=582) were from the National Institute of Mental Health (NIMH) Human Genetics Initiative and consist of non-Hispanic European-ancestry DNA samples made available through this resource. These samples and their corresponding data have been used by multiple investigators through permission and collaboration with the Center for Collaborative Studies of Mental Disorders resource (<https://www.nimhgenetics.org>). Finally, given that mtDNA is maternally inherited, we included the unaffected fathers (n=1,224) from the ascertained autism families (n=614 HHG; n=290 South Carolina; n=106 Vanderbilt; n=214 AGRE) as additional controls for these experiments.

### Phase 1 Dataset (Illumina Genotyping)

**Description and Genotyping**—A total of 2727 samples (964 cases, 1763 controls) were included in the phase 1 experiment (Table 1; Fig. 1). Cases were selected from among our ASD families, with a single case selected (probands were chosen preferentially) from each nuclear family. Additional cases were chosen from 76 extended multiplex families provided they did not share mitochondrial lineage (i.e. each case chosen had a unique maternal founder). Control data came from the fathers of probands (n=923) and the pediatric controls from the HHG and the preterm birth study (n=840).

**Analysis**—Individual sample data was selected for cases and controls following a comprehensive quality control (QC) analysis of these samples using the autosomal markers of the GWAS panel as previously reported (Ma et al., 2009; Wang et al., 2009). Of the total 163 mtDNA SNPs genotyped across both the Illumina 1M and Illumina 1M Duo BeadChip arrays, we examined the 132 common to both platforms. These SNPs were examined for call rate (requiring >95% for inclusion). Using the PLINK analysis software package, we next examined the call rate of samples and dropped any sample with a call rate below 95% in this set of SNPs (Purcell et al., 2007). We further checked both samples and SNPs by examining inconsistencies between mother and child genotypes at these 132 mtDNA SNPs with the use of the PLATO software package (Grady et al., 2010). This resulted in the removal of one case sample. Due to the insensitivity of the assay to detect heteroplasmy we set all heterozygous calls to missing.

To test for association while accounting for possible confounding by population stratification, we performed a stratified analysis, using the Cochran-Mantel-Haenszel (CMH) test, as implemented in PLINK, with genetically defined clusters generated with the software program CLUSTER (Table S1, Fig. S1). CLUSTER was developed in-house as an alternative to the STRUCTURE software application (Pritchard et al., 2000). It uses Ward's clustering algorithm to assign individuals to populations on the basis of information from multiple loci. In addition, to address the issue of population stratification on a finer scale, we conducted Eigenstrat analysis using the 132 mtDNA SNPs (Biffi et al., 2010; Price et al., 2006). This PCA was used to infer continuous axes of genetic variation which control for ancestry in the place of the categorical self-reported ethnicity variable. Eigenstrat analysis resulted in the exclusion of 49 samples with eigenvector values that were  $\geq 6$  standard deviations from the mean of principal components 1, 2 and 3 (mtPCs). The mitochondrial genomic inflation factor (mtGIF) was used as a measure of deviation from the median of the test statistic distribution. Association analysis was performed using logistic regression as implemented in PLINK, with mtPC1, mtPC2 and mtPC3 used as covariates in the analysis (Table 2). The mtGIF=1.0 with the inclusion of these principal components. We did not attempt to incorporate autosomal data to further correct for mitochondrial population substructure as data suggests it results in little improvement (Biffi et al., 2010). Furthermore, we conducted permutation testing to assess the significance of our results, using PLINK (--mperm 10000 --model-trend options) on the self-reported Caucasian non-Hispanic subset of the phase 1 dataset.

## Phase 2 Dataset (Sequenom Genotyping)

**Description and Genotyping**—A total of 2459 samples (818 cases, 1641 controls) were included in the phase 2 experiment (Table 1; Fig. 1), only five of which overlap with phase 1. Cases were selected as in phase 1 from among our ASD families (n=613), with additional cases from the AGRE resource (n=205). Control data for this experiment came from the mitochondrially unrelated fathers of probands (n=806), from the cord blood of the term pregnancies (n=253), and from the NIMH Human Genetics Initiative (n=582). In phase 2 of our experiment, we used both the Sequenom MassARRAY iPLEX and TaqMan genotyping platforms to genotype the major European mitochondrial haplogroup defining variants (Table S2). A total of 12 SNPs were genotyped in this effort for the main purpose of defining these haplogroups within our dataset.

**Analysis**—Cases and controls were chosen for analysis following a QC approach similar to phase 1. As we genotyped our entire ASD dataset on the 12 selected SNPs, our QC analysis in this phase benefited from mother-child as well as cross-platform genotype concordance. Due to the limited number of markers and their importance in haplogroup assignment the sample call rate threshold was set to 100%. After sample and marker checks, any remaining



instances of erroneous heterozygous genotype calls or platform discordant genotypes were set to missing for our analysis. Each sample was assigned to a specific haplogroup (Table 3) using information from the 12 genotyped SNPs (Table S2). Logistic regression analysis was conducted using SAS for both haplogroup and single marker tests of association (Table 4).

### Phase 3 Dataset (Affymetrix Genotyping)

**Description and Genotyping**—A subset of Caucasian samples was selected for resequencing of the entire mitochondrial genome (Table 1; Fig. 1). Before QC, the case group consisted of 400 samples that represented one affected individual per family (typically the proband), and the control group consisted of the 400 fathers of these individuals. We utilized the Affymetrix Human Mitochondrial Resequencing Array 2.0. (Affymetrix Inc. Santa Clara, CA) to resequence the entire mtDNA in this sample subset. This array uses microarray chip technology to sequence both strands of the entire mtDNA sequence after performing three long-range PCR amplifications. Each base position is interrogated with eight unique 25-mer probes on the resequencing array, and allows for the detection of both known and novel base substitutions.

**Analysis**—Haploid calls were initially made by setting the Affymetrix GSEQ software algorithm parameters to the haploid model and the quality score threshold to 12 (Coon et al., 2006). A summary of the sequencing results for each sample was generated with MSDAT, a powerful tool developed by our group for the analysis of mtDNA sequence data. Call rate thresholds were set at 95% for both samples and SNPs. As a result, 842 out of 16,544 positions and 49 out of 800 samples were dropped using the PLATO software package (Grady et al., 2010). Additional base positions were dropped depending on the analysis, with both tri-allelic ( $n=29$ ) and monomorphic base calls ( $n=14,732$ ) being dropped as part of the single marker analysis. In this phase, maternal genotypes were unavailable for conducting additional mitochondrial error checks. The sample call rate threshold reduced the final sample counts to 372 controls and 379 cases. We assessed significance by permutation testing for 183 common ( $MAF>0.01$ ), variants, and separately for 941 variants (no MAF filter) using the program RVASSOC (Kinnamon, 2010) which implements Cochran-Armitage (CA) max/sum tests (Kinnamon et al., 2012). Moreover, we tested for coding variation (Table 5) and heteroplasmy differences, examined our dataset for presumably rare variation that has been shown to be associated with other mitochondrial disorders (Table 6), and performed mutational burden assessments between our cases and controls. We qualitatively assessed the differences in heteroplasmic SNP distributions between cases and controls (Cutler et al., 2001) using the diploid GSEQ algorithm settings, a quality threshold of 3 and a call rate threshold of 95%. We were able to examine 16,010 out of the 16,544 mitochondrial positions. We followed the approaches set forth by Coon and colleagues to examine both heteroplasmy and mutational burden (Coon et al., 2006). Total mutational burden was calculated as the number of variants observed corrected for the number of sites/positions examined, multiplied by the number of cases or controls (Table 7).

**Additional QC across all phases**—The existence of overlapping sample and marker datasets made a cross-platform genotype concordance QC measure possible. This validation step provided additional confidence and validation of our genotype calls across the different platforms and phases of this project. Examining only non-missing data and comparing phase 1 with phase 2 data (6371 pair-wise comparisons across 1276 common samples and five common SNPs), yielded a genotype concordance rate of 0.998. Evaluation of phase 1 with phase 3 data considered 713 common samples and 132 common SNPs with a total of 92,516 pair-wise comparisons and yielded a concordance rate of 0.999. Finally, examination of phase 2 with phase 3 considered 6340 comparisons between 534 common samples and 12 common SNPs returned a concordance rate of 1.0.

## RESULTS

In phase 1, we had >80% power to detect a genotype relative risk (GRR) of 1.3 for any single SNP, with a type 1 error rate of 0.05 and an allele frequency of 0.10. The small number of individuals with African ancestry in our dataset significantly decreases the power for this subset (12% power for GRR of 1.3,  $\alpha=0.05$ , MAF=0.10). Our stratified CMH test identified six nominally significant variants ( $p < 0.05$ ) that do not survive Bonferroni correction (Table S1). Using a continuous variable to control for stratification in a logistic regression framework, we identified six SNPs with nominal p-values ( $p < 0.05$ ) through our single marker tests of association (Table 2; Fig. 2; Table S3). These nominal associations do not survive either Bonferroni correction or our less conservative assessment via permutation testing using RVASSOC in the homogeneous Caucasian non-Hispanic subset of phase 1 (observed max  $\chi^2=4.81$ ,  $p=0.93$ ). Of note, four of these six nominally significant variants overlap the stratified and logistic regression approaches; out of these four overlapping variants, two (mtDNA positions 9899 and 10589) tag subgroups of haplogroup L.

In phase 2, we examined both the major European haplogroups and European haplogroup defining SNPs for association to ASD susceptibility. Our haplogroup analysis yielded no significant difference in the frequency of cases versus controls for any particular haplogroup (Table 3). There are too few SNPs in this phase to control for population substructure using the principal components approach we used in phase 1. Instead, we used the haplogroups as covariates in a logistic regression analysis. We see no significant difference between our cases and controls for any of these single haplogroup defining SNPs (Table 4). Interestingly, individuals with self-reported African ancestry belong to a number of haplogroups other than haplogroup L.

Given that the samples in phase 3 are almost completely overlapping with those genotyped in phase 1 and 2, we chose not to perform single marker association tests of the 139 SNPs previously examined in those phases within this dataset. The advantage of this phase rests in capturing rare variants which cannot be powerfully tested with single marker tests of association. We confirmed the homogeneity of the phase 3 dataset which was based on self-report using the mtGIF calculated with  $\chi^2$  test statistics from all 941 polymorphic positions. Subsequently, we performed a joint test of these variants, many of which have MAF  $< 0.01$  and are spread across the mitochondrial genome; this returned no significant results (observed max  $\chi^2=6.34$ ,  $p=0.70$ ).

Furthermore, we examined whether we detected more rare variation in total among cases when compared to controls specifically in the coding regions of the 13 protein subunits of the mitochondrial electron transport chain. We found no significant difference between our cases and controls for either the number of synonymous or non-synonymous changes in these genes (Table 5).

We specifically analyzed our phase 3 dataset to determine if any of our samples contained any of these rare variations as reported and organized by the MitoMap project (<http://www.mitomap.org>). We identified 25 variants which were not previously examined in phases 1 or 2 that were in the MitoMap tables of 'mtDNA Mutations with Reports of Disease-Associations' from <http://www.mitomap.org> (Table 6). These tables contain any mtDNA variant that is reported in the literature of disease associations, but not necessarily replicated in subsequent investigations. A single variant (A13637G), previously associated to Lieber's Hereditary Optic Neuropathy (LHON), demonstrated a nominally significant p-value before multiple testing correction ( $p=0.02$ ) using the Fisher's Exact test. This rare allele was found in eight ASD cases and a single control.

Finally, we examined our phase 3 dataset for evidence of heteroplasmy and mutational burden differences between cases and controls. We examined the mutational burden in a specific subset of the variants detected, however failed to find any significant difference between our ASD case and control datasets (Table 7). No significant difference was found between cases and controls in the heteroplasmy analysis using individual raw intensity allele data (data not shown).

## DISCUSSION

In recent years, there has been considerable speculation that mitochondrial variation may play a role in autism spectrum disorders. We performed the first comprehensive investigation of this hypothesis. Because of common ancestry and maternal inheritance of mtDNA, the vast majority of humans can be assigned to a known haplogroup that arose during ancient migrations. Approximately 40% of Caucasians of European descent belong to haplogroup H, and the total prevalence of the next most common mitochondrial haplogroups (I, J, and K) is approximately 25% (Torroni et al., 1994). Although most persons of African descent belong to haplogroup L, this group is extremely diverse and can be divided into many sub-haplogroups (Chen et al., 1995). This genetic variation results in distinctive sets of human mitochondrial electron transport chains with different capacities for energy production, free radical generation and apoptosis (Swerdlow et al., 1996; Wallace, 2005). Despite the initial sequencing of the mitochondrion in the 1980s, routine full mitochondrial sequencing has been prohibitively expensive and thus the full complement of variation has not been routinely examined in any disease.

In each of the phases of this study, we searched for whether we find evidence of association of mitochondrial variation in ASD. While we do not detect strong evidence for significant main effects of any single mtDNA variation in our current dataset, we identify only a handful of variants demonstrating a mild level of significance. Furthermore, with our substantially larger dataset than the one explored by Kent and colleagues (Kent et al., 2008), we find that there is no compelling evidence that European mitochondrial haplogroups influence risk of developing ASDs. Our inability to detect a significant haplogroup association is not surprising based on the work by Samuels and colleagues noting that the reliable detection of haplogroup associations in complex disease is difficult under most study conditions where power is limited with sample sizes <10,000 (Samuels et al., 2006).

It is intriguing to note that a couple of the nominal hits in phase 1 tag subgroups of haplogroup L. These signals are generated by the small and diverse set of self-reported African-ancestry samples within our dataset. The inherent diversity, which this panel of markers is designed to capture, among individuals of African ancestry combined with probable admixture and consequent misclassification of their mitochondrial ancestry increases the difficulty of appropriately ascertaining matched cases and controls based on self-report. Importantly, our study underlines the need for larger, more powerful studies of patients with African ancestry. These will allow us to make a definitive statement on the involvement of mtDNA in ASD susceptibility in a more inclusive manner.

Beyond our examination of single marker tests for association, we have explored a number of additional possible mechanisms by which mitochondrial variation may play a role in ASD. Collectively, our investigations fail to provide any convincing evidence for a major contribution of mtDNA variation or heteroplasmy to ASD.

## Software

MSDAT source code and documentation is available for download at the Hussman Institute for Human Genetics website at <http://hihg.med.miami.edu/software-download>.



## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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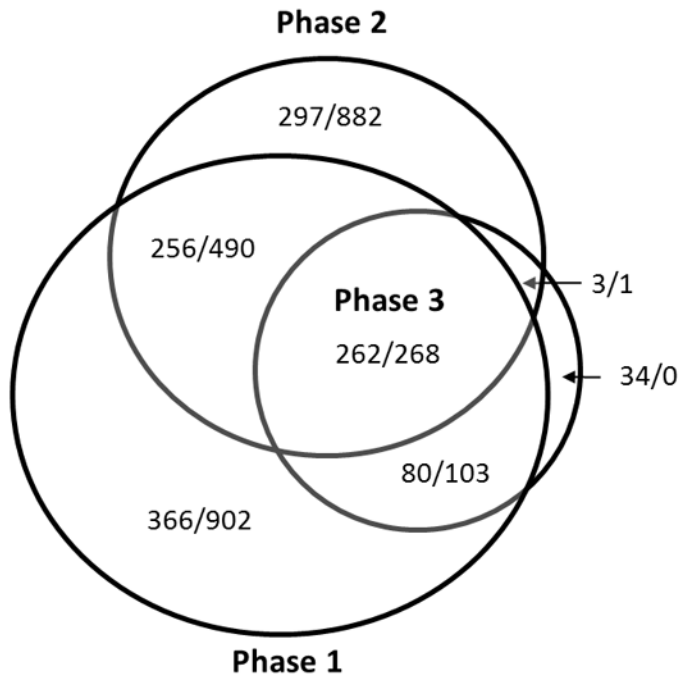
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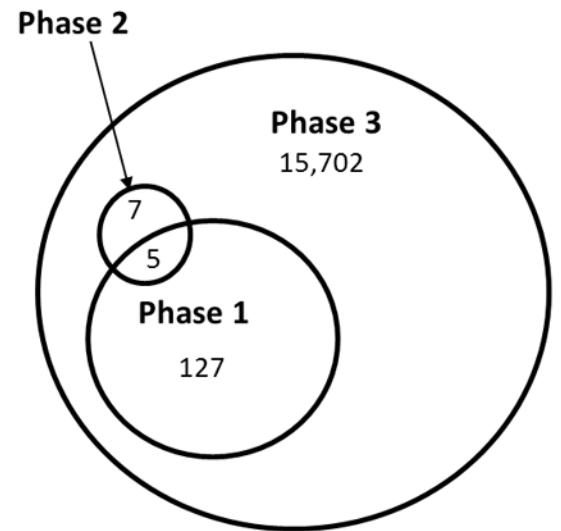
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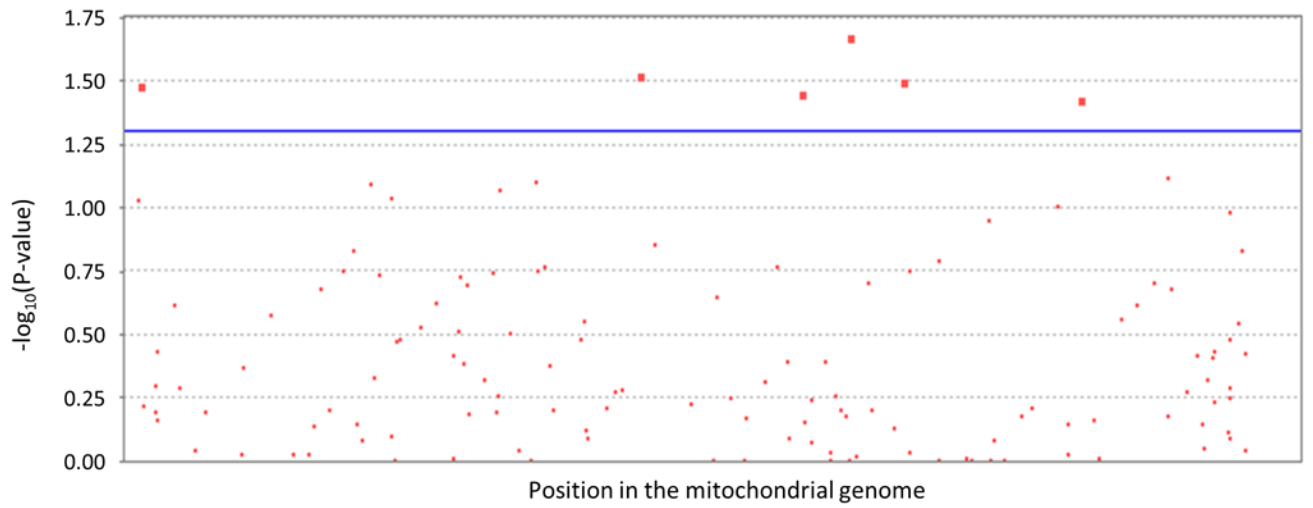
### A Cases/Controls



### B Markers



**Figure 1.**  
A graphical representation of the three phases of this study for a) samples and b) markers.



**Figure 2.**

A manhattan plot of the phase 1 logistic regression results.

The line represents the uncorrected significance threshold ( $P = 0.05$ )

Table 1

## Race/Ethnicity Dataset Descriptions

Phase (Platform)		* Overall Unique		Phase 1 (Illumina)		Phase 2 (Sequenom)		Phase 3 (Affymetrix)	
Race	Ethnicity	Cases	Controls	Cases	Controls	Cases	Controls	Cases	Controls
White	Non-Hispanic	885	1524	596	1212	551	1391	379	372
White	Hispanic	78	163	63	163	29	22	0	0
White	Unknown	145	708	167	182	165	172	0	0
Black	Non-Hispanic	41	139	40	143	6	3	0	0
Black	Unknown	32	21	32	21	2	1	0	0
**Other/Unknown		117	91	66	42	65	52	0	0
TOTAL		1298	2646	964	1763	818	1641	379	372

\* As different phases of this project contain overlapping samples, this represents the non-overlapping unique sample set.

\*\* Contains individuals of self-reported race and ethnicity of American Indian, Asian, and other/unknown.

**Table 2**

Nominally significant ( $p < 0.05$ ) logistic regression results of mtDNA SNPs analyzed in phase 1

SNP	Basepair Position	Minor Allele	MAF in controls	MAF in mtDB	P-value	Amino Acid Change	Region
MitoG228A	228	A	0.052	0.032	0.03	na	D-loop
MitoG7522A	7521	A	0.087	0.060	0.03	na	t-RNA (Asp)
MitoT9900C	9899	C	0.020	0.010	0.04	syn	COX 3
MitoG10590A	10589	A	0.013	0.017	0.02	syn	ND4L
MitoG11378A	11377	A	0.011	0.010	0.03	syn	ND4
MitoT13966C	13965	C	0.016	0.006	0.04	syn	ND5

**Table 3**

Phase 2 Haplogroup Results

Haplogroup	Counts		Frequency		P-value	OR	L95	U95
	Cases	Controls	Cases	Controls				
H	358	762	0.44	0.46	0.21	0.90	0.76	1.06
I	21	36	0.03	0.02	0.56	1.18	0.68	2.03
J	83	146	0.10	0.09	0.32	1.16	0.87	1.54
K	64	134	0.08	0.08	0.77	0.96	0.70	1.30
T	82	173	0.10	0.11	0.69	0.95	0.72	1.25
U	147	272	0.18	0.17	0.39	1.10	0.88	1.38
V	32	56	0.04	0.03	0.53	1.15	0.74	1.79
W	18	36	0.02	0.02	0.99	1.00	0.57	1.78
X	13	26	0.02	0.02	0.99	1.00	0.51	1.96
Total	818	1641	1.00	1.00	-	-	-	-

OR, odds ratio; L95/U95, lower and upper bounds of the 95% confidence interval for the OR



**Table 4**

Logistic regression results for haplogroup defining mtDNA SNPs analysed in Phase 2

SNP	Basepair position	Minor Allele	**MAF	P-value	OR	L95	U95
*MitoA1719G	1719	G	0.04	0.44	1.18	0.78	1.79
MitoC4216T	4216	T	0.17	0.58	1.07	0.85	1.34
MitoA4580G	4580	G	0.04	0.79	0.94	0.58	1.52
*MitoG4917A	4917	A	0.09	0.34	0.86	0.63	1.17
MitoC7028T	7028	T	0.39	0.30	0.83	0.58	1.18
MitoA8251G	8251	G	0.04	0.73	1.08	0.71	1.63
MitoA9055G	9055	G	0.09	0.93	0.99	0.73	1.33
*MitoG10398A	10398	A	0.26	0.31	1.12	0.90	1.41
*MitoG12308A	12308	A	0.22	0.99	1.00	0.78	1.27
MitoA13368G	13368	G	0.1	0.66	0.94	0.69	1.26
MitoA13708G	13708	G	0.09	0.09	1.28	0.97	1.71
*MitoA16391G	16391	G	0.02	0.68	1.13	0.63	2.03

\* These 5 SNPs were also genotyped in Phase 1

\*\* Minor allele frequency in controls

OR, odds ratio; L95/U95, lower and upper bounds of the 95% confidence interval for the OR

Table 5

Coding changes per gene in Phase 3 analysis

Mito Gene	*Synonymous Changes		**Non-synonymous Changes		P-Value
	Controls	Cases	Controls	Cases	
ATPase8	12	18	24	19	0.41
ATPase6	80	66	93	101	0.66
ND1	100	86	124	117	0.55
ND2	136	152	92	99	0.71
ND3	36	30	100	89	0.35
ND4	525	544	46	32	0.10
ND7	56	63	6	3	0.46
ND5	357	364	131	129	0.78
ND6	133	136	23	20	0.60
CO1	302	306	37	26	0.14
CO2	66	71	10	9	0.79
CO3	117	103	55	70	0.22
Cytb	162	151	448	452	0.88

\* Coding changes per gene that DO NOT result in an amino acid change from rCRS

\*\* Coding changes per gene that DO result in an amino acid change from rCRS

**Table 6**

mtDNA variants shown to be associated with other diseases

*SNP	Basepair position	Minor Allele	**P-value
MitoT1005C	1005	C	0.50
MitoG1438A	1438	A	0.14
MitoA1555G	1555	G	0.12
MitoG3316A	3316	A	0.50
MitoA3796G	3796	G	0.25
MitoC4025T	4025	T	0.50
MitoC4171A	4171	A	0.50
MitoA4295G	4295	G	0.38
MitoC4640A	4640	A	0.49
MitoT5814C	5814	C	0.50
MitoC6489A	6489	A	0.25
MitoG7444A	7444	A	0.50
MitoA8348G	8348	G	0.50
MitoG9804A	9804	A	0.10
MitoT9957C	9957	C	0.50
MitoT10237C	10237	C	0.49
MitoA11084G	11084	G	0.50
MitoT11253C	11253	C	0.16
MitoG11696A	11696	A	0.51
MitoA12026G	12026	G	0.50
MitoT12297C	12297	C	0.50
MitoT12811C	12811	C	0.25
MitoA13637G	13637	G	<b>0.02</b>
MitoT14325C	14325	C	0.38
MitoG14831A	14831	A	0.25

\* Novel Phase 3 interrogated variation was compared against the “mtDNA Mutations with Reports of Disease-Association” from [www.mitomap.org](http://www.mitomap.org)

\*\* Fisher's exact test with 1 degree of freedom

**Table 7**

The distribution of mtDNA SNPs between ASD Cases and Controls in the Phase 3 resequencing dataset

	*Total sites possible		Cases		Controls		$\chi^2$ P-value
	Total observed	Per individual (N = 379)	**Total mutational burden	Per individual (N = 372)	Total observed	**Total mutational burden	
Sequence variants <sup>1</sup>	16,010	53.0	0.0033	49.6	18435	0.0031	NS
"n" SNPs	16,010	34.7	0.0022	31.1	11566	0.0019	NS
Homoplasmic SNPs <sup>2</sup>	16,010	18.3	0.0011	18.5	6869	0.0012	NS
Potential heteroplasmic SNPs <sup>3</sup>	16,010	37.5	0.0023	33.9	12610	0.0021	NS
Heteroplasmic SNPs <sup>4</sup>	16,010	2.8	0.0002	2.8	1044	0.0002	> 0.05

\*\* rCRS probes to 16,544 positions, 534 of which failed the 95% call rate threshold using the diploid algorithm

\* Total mutational burden = total variants observed/(total sites possible x N of cases or controls)

<sup>1</sup> Homoplasmic SNPs and n calls

<sup>2</sup> IUPAC codes "a", "c", "g", "t"

<sup>3</sup> IUPAC codes "r", "y", "k", "m", "s", "w", "n"

<sup>4</sup> IUPAC codes "r", "y", "k", "m", "s", "w"