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Project Title:	Astronaut Vision Issues and One Carbon M Potential Analog Population	letabolism: Expanded Polymorphism Evaluat	tion and Evaluation in a
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Program/Discipline Element/Subdiscipline:	HUMAN RESEARCHBiomedical counter	ermeasures	
Joint Agency Name:		TechPort:	No
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Human Research Program Risks:	(1) SANS:Risk of Spaceflight Associated N	Jeuro-ocular Syndrome (SANS)	
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Zip Code:	77058-3607	Congressional District:	36
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Flight Program:			
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Task Description:	 We have documented a genetic predisposition for some astronauts to develop ophthalmologic findings as described in 2011 by NASA's Space Medicine Division (1) known as VIIP (vision impairment/intarcanial pressure). From a limited study of 5 single-nucleotide polymorphisms (SNPs), we found one SNP associated with a greater risk of ophthalmic findings (e.g., choroidal folds, cotton wool spots), and another SNP that was protective against optic dise edema. In light of these findings, we propose to evaluate a wider range of SNPs from the same metabolic pathway, to clarify and verify the relationship of genetics in some astronauts that may predispose to ophthalmic anomalies. Furthermore, we have identified a clinical population patients with polycystic ovary syndrome (PCOS) with several characteristics similar to those described in astronauts. The clinical findings shared by PCOS patients and astronauts with ophthalmic anomalies include higher homocysteine, increased retinal nerv fiber layer, increased incidence of white matter hyperintensities on MRI, increased androgen concentrations (or responses), and altered carbohydrate metabolism. PCOS patients have not been evaluated in detail regarding the occurrence of other anomalies in eycloplegic refraction, and optic nerve sheath diameter. While researchers have evaluated one-carbon polymorphisms, none have looked at the complete set of SNPs we propose here. Thus, this study has two major goals 1. To extend the one-carbon natway SNP assessment as related to astronaut ophthalmologic findings. 2. To evaluate patients with PCOS and/or Idiopathic intracranial hypertension (IIH) to assess one-carbon biochemistry and genetics and their possible correlation with ophthalmologic findings. While these studies alone will not identify the mechanism(s) of VIIP, we aim to clarify the genetic relationship to ophthalmic findings, and to document the utility of PCOS as a clinical population that could be used for studies that may ultimately allow for the definition of
	2. Compare the same one-carbon metabolism genetics and biochemistry and ophthalmologic data from patients in one of four treatment groups:
	i. women diagnosed with PCOS without IIH
	ii. women diagnosed with PCOS and IIH
	iii. women diagnosed with IIH without PCOS
	iv. controls (neither PCOS nor IIH)
	Subjects will be matched by age and body mass index (BMI). Statistical analyses will be used to evaluate the independent and shared contributions of age, body mass (BMI), and genetics on biochemical, and ophthalmologic outcomes, with False Discovery Rate adjustments to account for multiple comparisons.
	3. A secondary aim is to combine the patient and control data from this study with ISS crewmember data in order to help inform us on whether or not these two cohorts (VIIP, PCOS) share similar associations among one-carbon metabolism genetics and biochemistry and ophthalmologic data identified in our earlier analyses.
Rationale for HRP Directed Research:	This research is directed because it contains highly constrained research. This study has two major goals: 1. To utilize existing samples where possible to extend the scope of the initial One Carbon study. This was initially submitted and reviewed in the NNJ14ZSA001N-OMNIBUS NRA. HRP Management has now asked we submit this as directed research. 2. To add testing to an ongoing clinical trial at the Mayo Clinic. Timing is critical given that study is ongoing. The primary study is a clinical trial of pharmaceutical treatment for PCOS. We propose to extend this study by collecting a blood sample for one carbon biochemical and genetic testing, along with ophthalmologic exams, with the aim of documenting the utility of this population as an analog group for future VIIP research.
Research Impact/Earth Benefits:	While much research is in progress to understand vision issues in astronauts, a key question remains as to why only some individuals are affected. Our preliminary data suggest that some individuals may have a genetic predisposition for vision issues, related to one-carbon metabolism. Our initial study was intentionally constrained given our concerns about it being the first study involving individual genetic testing at NASA. In light of the crewmember response to that study (>97% participation) and the initial findings from that effort, we now propose to evaluate a wider range of one-carbon metabolism SNPs, to help clarify and verify that one-carbon metabolism is indeed the source of this effect, and to identify possible associations with ethnicity. The results of this study could be profound, and may have significant implications for the direction of NASA vision countermeasure research, for operational decisions regarding treatment of affected astronauts, and for informing the general medical and scientific communities, where research is ongoing to understand the role of one-carbon metabolism genetics in other cerebrovascular issues.

Task Progress:	For the flight study: we have obtained IRB (Institutional Review Board) approval, and Human Research Program (HRP) select for flight approval. We are currently waiting for International Space Station Medical Projects (ISSMP) element to determine if we can consent crewmembers by email. For the PCOS study: we obtained IRB approvals at Johnson Space Center (JSC) and Mayo Clinic. Subject recruitment, screening, and data collection have been initiated.
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