

Status of Department of Defense Funded Suicide Research

Peter M. Gutierrez, Ph.D. (moderator), Diana J. Fitek, Ph.D., Thomas Joiner, Ph.D., Dave Jobes, Ph.D., Marjan Holloway, Ph.D., and M. David Rudd, Ph.D.

Brief Cognitive Behavioral Therapy (BCBT)

M. David Rudd, Ph.D., ABPP

ASPIRE-1 Team

(Army Suicide Prevention & Intervention Research at Evans)

University of Utah

M. David Rudd, PhD, ABPP (PI)
Craig J. Bryan, PsyD, ABPP (PM)
Kim Arne, LMSW (Therapist)
Sharon Stone, LCSW (Therapist)
Sean Williams, LMSW (Evaluator)

UTHSCSA

Alan L. Peterson, PhD, ABPP (Co-I)
Jim Mintz, PhD (Biostat)
Stacey Young-McCaughan, RN, PhD (Co-I)
Deanne Hargita (Regulatory)

Ft. Carson

Evelyn Wertenberger, PhD, LCSW (Site PI)
MAJ Jill Breitbach, PsyD (Collaborator)
Travis Bruce, MD (Collaborator)
LTC Erin Wilkinson, PsyD (Collaborator)
Kenneth Delano, PhD (Collaborator)

Army Warrior Resiliency Program

COL Bruce Crow, PsyD (Consultant)
MAJ Monty Baker, PhD (Consultant)

Brief Cognitive Behavioral Therapy for Military Populations

W81XWH-08-MOMRP-SPRC, Suicide Prevention and Counseling Research
W81XWH-09-0569



PI: M. David Rudd

Org: The University of Utah

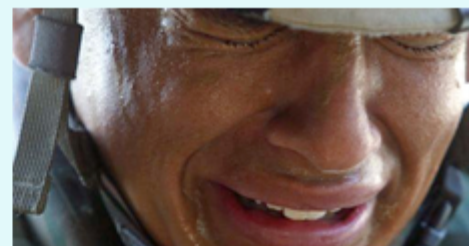
Award Amount: \$1,987,035.00

Study/Product Aim(s)

1. To evaluate the effectiveness of brief cognitive-behavioral therapy for suicidality (BCBT-S), including suicidal ideation and attempts, among active duty military personnel.
2. To engage in prospective investigation of suicide risk factors and warning signs, exploring their ability to predict subsequent suicidal behavior.
3. To explore the effectiveness of BCBT-S for increasing appropriate utilization of and compliance with medical, mental health, and substance abuse treatment.
4. To develop a risk management software program for the initial risk assessment, ongoing monitoring, and clinical management of high-risk suicidal patients.

Approach

Randomized controlled trial randomizing 150 Soldiers to either BCBT-S or treatment as usual (TAU), with follow-up for 24 mos.



Reducing suicidal behavior among military personnel through targeted treatment.

Number of enrolled/randomized participants has increased to 90 out of 150, the first 12-month follow-up assessments have been initiated, and several data analyses have been initiated and submitted for presentation and/or publication.

Timeline and Cost

Activities	FY	10	11	12
Staff hiring and training		█		
Randomized clinical trial			█	█
Data entry/deaning			█	█
Data analysis / dissemination				█
Estimated Budget (\$K)		\$697	\$626	\$644

Updated: 6 March 2012

Goals/Milestones

CY10 Goal – Staff hiring and training

- Obtain IRB approvals
- Interview, hire, train, and supervise therapists and evaluator

CY11 Goals – Conduct randomized clinical trial

- Begin recruiting participants and conducting intake evaluations
- Begin randomizing participants and administering BCBT-S
- Begin follow-up assessments

CY12 Goal – Complete randomized clinical trial & disseminate results

- Complete recruiting participants and intake evaluations
- Complete randomizing participants and administering BCBT-S
- Complete follow-up assessments

- Analyze data, publish papers, present results

Comments/Challenges/Issues/Concerns

- None – study is on schedule and on budget

Budget Expenditure to date

Projected Expenditure: \$1,845,000 Actual Expenditure: \$1,208,523

Phase I:

Crisis management, distress tolerance

Phase II:

Cognitive restructuring of suicidal belief system,
problem solving, cognitive flexibility

Phase III:

Relapse prevention

How BCBT differs from TAU

TAU ($n = 75$)

- Suicide as symptom of psychiatric dx
- Focus on psych dx
- Emphasizes external sources of self-mgt, including hospitalization
- Clinician responsibility for preventing suicide

BCBT ($n = 75$)

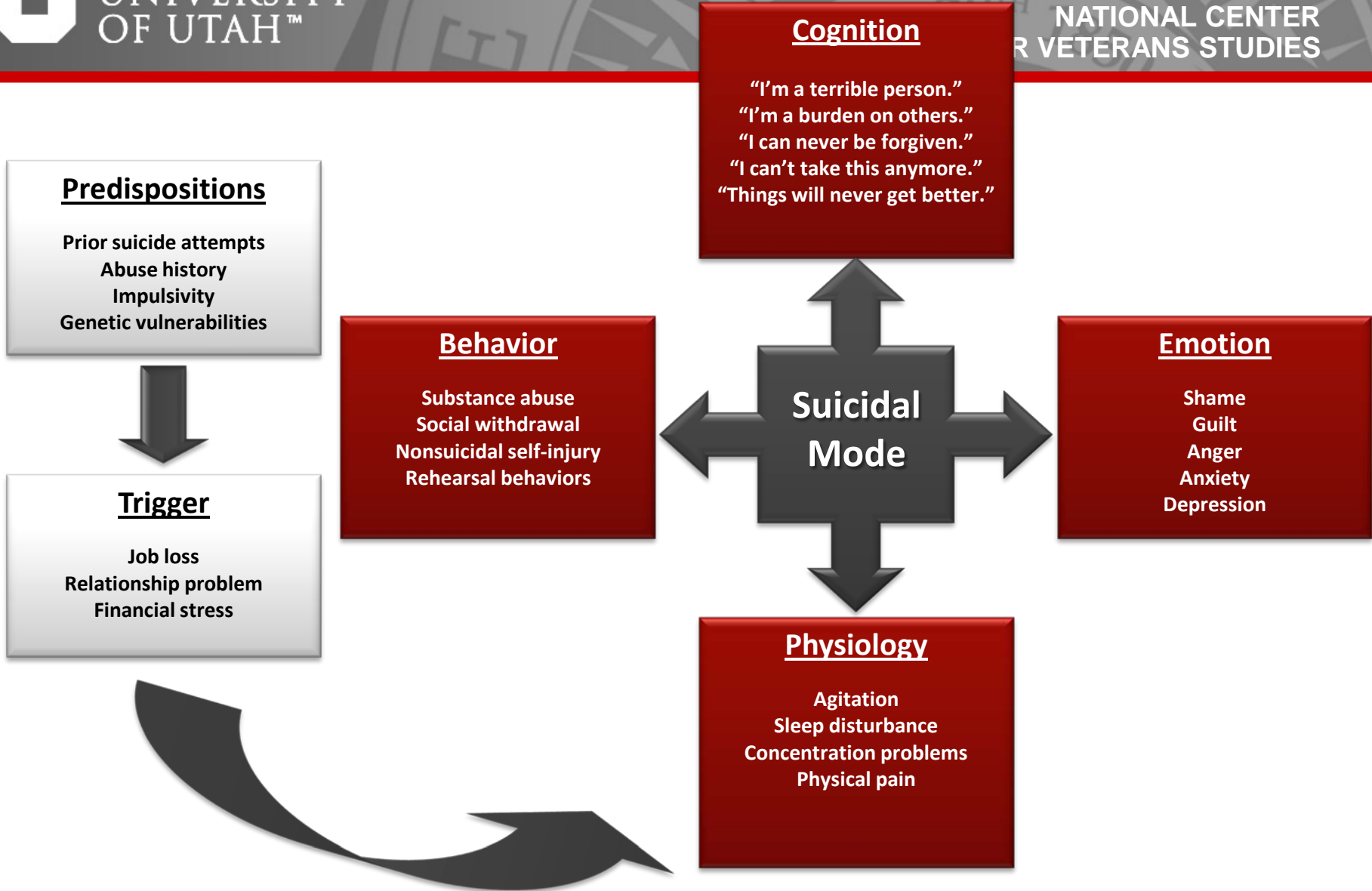
- Suicide as problem distinct from psych dx
- Focus on suicide risk
- Emphasizes internal sources of self-mgt to minimize hospitalization
- Shared patient-clinician responsibility for preventing suicide

Competency-based progress

- Progress through treatment is determined based on patient skill mastery
- Patient must demonstrate skill mastery for each phase before progressing to next phase
- If patient demonstrates insufficient skills mastery at later phase, clinician returns to earlier phase
- Final competency check is relapse prevention task

Primary treatment tasks

1. Describe treatment
2. Conduct assessment of index suicidal episode
3. Educate patient about suicidal mode
4. Develop crisis response plan
 - Means restriction counseling
5. Develop treatment plan & obtain commitment
 - Commitment to treatment agreement
6. Emotion regulation skills training



Emotion regulation strategies

- Relaxation training
- Mindfulness training
- Reasons for living list
- Survival kit
 - Including Reasons for Living
- Sleep hygiene / stimulus control
- Recognize critical role of shame/guilt/grief

Completion Coin



Early observations

- Service members take numerous medications
- Providing patients with treatment log (or “smart book”) is a highly effective method for obtaining buy-in, skills training, and relapse prevention
- Framing treatment as occupational skills training
- Phase I must target emotion regulation
- Guilt/shame common themes & targets of Phase II
- BCBT appears to retain patients at a higher rate
- Combat exposure /trauma are distal contributors

Operation Worth Living Project: A Randomized Clinical Trial of CAMS vs. E-CAU at Ft. Stewart GA

David A. Jobes, Ph.D., ABPP

Principal Investigator

Professor of Psychology

The Catholic University of America

Washington, DC

American Association of Suicidology

Annual Conference

Perspectives from DOD Task Force



DoD Task Force on the Prevention of Suicide By Members of the Armed Forces

10 November 2009

Bethesda, MD



Evidence-Based Treatments for Suicidality



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

National Institutes of Health
National Institute of Mental Health
6001 Executive Boulevard Rm 8235
Bethesda, Maryland 20892

Opportunities to Improve Interventions to Reduce Suicidality: Civilian “Best Practices” for Army Consideration

Michael Schoenbaum, PhD
schoenbaum@mail.nih.gov

Robert Heinsse, PhD
rheinsse@mail.nih.gov

Jane Pearson, PhD
jpearson@mail.nih.gov

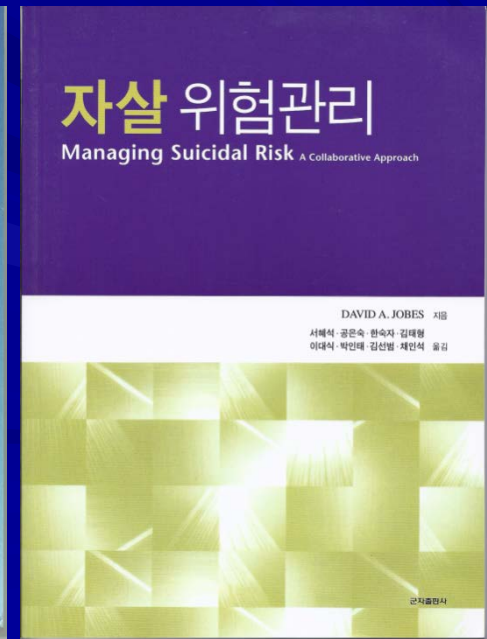
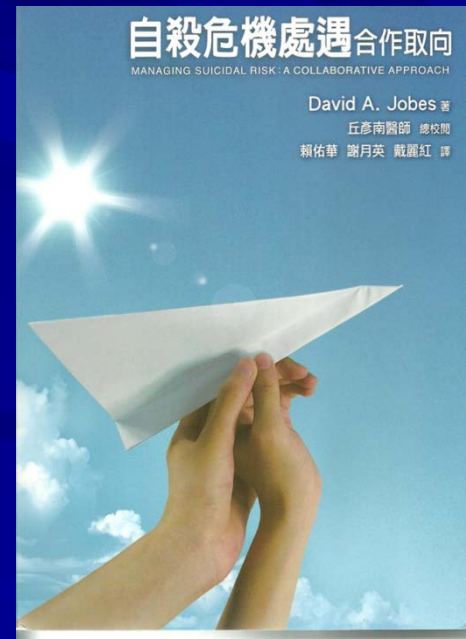
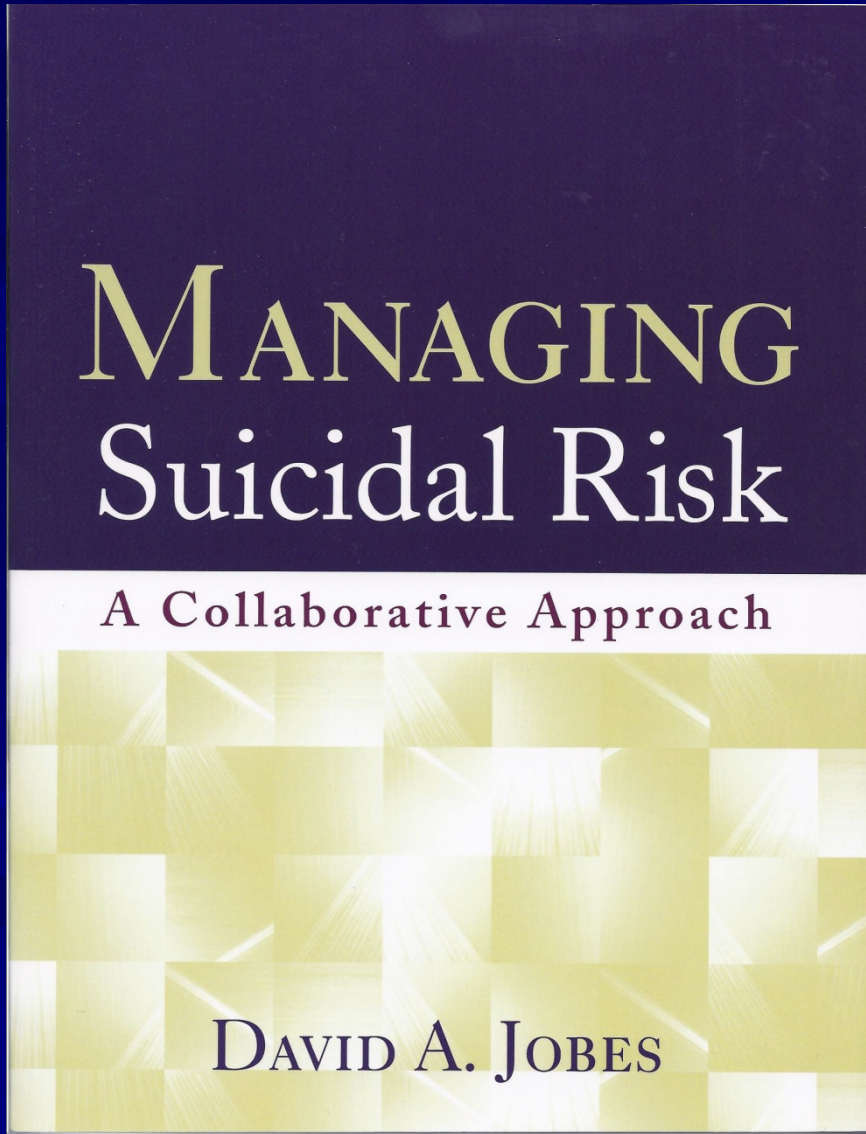
Author affiliations: All authors are affiliated with the Division of Services and Intervention Research, National Institute of Mental Health, 6001 Executive Blvd, Bethesda, MD 20892

The views expressed in this document do not necessarily represent the views of the National Institute of Mental Health, the National Institutes of Health, the Department of Health and Human Services, or the United States government.

* This document was developed in support of the US Army's ongoing efforts to reduce suicides and suicidality among Army Soldiers, and submitted to General Peter Chiarelli, Vice Chief of Staff of the Army, in December 2009. An earlier version of this document was submitted to Mr. Robert Andrews, Special Assistant to the Secretary of the Army, and General Chiarelli, in May 2009. The current version has been updated and somewhat expanded, in particular by adding a section on quality assurance and performance metrics (Section III).

- With n=49 studies (in the world literature), there are remarkably few evidence-based treatments and interventions for suicidal risk
- We mostly know what does not work (e.g., medication only)
- What does work:
 - Dialectic Behavior Therapy (DBT)
 - Cognitive Therapy
 - Brief interventions with non-demand follow-up

The Collaborative Assessment and Management of Suicidality (CAMS)



TEAM STEWART

Fort Stewart / Hunter AAF / Kelley Hill



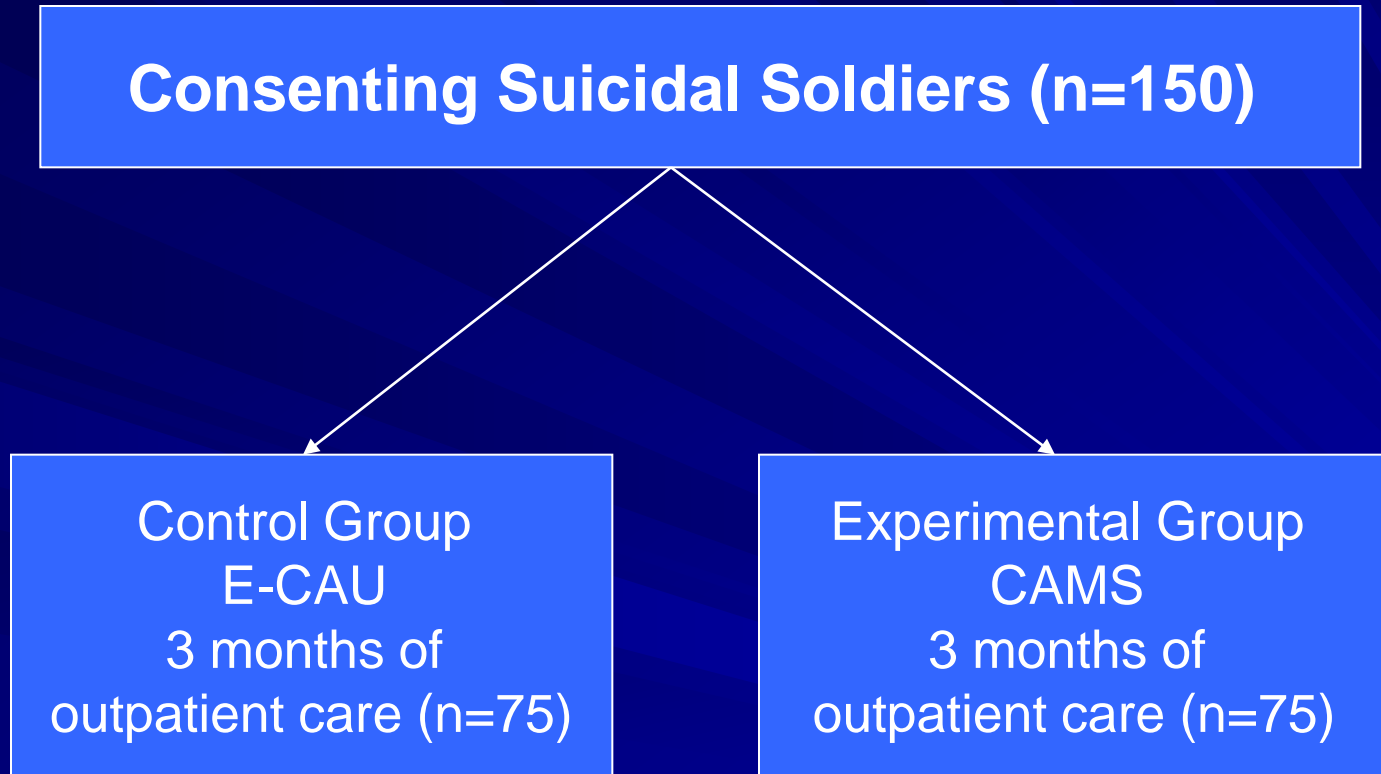
Operation Worth Living (OWL) Research Team



Overview of the OWL Project

- MOMRP awarded the CUA team a grant to conduct a large randomized clinical trial of the Collaborative Assessment and Management of Suicidality (2011-2015).
- In support of the project at Ft. Stewart, the Geneva Foundation will be hiring 3 FTE research staff and “back-fill” clinicians (to off-set study impact).
- Existing project with WRP/BAMC and new project WRNMMC are directly informing research methodology that will be used at FSGA.
- Thus far this research has received unparalleled support from Command.
- Members of the research team:
 - Dr. David Jobes (PI)—The Catholic University of America
 - Dr. Kate Comtois (Co-PI)—The University of Washington
 - Dr. Lisa Brenner (Co-PI)—Denver VAMC; University of Colorado
 - Dr. Peter Gutierrez (Co-PI)—Denver VAMC; University of Colorado
 - COL Bruce Crow, Psy.D. (Co-PI)—Warrior Resiliency Program
 - Mr. Brad Singer (Site-PI)—Ft. Stewart
 - Ms. Gretchen Ruhe (PC)—Geneva Foundation/Ft. Stewart
 - CAPT Philip McRae, Psy.D. (Chief, DBM)—Ft. Stewart

CAMS RCT at Ft. Stewart, GA



Dependent Variables: Suicidal Ideation/Attempts, Symptom Distress, Resiliency, Primary Care visits, Emergency Department Visits, and Hospitalizations.

Measures: SSI, OQ-45, SHBQ, SASIC, CDRISC, PCL-M, SF-36, NFI, THI (at 1, 3, 6, 12 months)

Reality of Progress

- IRB approval from four different institutions (11 months).
- Team wrote new CAMS Treatment Manual and revised existing versions of SSF and CAMS Rating Scale.
- Hired the study's participant coordinator; currently searching for "back-fill" clinicians to off-set impact of the study.
- Experimental arm training scheduled for 30 April to 2 May.
- Pilot phase of adherence consultation/training to begin in May.
- We estimate that study patients will be recruited and enrolled in late summer/early fall.

Recent site visit to FSGA



- Visit re-introduced the study.
- Recruited and consented five research therapists.
- Brigade command briefing.
- Discussed procedural and methodological study details.
- Oriented new our Participant Coordinator.





Bottom line:

We are making progress to conducting the study!