ACCEPTANCE AND COMMITMENT THERAPY FOR DEPRESSION WITH COMORBID SOCIAL ANXIETY: RESULTS FROM A PILOT RANDOMIZED TRIAL

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DISCLOSURES

• Grant support:

• The National Institute of Mental Health (K23MH085730; PI: Dalrymple) DEPRESSION AND SAD ARE HIGHLY COMORBID
SAD present in 1/3 of patients with depression

• Greater severity of symptoms

• Poorer functioning

• Shared clinical features

• Interpersonal rejection sensitivity, avoidance behaviors, social disengagement

EFFECTS ON TREATMENT OUTCOMES

• Pharmacotherapy studies:

- More severe course of illness (Mulder et al., 2006)
- Treatment-resistant depression (Souery et al., 2007)
- Greater number of depressive recurrences (Holma et al., 2008)

• Psychotherapy studies (CBT):

- Findings somewhat mixed
- Poorer outcomes (DeRubeis et al., 2005)
- No difference in rates of improvement (e.g., Erwin et al., 2002)
- One year follow-up: SAD severity worsened (Marom et al., 2009)

WHY ACT?

- Transdiagnostic approach
- Empirically-supported for depression (APA, SAMHSA)
- Preliminary support for SAD (e.g., Dalrymple & Herbert, 2007; Kocovski et al., 2009)
- ACT>CBT for comorbid anxiety and mood disorders at post-treatment and 1 year follow-up (Wolitzky-Taylor et al., 2012)

OPEN TRIAL STUDY (DALRYMPLE ET AL., 2014)

- Adults (n=38) recruited from routine outpatient psychiatric practice
- 16 sessions of ACT + medication treatment as usual
- High satisfaction with treatment
- Medium-to-large effect size changes on symptoms, functioning, and processes
- Psychological flexibility and behavioral activation correlated with symptoms and functioning

CURRENT STUDY AIMS

• Further assess feasibility and acceptability of ACT for Depression+SAD

• Test as adjunctive to medication as usual in a typical outpatient psychiatry setting

• Assess preliminary efficacy compared to medication as usual alone

Hypotheses

• ACT would continue to demonstrate feasibility and acceptability

- Initial signal efficacy:
 - Greater effect size change patterns compared to medication as usual alone

METHODS

PARTICIPANTS

• Recruited from outpatient psychiatry practice

• July 2012-January 2016

• Inclusion:

- Adults 18-65
- Diagnoses of depressive disorder and SAD
- Receiving medication management

• Exclusion:

- Bipolar/psychosis/borderline PD/severe substance
- Currently in psychotherapy

MEASURES

- Feasibility:
 - Recruitment targets and therapist training
- Acceptability:
 - Treatment Credibility (RTQ) and patient satisfaction (CSQ-8)
 - Study retention, completed sessions
- Symptoms, functioning, and processes:
 - Diagnoses (SCID & SIDP)
 - Depression severity (QIDS)
 - Social anxiety severity (LSAS)
 - Quality of life (QOLI) & functioning (WHODAS)
 - Behavioral activation (BADS) & psychological flexibility (AAQ-II)

TREATMENTS

• Medication As Usual (MAU)

- Med management by board-certified psychiatrist
- Flexible frequency of sessions
- Typical session length: 15-30 minutes

• MAU+ACT

- Med management as above
- Plus 16 individual sessions of ACT
- Flexibly designed protocol
- Therapists: doctoral students, residents, postdoctoral fellows

ACT SESSIONS

• Phases:

- Sessions 1-3: Identify avoidance, assess workability, introduce willingness
- Session 4: introduce exposure linked to values
- Sessions 5-9: defusion
- Sessions 10-11: self-as-context
- Sessions 12-13: formal values clarification
- Sessions 14-15: committed action
- Sessions 15-16: relapse prevention & post-tx planning

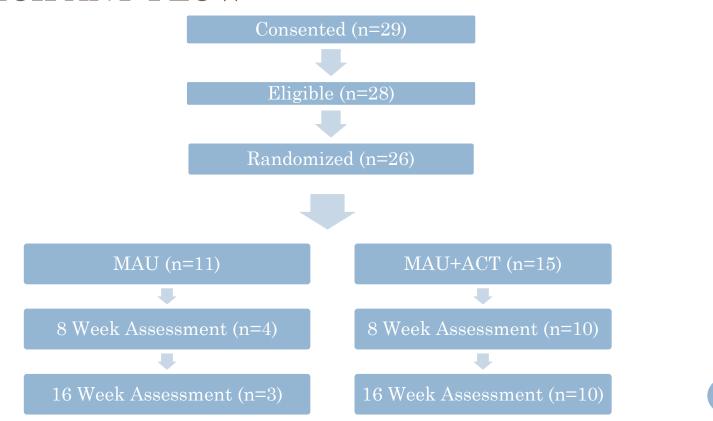
• Metaphors/experiential exercises used throughout

SAMPLE SESSION OUTLINE

- Session 9: Defusion
 - Review homework (e.g., engagement in valued behaviors, address barriers)
 - Experiential Exercise
 - e.g., Physicalizing, Content on Cards, Tin Can Monster
 - In-session exposure exercise
 - Choose valued behavior to practice
 - Link to personally-identified value
 - Role play, with emphasis on practicing defusion & orienting to value

RESULTS

PARTICIPANT FLOW



BASELINE CHARACTERISTICS

- Demographics:
 - Mean age = 38
 - 46% female
 - Majority Caucasian, single, <college degree

- Diagnoses:
 - Avg = 4 total current/partial remission diagnoses
 - Next most common: GAD (52%)
 - Other common: SUD, PTSD, panic, impulse control

BASELINE COMPARISONS

- Completers vs. dropouts
 - No differences on demographics or baseline assessments

• MAU vs. MAU+ACT

• No differences on demographics or baseline assessments

PHARMACOTHERAPY BETWEEN CONDITIONS

Variable	MAU	MAU+ACT
Number of medication visits (<i>M</i>)	2.2	3.1
Medication changes made (% of visits)	77.8%	46.2%

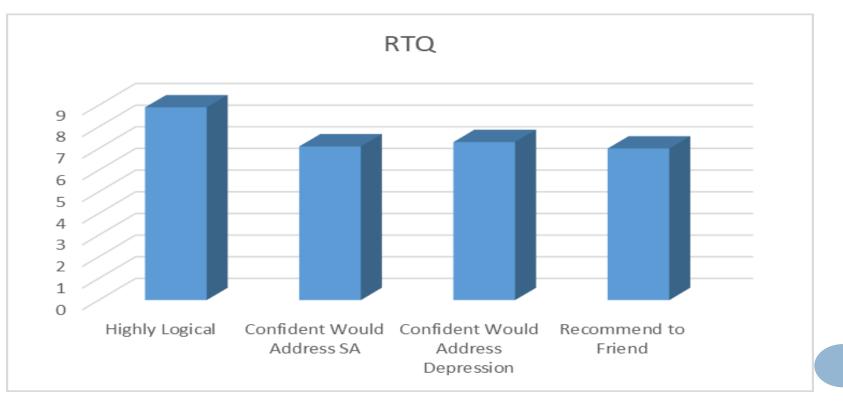
FEASIBILITY

- Able to feasibly recruit
- Trained 7 therapists on protocol
- Drop-out rates:
 - MAU = 64%
 - MAU+ACT = 33%
- Of MAU drop-outs:
 - 3 discontinued treatment completely
- MAU+ACT session completion:
 - 6 patients completed at least 10 sessions over 16 weeks
 - 6 completed all 16 sessions, over avg. of 20 weeks

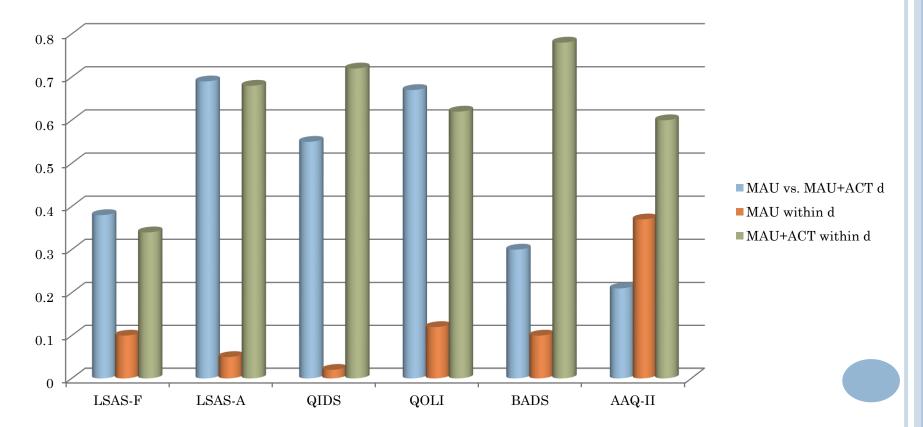
ACCEPTABILITY (MAU+ACT)

- Patient Satisfaction (CSQ-8):
 - Assessed after final session
 - Highly satisfied with treatment (*M* = 30; out of 32 pts.)

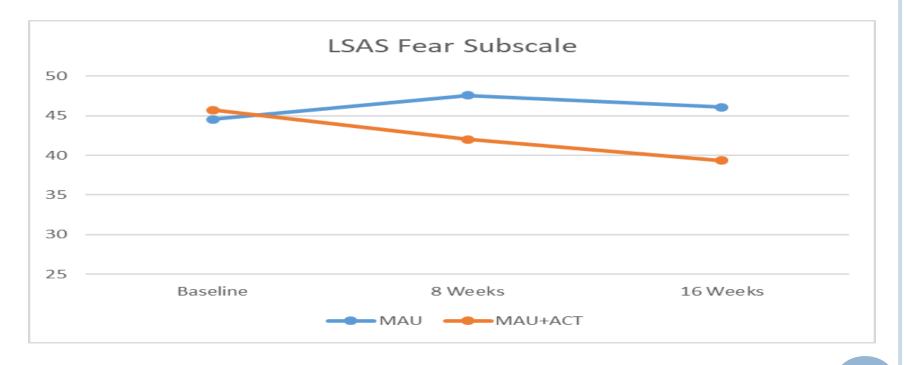
TREATMENT CREDIBILITY



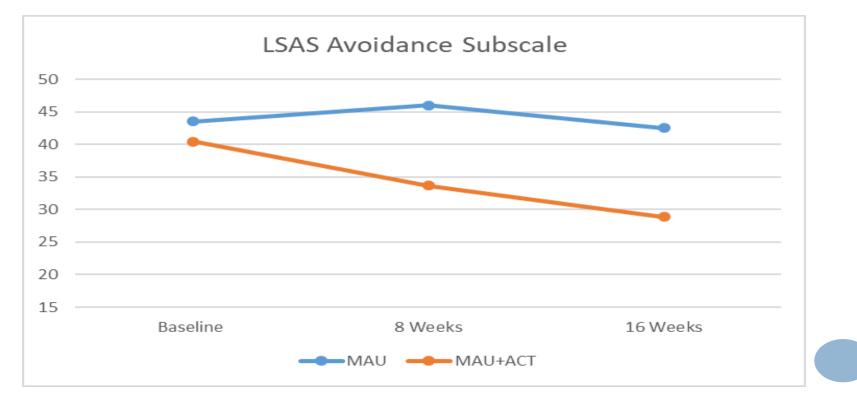
PRELIMINARY TREATMENT OUTCOMES



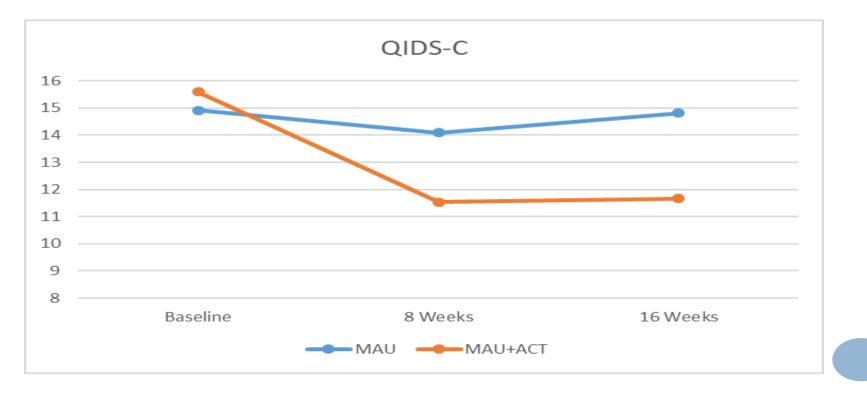
SOCIAL FEAR



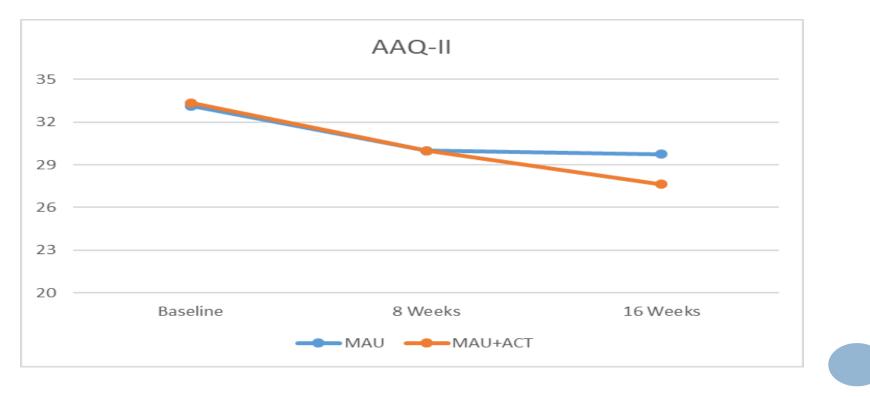
SOCIAL AVOIDANCE



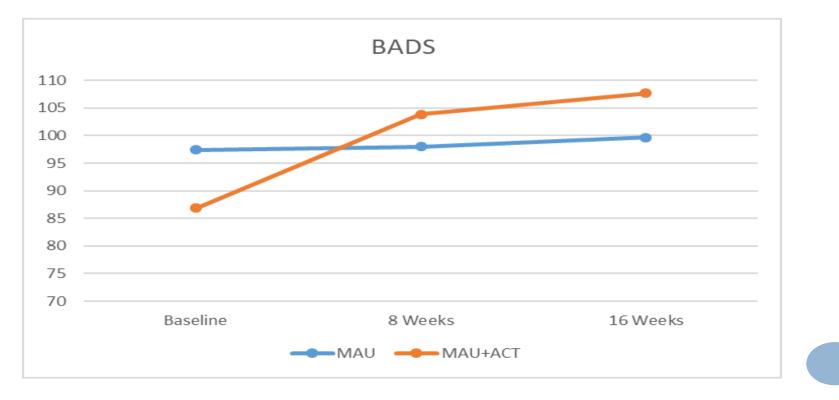
DEPRESSION SEVERITY



PSYCHOLOGICAL FLEXIBILITY (LOWER=BETTER)



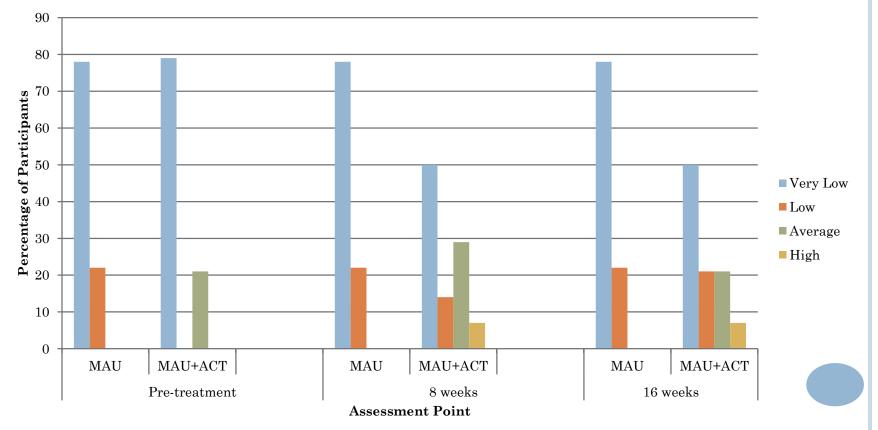
BEHAVIORAL ACTIVATION (HIGHER=BETTER)



Reliable Change

Measure	Completer		ITT	
	MAU	MAU+ACT	MAU	MAU+ACT
QIDS-C (Depression)	0%	77.8%	9.1%	46.7%
LSAS-F (Social Anxiety Fear)	0%	22.2%	0%	13.3%
LSAS-A (Social Anxiety Avoidance)	33.3%	55.6%	9.1%	33.3%

QUALITY OF LIFE



DISCUSSION

SUMMARY OF RESULTS

- Good feasibility and acceptability
- Preliminary signal of efficacy
 - Within MAU+ACT Cohen's *d*:
 - Depression (0.72), SA avoidance (0.68)
 - Psych flex (0.60), BA (0.78)
- But high attrition in MAU (~66%)
 - Nearly $\frac{1}{2}$ of MAU dropouts ended treatment completely at the practice

PREMATURE TERMINATION IN MAU

• Potential reasons:

- Receiving non-preferred treatment
 - Depression: patients prefer psychotherapy over medication (McHugh et al., 2013)

- Consistent with recent meta-analysis (Swift et al., 2017)
 - Rates as high as 69% in comparative treatment trials
 - SAD: 1.29 OR, with higher rate in meds alone

PRELIMINARY EFFICACY

- Results comparable to open trial
- Encouraging results with quality of life
- Consistent with effectiveness studies
 - Effect sizes 1.06-1.13 for depression (Hans & Hiller, 2013)
 - Reliable change 50%, 15-30% (McEvoy & Nathan, 2007; Westbrook & Kirk, 2005)

LIMITATIONS

- Small sample size
- Large confidence intervals
- High MAU attrition rate
- Intention-to-Treat: LOCF method
- Unstructured MAU

FUTURE IMPLICATIONS

• Comparison with other active psychotherapies

• Improve study retention, especially in MAU

• Further examination of target mechanisms

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• Questions?

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