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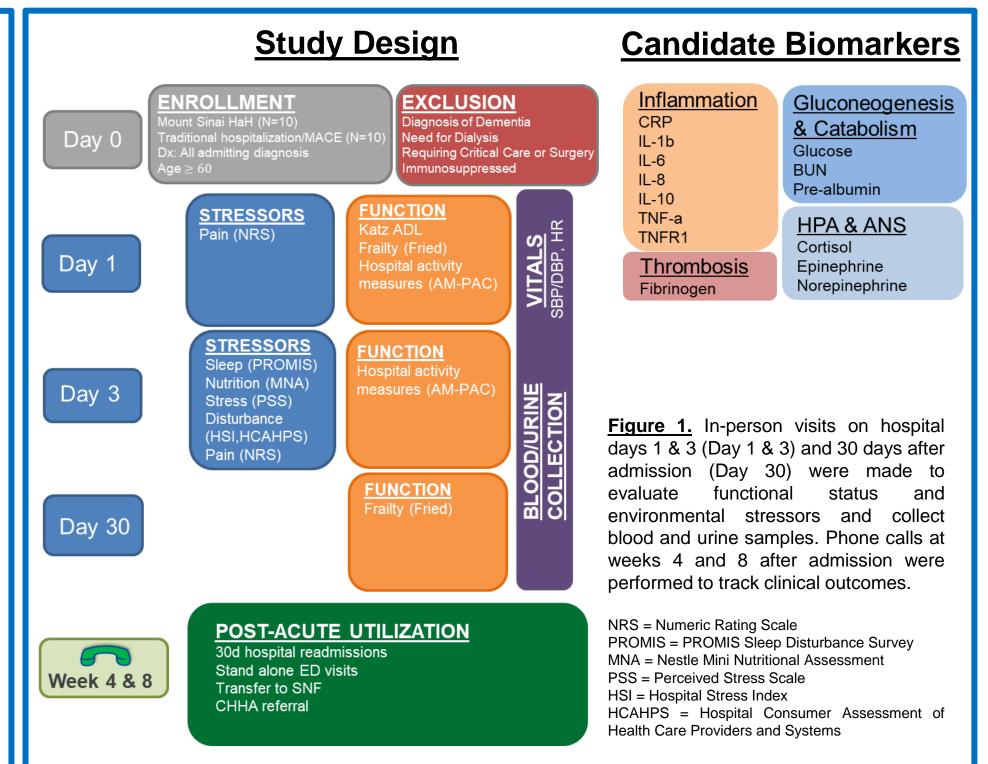
Medicine at

# Environmental Stressors and Biological Markers of Post-Hospital Syndrome: A Feasibility Study

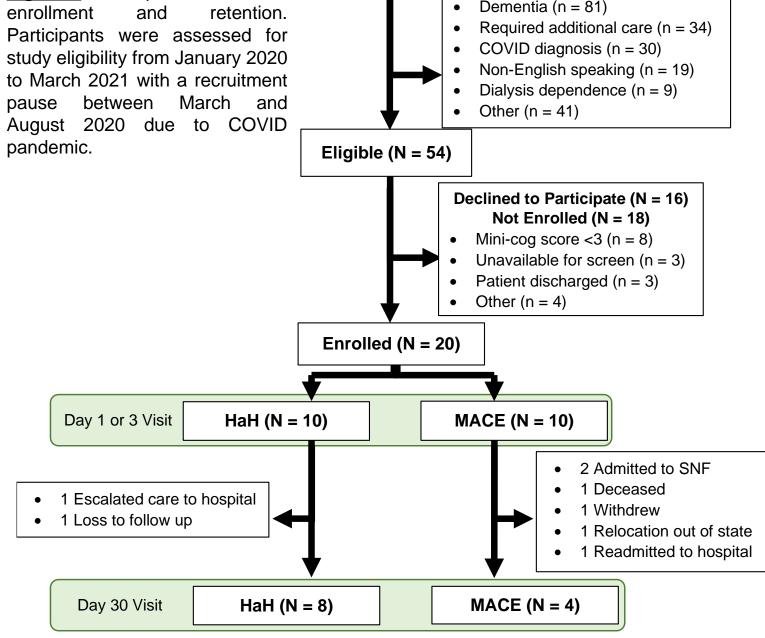
I. Chun<sup>1</sup>, C. Park<sup>2</sup>, G. Schiller<sup>3</sup>, D. Goldwater<sup>4</sup>, D. Levine<sup>5</sup>, N. Latham<sup>5</sup>, B. Leff<sup>6</sup>, A. Siu<sup>3</sup>, F. Ko<sup>3</sup>
 1. University of Hawai'i, John A. Burns School of Medicine, 2. Stanford School of Medicine, 3. Icahn School of Medicine at Mount Sinai,
 4. David Geffen School of Medicine at UCLA, 5. Brigham and Women's Hospital, 6. Johns Hopkins School of Medicine

### **Background**

- Post-Hospital Syndrome (PHS) in older adults is characterized by a transient period of heightened vulnerability to adverse outcomes (ED revisits, hospital readmissions, mortality) following hospitalization.
- We hypothesize that: (1) environmental stressors associated with hospitalization (sleep disruption, poor nutrition, mobility restrictions) are linked to biological mechanisms underlying PHS-associated vulnerability; and (2) dysfunction of stressresponse systems such as the hypothalamicpituitary-adrenal (HPA) axis, inflammatory, and metabolic systems are implicated in PHSassociated vulnerability.
- Hospital at home (HaH) provides comparable levels of care to traditional inpatient hospitalization but is associated with reduced lengths of stay, ED revisits, hospital readmissions, and admissions to skilled nursing facilities suggesting that treatment in the home environment may reduce PHS.
- We implemented a feasibility pilot comparing HaH vs traditional geriatric inpatient hospitalization (Mobile Acute Care for the Elderly/MACE), in order to evaluate differences in hospital-induced stress exposures and biological markers of stress.



Results Sleep GROUP								
Assessed for Eligi					· · · · · · · · · · · · · · · · · · ·	9) MACE $(n = 8)$	р	
Jan 2020 – Mar 2020 Aug 2020 – Mar 2021				No sleep distress	7	4		
Aug 2020 – 1			Sleep Disturbance Level	Mild sleep distress	2	0	0.034	
Figure 2. Participant recruitment,	Excluded (N = 214)			Moderate sleep distress	0	4		



#### **Feasibility**

Determined by adherence of  $\geq$ 80% of enrolled participants who were available to be approached (e.g. not deceased, admitted to skilled nursing facility, readmitted to hospital) in completing study procedures.

- 100% (19/19) completed a Day 1 or 3 visit
- 92% (12/13) completed a Day 30 visit
- 84% (16/19) completed a Day 1 or 3 blood collection

### **Biological Markers**

Day 3 Serum Prealbumin Correlation

	Pearson Correlation	р
Changes in Meals	0.678	0.031
AMPAC Day 3	0.644	0.045

**Table 4.** Serum prealbumin level positively correlated with meal intake and mobility and transfer score on hospital day 3.

## **Conclusion**

In this pilot study, HaH patients reported less sleep disturbance, fewer changes to nutritional intake, and retained mobility and transfer ability compared to patients that experienced traditional

Disturbed by procedures	Unbothered by procedures	8	1	0.003
Distuibed by procedures	Bothered by procedures	1	7	0.005

**Table 1.** MACE patients were more likely to report sleep distress and being woken up in the middle of the night for procedures (e.g. check vitals, administer medications) between hospital days 1 and 3.

### <u>Nutrition</u>

		GF		
Changes in amount eaten		HaH(n = 9)	MACE $(n = 8)$	р
	Less	2	4	
Meals and meal size	Same	6	1	0.074
	More	1	3	
	Less	0	3	
Dairy	Same	8	2	0.023
	More	1	3	
	Less	1	5	
Beans or eggs	Same	8	2	0.027
	More	0	1	
	Less	3	3	
Meat, fish, or poultry	Same	6	2	0.149
	More	0	2	
	Less	1	4	
Fruits or vegetables	Same	8	2	0.025
	More	0	2	

**Table 2.** MACE patients were more likely to report nutritional intake changes in meals both in quantity and content between hospital days 1 and 3 compared to prehospitalization baseline.

#### **Mobility and Transfer Ability**

		Ν	Mean (Std. dev)	р	
AMPAC Day 1	HaH	10	17.40 (6.275)	0.920	
	MACE	9	17.11 (6.009)		
AMPAC Day 3	HaH	9	17.78 (6.888)	0.357	
	MACE	8	14.50 (7.329)		
Day 1 to Day 3 Change	HaH	9	0.55 (1.42)	0.040	
	MACE	7	-2.57 (2.82)	0.012	

**Table 3.** MACE patients reported decreased mobility and transfer (Activity Measure for Post Acute Care scores) between hospital days 1 and 3 compared to HaH patients.

### **Acknowledgments**

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hospitalization.
Serum prealbumin level on hospital day 3 may be a correlate of nutritional intake and biomarker of ambulatory and transfer ability.
This pilot study demonstrates the feasibility of a larger, adequately powered trial to definitively quantify hospital-induced stress and identify their effects on multisystem dysfunctions and ikho impacts on PHS in vulnerable older adults.

Contact InformationIan Chun, BSFred Ko, MDJohn A. Burns School of Medicine<br/>at the University of Hawai'iGeriatrics & Palliative Medicine<br/>Icahn School of Medicine at<br/>Mount Sinai<br/>Fred.ko@mssm.edu