

# CLINICAL WEBINARS FOR HEALTH SERVICE PSYCHOLOGISTS

TRANSLATING RESEARCH TO PRACTICE

## Recent Advances in the Treatment of PTSD

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#### Mark B. Powers, PhD



Dr. Mark Powers is a licensed clinical psychologist and the Director of Trauma Research at Baylor University Medical Center. His research is funded by NIH, DoD, DARPA, and HHS spanning three areas: a) treatment and prevention of PTSD, b) resilience training, and c) non-opioid pain management. Dr. Powers was certified in prolonged exposure therapy for PTSD at the University of Pennsylvania with Dr. Edna Foa and he was a Beck Scholar at the Beck Institute for Cognitive Therapy and Research. He has over 150 publications and is the Editor-in-Chief of the journal *Cognitive Behaviour* Therapy.



### Disclosures/Conflicts of Interest

• Dr. Powers receives book royalties from Oxford University Press and Academic Press.



#### Learning Objectives

At the conclusion of this webinar, participants will be able to:

- Describe the nature and causes of PTSD
- Discuss the efficacy of Prolonged Exposure Therapy (PE) for PTSD
- Apply the principles of Prolonged Exposure Therapy in trauma focused care



#### Overview

- A word about anxiety
- PTSD Past, Present, Future
  - Definition of "Trauma"
  - PTSD Symptoms & Diagnosis
  - What causes/maintains PTSD?
  - What are the most effective treatments for PTSD?
  - Therapy augmentation strategies
  - PTSD prevention
- Summary



## A word about anxiety





- PTSD
- Panic Disorder (and agoraphobia)

#### Disorders:

- OCD
- Social Anxiety Disorder (& public speaking)
- Generalized Anxiety Disorder
- Specific Phobias (e.g. storms, dark, spiders, heights, flying, claustrophobia)



#### Introduction to PTSD

- How PTSD is Unique
  - Only disorder we know how/when it started
    - Criterion A
  - Only disorder with Re-experiencing
  - PTSD & OCD = "Specific Phobia with a much cooler story"



#### Trauma Definition: Criterion A

- Exposure to <u>actual or threatened death</u>, <u>serious injury</u>, <u>or sexual</u>
   <u>violence</u> in one of the following ways
  - Direct experience
  - Witnessing the event
  - Learning that the trauma occurred to a close family member or friend
  - Experiencing repeated or extreme exposure to details of traumatic events (e.g. trauma surgeons, first responders, police officers).



## **Example Traumas**

- Natural Disasters
- Collisions (e.g. motor vehicle, motorcycle)
- Physical Assault
- Molestation
- Rape
- Combat
- Working directly with child suicides, homicides, sex trafficking
- NOT
  - Mental/Verbal Abuse
  - Difficult Childhood
  - Parental Divorce
  - Extreme Disappointments
  - Difficult breakups
  - Television Coverage of Traumas (unless informing you of a loved one etc.)



## Most People will Experience a Trauma

- PTSD Diagnosis first appears in DSM-III (1980)
- DSM-III & DSM-III-R
  - "Outside the range of usual human experience"
  - Removed now (DSM IV & 5) given 80-90% of Americans will experience a
     Trauma

• (Kilpatrick, Resnick, Milanak, Miller, Keyes, & Friedman, 2013; Sledjeski, Speisman, & Dierker, 2008 from the NCS-R)



## PTSD Symptoms & Diagnosis (1 month or more)

- Intrusion (need 1) Criterion B
- Avoidance (need 1) Criterion C
- Negative Mood and Cognitions (need 2) Criterion D
  - World is dangerous and I cannot cope
- Hyperarousal (need 2) Criterion E

\*Acute Stress Disorder: Symptoms 3 days to 1 mo



#### Prevalence

• Lifetime: 8.7%

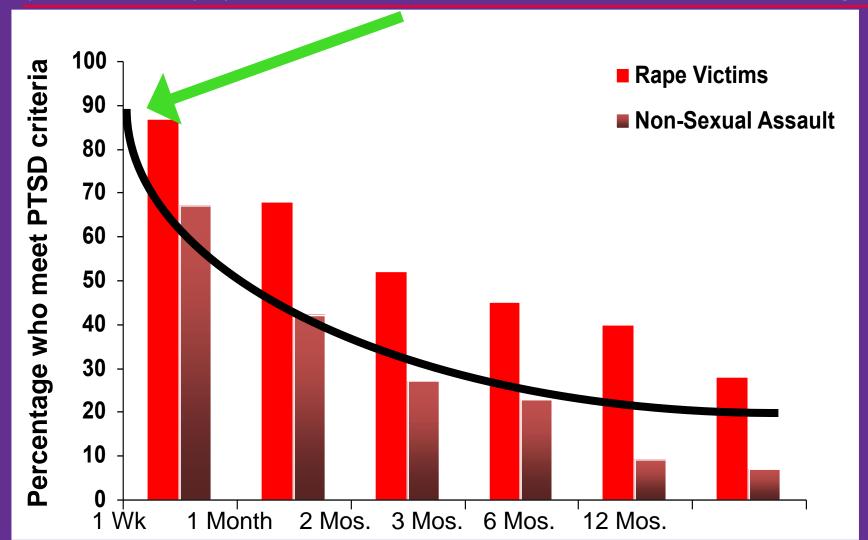
• 12-Month: 3.5%

- \*30-50% among rape, combat, and captivity survivors
- \*Better with age? Rates are lower among older adults

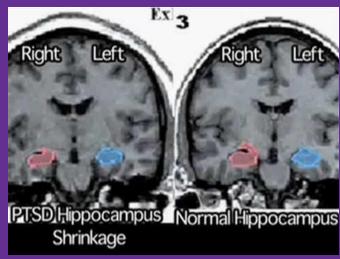


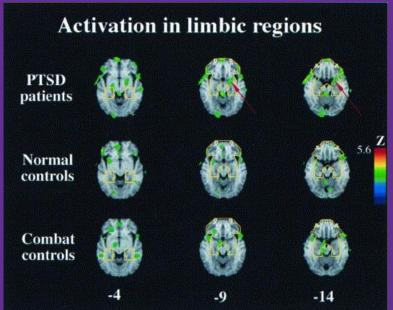
#### **After Trauma**

Most people have PTSD symptoms (Re-experiencing, Avoidance, Changes in Cognition and Mood, & Hyperarousal) but get better



#### **Predictors of PTSD**





- Lower SES
- Female
- Younger
- Lack Hlth Ins
- Smokers
- Hx of MDD, other Anx Dis, and SUDs
- # of traumas/prior trauma Hx
- Trauma experienced directly
- Penetrating wounds
- Hippocampus Atrophy
- Norepinephrine & Cortisol
- Genes (BDNF SNP Val66Met)
- HPA Axis
- Low SES Social & Family Psych Hx
- Gunshot, younger age, psych Hx = 78% acuracy
- Peritraumatic Dissociation largest in one meta-analysis but still only explained 9% of the variance VIDEO

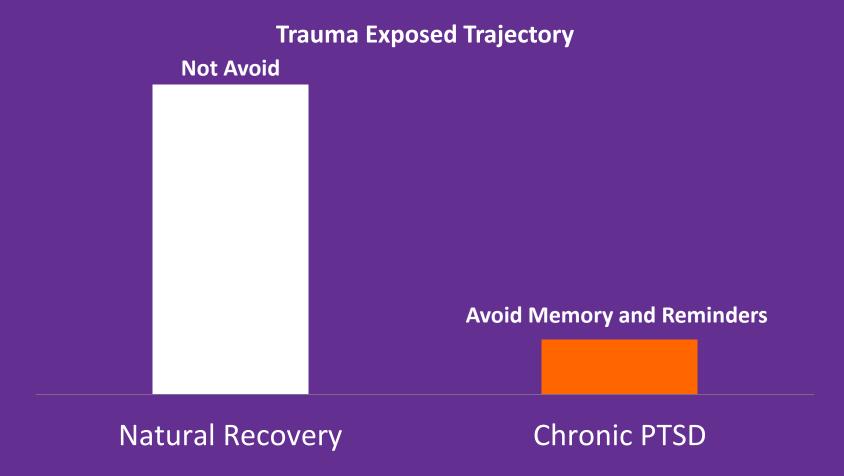


### **Summary of Predictors**

- Younger victims with pre-existing psychiatric histories
- Higher injury severity
- Females
  - Males experience more traumas externalizing
  - Females are most likely to maintain symptoms following trauma internalizing
- Victims of Interpersonal & Intentional Traumas (Rape, assault, combat)
- Comorbid Substance Use Disorders
- Lack of social support



### What is the most reliable predictor?



Breslau et al. 1999; Foa, Stein, & McFarlane, 2006; Gil & Caspi, 2006; Karamustafalioglu et al. 2006; Maes et al. 1998; Myers, VanMeenen et al. 1012; Myers, VanMeenen, McAuley et al. 2012; North et al. 2004; North et al. 1999; O'Donnell et al. 2007; Solomon et al. 2009; Yoon et al. 2009



## How do we measure/diagnose PTSD?

#### Foa Measures

- PSSI-5: Posttraumatic Stress Disorder Symptom Scale Interview for DSM-5 (Interview)
- PDS-5: Posttraumatic Diagnostic Scale for DSM-5 (Self-report)

#### Other Example Measures

- CAPS-5: Clinician-Administered PTSD Scale for DSM-5 (Interview)
- PCL-5: PTSD Checklist for DSM-5 (Self-report)



#### How do we Treat PTSD?



## Treatments for PTSD Learn To:

- Live/cope with PTSD
  - Medications
  - Talk Therapy/Counseling
  - Support Groups
  - Service Dogs
  - Relaxation
  - Yoga
  - Meditation
  - Coping Strategies
  - Exercise

- Live without PTSD
  - Trauma focused Exposure Therapies
    - PE, CPT, EMDR



## Prolonged Exposure Therapy (PE) for PTSD

9-12 weekly or twice weekly 90-minute sessions

- Psychoeducation (sessions 1-2)
- In vivo exposure (sessions 2-12)
  - to trauma reminders in life as homework
- Imaginal exposure (sessions 3-12)
  - revisiting the trauma memory (30-45 minutes during sessions)
- Processing (sessions 3-12)
  - Following each imaginal exposure session (15 minutes)

- \*Exposure needs to be:
  - Systematic, Deliberate, Repeated, Prolonged
- \*While fading safety behaviors/avoidance

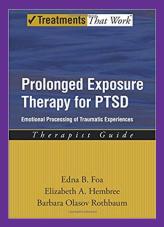


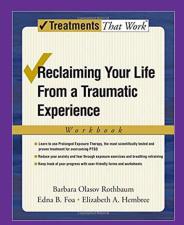
Session	Components	Homework
Phone Contact	Identify Patient's Core Threats	<ul> <li>Fill out Intake Packet Including Initial Measures</li> </ul>
Intake	<ul> <li>Set agenda</li> <li>Presenting Problem</li> <li>Confirm Patient's Core Threats</li> <li>Working Diagnoses</li> <li>Initial Treatment Plan</li> </ul>	Fill out Domain Specific     Measures
1	<ul><li>Set agenda</li><li>Overall Treatment Rationale</li><li>Gather information about symptoms</li><li>Assign Homework</li></ul>	<ul> <li>Listen to the session recording once</li> <li>Fill out the PDS-5, PTCI, and PHQ-9</li> </ul>
2	<ul> <li>Set agenda</li> <li>Homework review</li> <li>Common reactions to trauma</li> <li>Rationale for in vivo exposure</li> <li>Introduce SUDs scale</li> <li>Create in vivo hierarchy</li> <li>Assign homework</li> </ul>	<ul> <li>Listen to the session recording once</li> <li>In vivo exposure (60mins/day)</li> <li>Fill out the PDS-5, PTCI, and PHQ-9</li> </ul>
3	<ul> <li>Set agenda</li> <li>Homework review</li> <li>Rationale for imaginal exposure</li> <li>Imaginal exposure</li> <li>Processing</li> <li>Assign homework</li> </ul>	<ul> <li>Listen to the session recording once</li> <li>In vivo exposure (60mins/day)</li> <li>Imaginal exposure (60mins/day)</li> <li>Fill out the PDS-5, PTCI, and PHQ-9</li> </ul>
4-11	<ul><li>Same as session 3</li><li>Session 6 or 7 imaginal exposure Hot Spots</li></ul>	Same as session 3
12	<ul> <li>Termination</li> <li>Homework review</li> <li>Imaginal exposure (entire memory)</li> <li>Review of the hierarchy</li> <li>Processing</li> <li>Relapse prevention</li> </ul>	

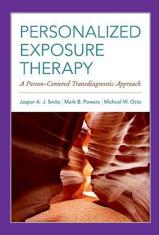


#### PE Training Opportunities & Resources

- PE Training by Dr. Edna Foa
  - https://www.med.upenn.edu/ctsa/workshops\_ptsd.html
- PE for Veterans: STRONG STAR Training Initiative
  - <a href="https://www.strongstartraining.org/what-we-do">https://www.strongstartraining.org/what-we-do</a>
- Manuals by Dr. Foa
- Chapter on PE by Dr. Powers







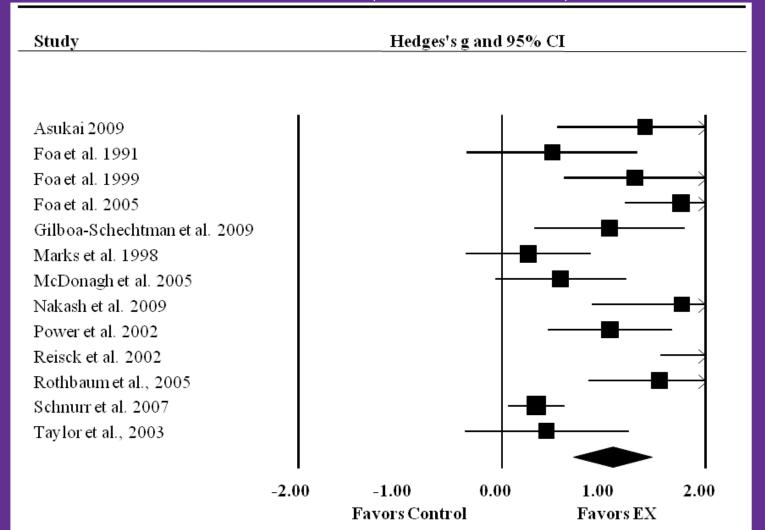


#### How Effective is PE?



#### Meta Analysis of 13 PE RCTs

Large effect size (g = 1.08) relative to control conditions Average PE-treated patient fared better than 86% of those in the control conditions NNT = 2.49 (SSRI NNT for MDD = 9)



#### Overall

- Posttreatment:
  - 85% response rate (significantly better)
  - Loss of diagnosis (no longer meet criteria for PTSD)
    - 50% (Intent to treat)
    - 70% (Completers)
  - 20% dropout rate
- Long-term follow-up
  - Loss of diagnosis 80%
- \*Recent data may suggest more modest outcomes with military personnel
- Relative Efficacy/Effectiveness & Dropout
  - Similar to other trauma focused therapies (e.g. CPT, EMDR)

(Bisson et al. 2013; Bradley et al. 2005; Cusa ck et al. 2016; \*Foa et al. 2018; Powers et al. 2010; Resick et al. 2012; Watts et al. 2013)

#### Medications

- FDA Approved
  - Zoloft
  - Paxil



## Room for Improvement?

- 20% of patients dropout
- 15% of completers do not respond
- More modest outcome in military personnel



#### Augmentation

Improve:
Tolerability, Efficacy, Speed,
Generalizability



## Choose between Efficacy OR Tolerability

#### **Increase Tolerability = Decrease Efficacy**

- Benzodiazepines, Alcohol, Safety Behaviors, Avoidance
  - Interfere with consolidation of extinction learning
  - Misattribution of treatment gains (I'm ok as long as I have my benzo) patient learns conditional safety



## Choose between Efficacy OR Tolerability

#### **Increase Efficacy = Decrease Tolerability**

- Cognitive Enhancers: Yohimbine hcl, DCS, Methyline blue, Exercise
- Increase efficacy and durability of gains
- But can decrease tolerability (Yohimbine/Exercise) and enhance whatever is learned:
  - Good sessions (e.g. low end fear)
  - Bad sessions (e.g. high end fear)



## Vagus Nerve Stimulation may accomplish both



#### Vagus Nerve Stimulation

- Anxiolytic: 20% efferent <u>stimulation of parasympathetic NS</u>
- Cognitive enhancer: 80% afferent stimulation of brain plasticity
- Preclinical studies show 10-30 times faster/more effective extinction in animal model of PTSD
- VNS in 80,000 patients for other indications (epilepsy, depression, tinnitus, stroke rehab)
- **But:** Existing Device is <u>permanent</u>, <u>wires get damaged</u>, <u>clavicle</u> <u>stimulator large</u>



## DARPA, BUMC, UTSW, UTD Trial

#### New device

- Smaller (1-inch incision left side)
- Single incision (30 minutes or less)
- Removable
- No wires
- No clavicle implant

#### New Stimulation Parameters

- .5 second .8mA every 7 seconds during exposure
  - Most effective
  - Prevents tolerance
- First in human trial for device
- First in human trial of VNS for PTSD



#### **OFFICE THERAPY**



The patient participates in therapy with a licensed therapist once a week.



The therapist uses a smartphone app to record the therapy and the triggers, which activate the vagus nerve stimulator.

#### **HOME THERAPY**



The patient uses a smartphone app to listen to their weekly recorded therapy sessions while receiving vagus nerve stimulation at home.



## Final Research Target: Prevention

- Multisite RCT of secondary prevention of PTSD in Level 1 Trauma Centers
- Intervention delivered via tablet:
  - Trauma/PTSD Psychoeducation

+

 Visuo-spatial task to interrupt consolidation/reconsolidation of trauma memory



### Summary

- Who is at risk for PTSD?
  - People who push away trauma reminders
- What causes PTSD?
  - Primarily Avoidance (among a host of other predictors)
- Effective treatments for PTSD?
  - Variants of trauma focused therapy (e.g. variants of CBT: PE, CPT, EMDR)
- Recent Advances
  - VNS augmentation may increase tolerability and efficacy of trauma focused therapy for PTSD
  - Visuo-Spatial tasks and psychoeducation may lead to secondary prevention of PTSD



#### References/Citations

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#### Q&A



- Dr. Sammons will read select questions that were submitted via the Q&A feature throughout the presentation.
- Due to time constraints, we will not be able to address every question asked.



### Thank You for Joining Us!

- Attendees eligible for CE credit may refer to the chat box for a direct link to the CE evaluation—or you may type this URL into a new browser window: bit.ly/ptsdeval
- If you have comments or feedback regarding this webinar, please email CESupport@nationalregister.org
- We hope you can attend our next webinar on June 12: Clinical Exemplars for Primary Care and Brief Treatment Settings with Bridget Beachy, PsyD.





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