The Transcriptional Response of Diverse Saccharomyces cerevisiae Strains to Simulated Microgravity LILY S. NEFF^{1,2}, SAMANTHA T. FLEURY^{3,4}, JONATHAN M. GALAZKA⁵

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Stresses of Spaceflight



Microgravity



Space exploration missions place stresses on the space crew and their supporting microbial commensals

Reveal a conserved response to the stress of microgravity, measure physiological response



Why yeast? Why S. cerevisiae?

- Powerful microbial model
- Easy to grow and allows for transcriptomes to be recorded cheaply
- Part of human microbiota
- S. cerevisiae

Yeast

- 🕨 Human colonizer
- > Opportunistic pathogen
- Diverse set of strains are readily available



Credit: Biotium

Studying Simulated Microgravity

- Environmental stress of SMG causes an increased growth of Candida albicans in filamentous forms¹
 - Alteration in two genes associated with this hyphal transition
 - Evidence of enhanced pathogenicity, fungal pathogen becomes more virulent
- ~300 million year divergence from S. cerevisiae²

¹(Altenburg et al. 2008 Geno. Prot. Bioinfo. Increased Filamentous Growth of Candida albicans in Simulated Microgravity)

²(Hedges SB et al., Tree of life reveals clock-like speciation and diversification. Mol Biol Evol. 2015 Apr32(4):835-45.)

HARV Control 25 Generations

HARV SMG 25 Generations



Studying Simulated Microgravity cont.

- Studies conducted show that cells perceive and respond to variations in mechanical forces, i.e. gravity¹
- S. cerevisiae, under SMG, demonstrates random budding than typical bipolar budding pattern¹





Hierarchical clustering based on 600 growth rate variables

- clustering represents patterns
- shows diversity in phenotype and physiology

Credit: Warringer J, Zoʻrgoʻ E, Cubillos FA, Zia A, Gjuvsland A, et al. (2011) Trait Variation in Yeast Is Defined by Population History. PLoS Genet Neff 6



Yeast are Diverse!



Project Work Flow

Central Objective: Reveal a conserved response across all strains or unique to lab strain, \$288c

► How to accomplish this:



Screening Procedure: YPD Plate

- Create YPD plates (1% yeast extract, 2% peptone, 2% glucose) and YPD liquid culture
- Inoculate YPD plates with strains from cryogenic stock
- Observe for unusual growth (different morphologies):







YJM996 (normal)



Screening Procedure: Liquid Culture

- Inoculate 5mL YPD broth from overnight "normal" cultures on YPD plates, incubate overnight
- ▶ Dilution 10µL:100µL to test OD₆₀₀ using NANODROP 2000 Spectrometer

> 24 hour incubation for microscopy check

▶ 48 hour incubation for HARV Vessels





Neff 11

Screening Procedure: Microscopy

Top Row (left to right):

YJM1439 (West African, Clinical)

YJM1388 (Sake, Non-clinical)



Bottom Row (left to right):

YJM1248 (West African, Nonclinical)

YJM627 (West African, Nonclinical)

Results to Date

Color Code Key:

	Gray : cryogenic stock DNE	Strain	Clade	Plate	Flask	Microscope	Cleared for HARV
		YJM1078	European, Clinical				
	Red : Did not have normal phenotype, cannot use	YJM1450	European, Clinical				
		YJM1526	European, Clinical				
		YJM244	European, Clinical				
	Light Green : Normal Phenotype so far; TBD	YJM248	European, Clinical				
		YJM453	European, Clinical				
		YJM969	European, Clinical				
Denote Indenote Methods Performance Verify	Dark Green : All normal, including Microscopy	YJM972	European, Clinical				
		YJM978	European, Clinical				
	Phenotype; can	YJM984	European, Clinical				
	vessel	L	•	•	•	·	





94 S.

and

Cleared for HARV Use





▶ isolated from clinical and environment[•] al settings ▶ multiple

strains:

locations around the world to encompass evolutionary divergence

West African

High Aspect Ratio Vessel (HARV)

- Simulates microgravity conditions by rotating on vertical plane
- "functional weightlessness"*
 - randomizes the gravitational effect
 - minimizes turbulence (fluid undergoes irregular fluctuations) over surface of cell
- Remain suspended in liquid culture

*(Altenburg et al. 2008 Geno. Prot. Bioinfo. Increased Filamentous Growth of Candida albicans in Simulated Microgravity)





NANO Chip Bioanalyzer

- Purity (degree of contamination) and quality (intactness/integrity) of RNA are essential for examining gene expression
- Degraded samples lead to misrepresentative data and inconsistency in reproducibility



RNA Nano Chip

LOT#: UM098K20 EXP:09.DEC:2017





- Allow for generation of transcriptome information cheaply
- Allows for the investigation of known transcripts and new ones (important for the comparisons)
 - Analyze physiology and phenotype
- Allows for the identification of conservation with gene expression profiles



Future Plans of Progression

- Complete simulated microgravity runs for the 32 strains (along with control experiments)
- Complete RNA Extraction and Illumina sequencing of samples using the KAPA mRNA HyperPrep Kit
- Send samples to be sequenced at UCSF and analyze data

Significance

- Systemic understanding of how microbes respond to simulated space flight environment
- Serve as a platform for future flight experiments



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