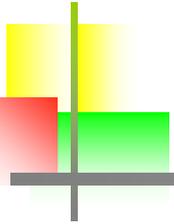


Care for Patients with Complex and High-Impact Chronic Pain



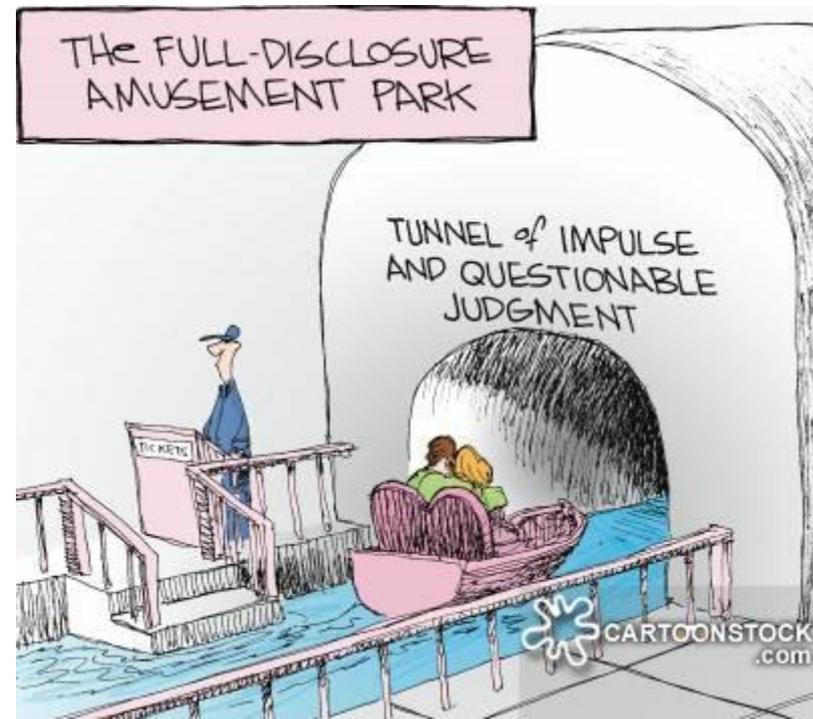
*Rob Edwards, Ph.D.
Pain Psychologist
BWH Department of
Anesthesiology*



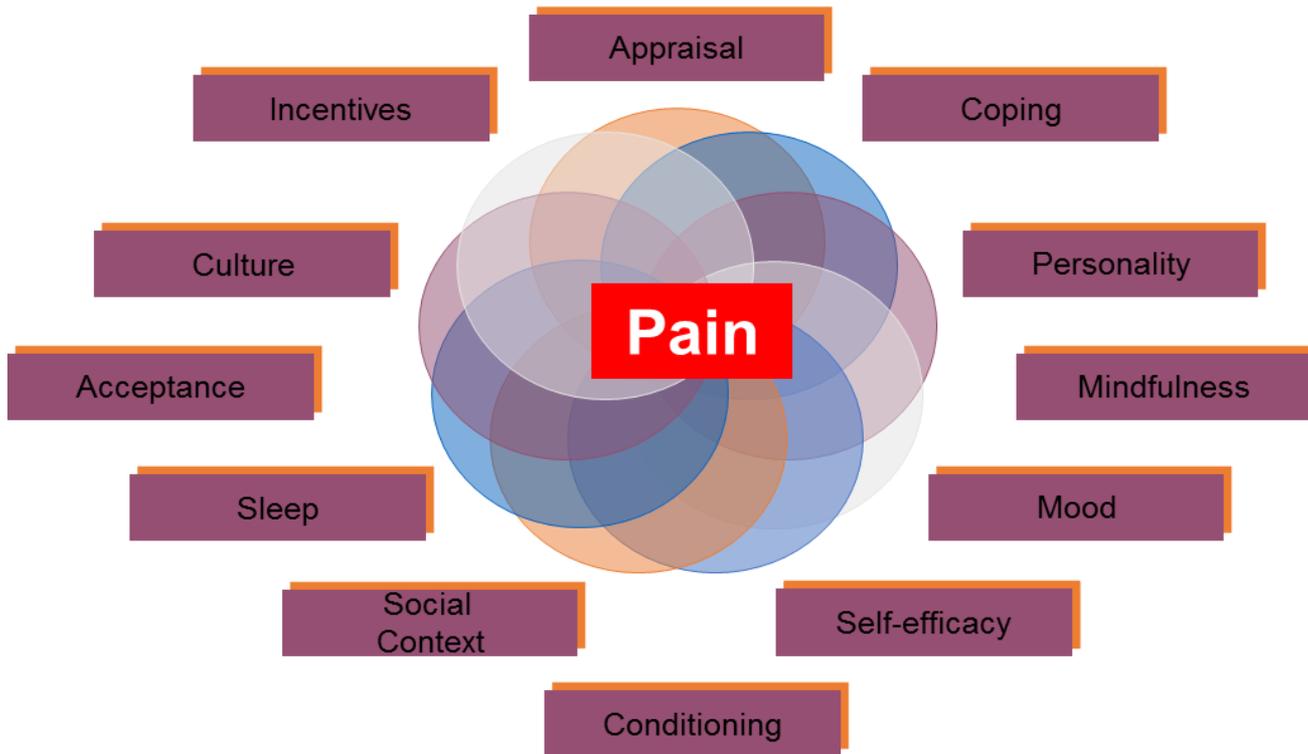
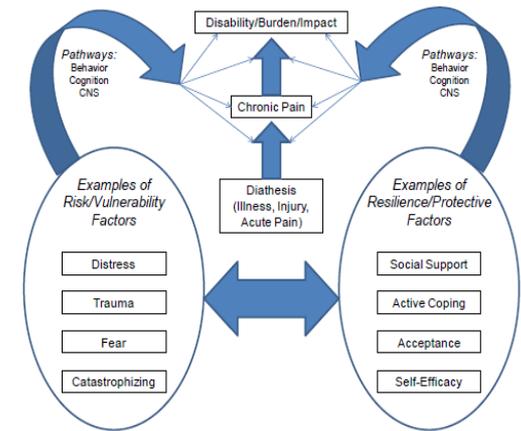
Disclosures

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- National Institutes of Health
- Patient-Centered Outcomes Research Institute



Depicting the Biopsychosocial Model of Pain



American Pain Society RESEARCH EDUCATION TREATMENT ADVOCACY PUBLISHED BY The Journal of Pain, Vol 17, No 9 (September), Suppl. 2, 2016; pp T70-T92 Available online at www.pain.org and www.sciencedirect.com ELSEVIER

The Role of Psychosocial Processes in the Development and Maintenance of Chronic Pain 

Robert R. Edwards,^{*} Robert H. Dworkin,[†] Mark D. Sullivan,[‡] Dennis C. Turk,[§] and Ajay D. Wasan[¶]

Clusters of risk factors are associated with high-impact chronic pain

Women
Older age
Minority
HS Edu
Not married
Poverty
Rural
Public Health
Insurance

Comorbid
medical
conditions
Past surgeries

Depression
Anxiety
Fatigue
Cognitive
limitations

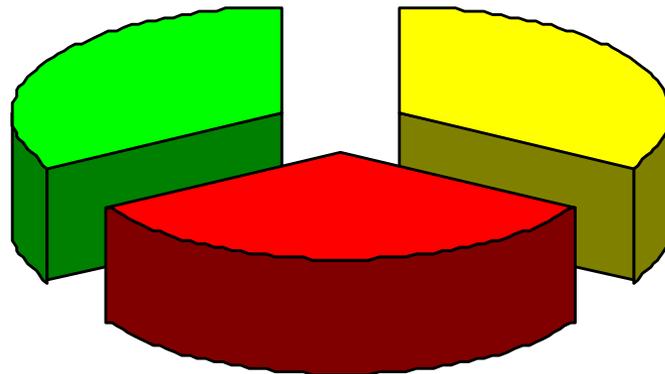


Catastrophizing

I worry all the time about whether the pain will ever stop

I feel I can't stand the pain

I wonder whether something serious may happen



-  Rumination
-  Magnification
-  Helplessness



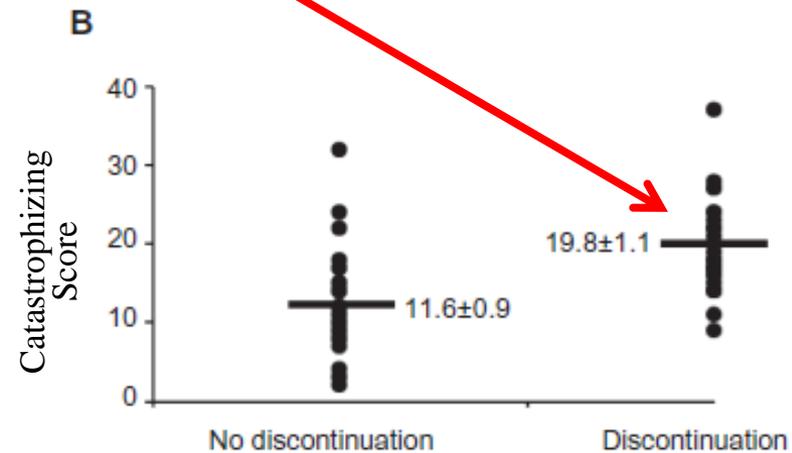
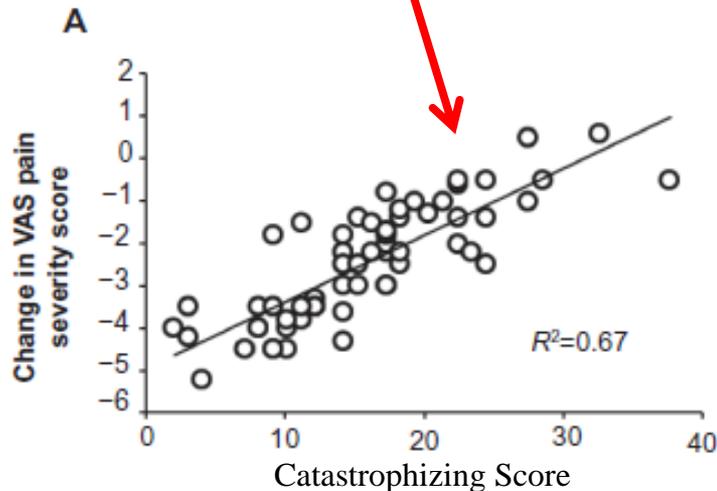
Risk factor for poor treatment outcomes

The importance of catastrophizing for successful pharmacological treatment of peripheral neuropathic pain

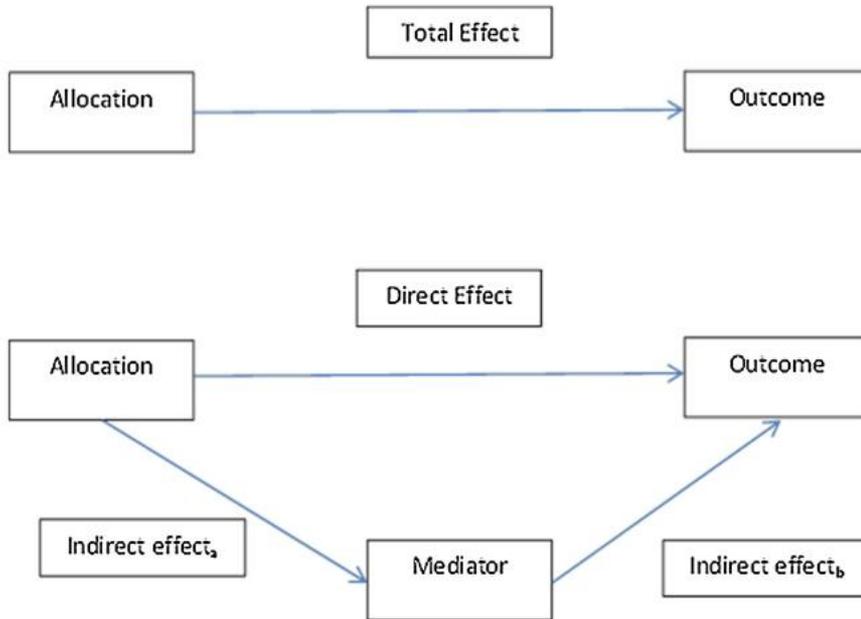
Cory Toth
Shauna Brady
Melinda Hatfield

Table 3 Medication interventions provided to patients

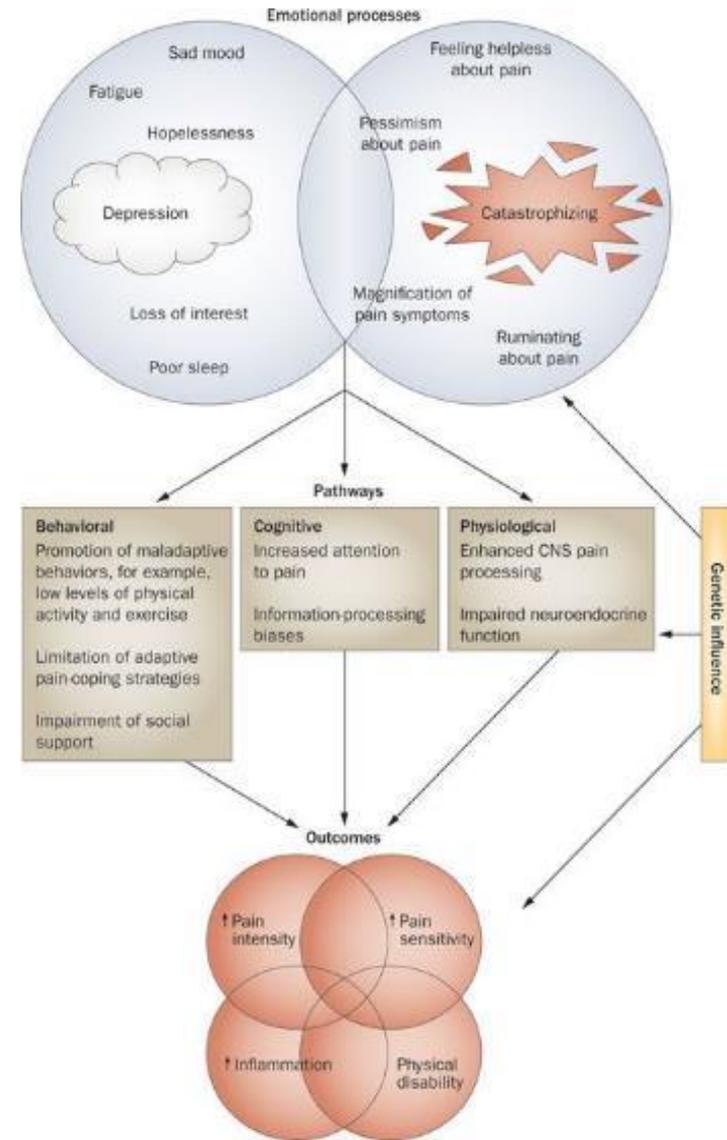
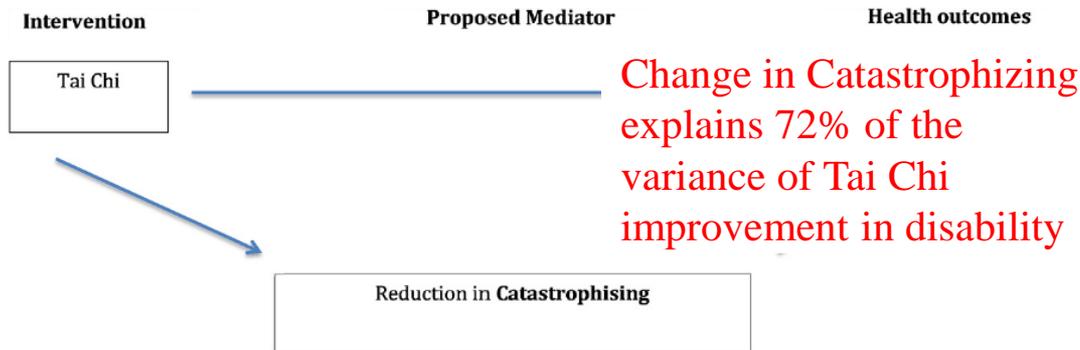
	Number of patients receiving specific medication	Initial medication daily dose	Final medication daily dose	Discontinuations
Amitriptyline	25 (40%)	15.1±5.2 mg	37.3±12.7 mg	14/25 (56%)
Nortriptyline	10 (16%)	25.5±11.3 mg	55.7±13.3 mg	5/10 (50%)
Gabapentin	17 (27%)	667.0±112.3 mg	1,562.7±147.8 mg	7/17 (41%)
Pregabalin	10 (16%)	120.5±22.3 mg	363.3±78.8 mg	4/10 (40%)



Process Variable



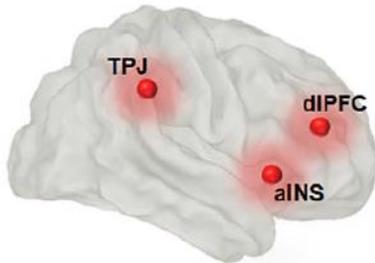
A.M. Hall et al. / Complementary Therapies in Medicine 25 (2016) 61–66



Neural Underpinnings/ Biomarkers?

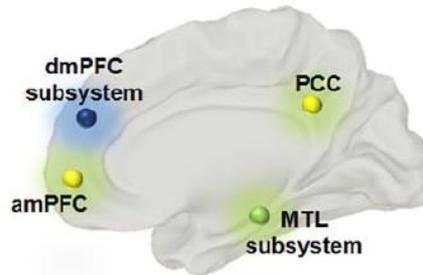
Salience Network

Sustained activation during attention to pain



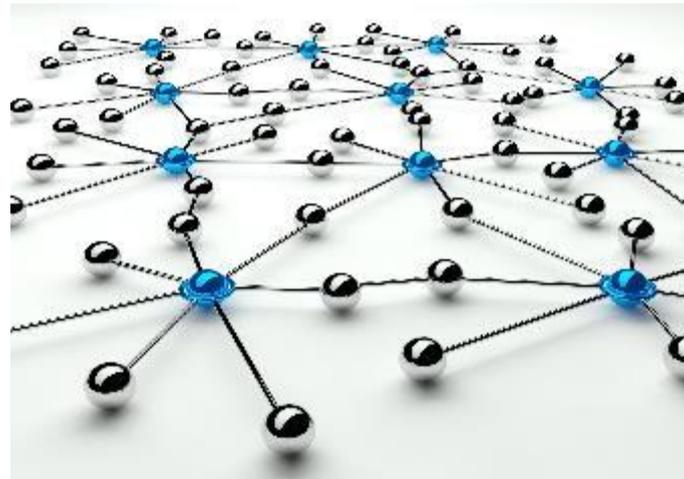
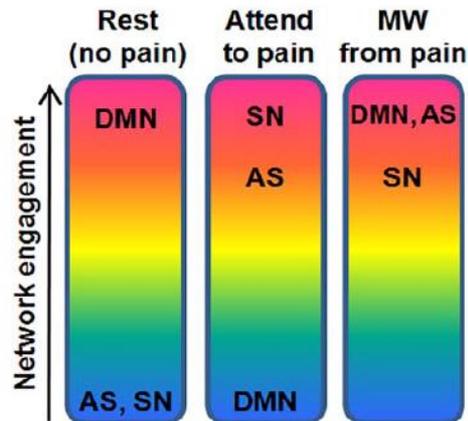
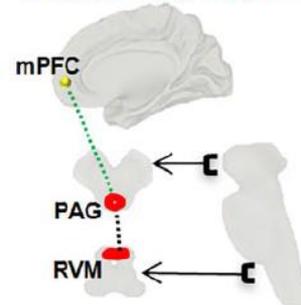
Default Mode Network

Suppressed when attending to pain but not when mind wandering away



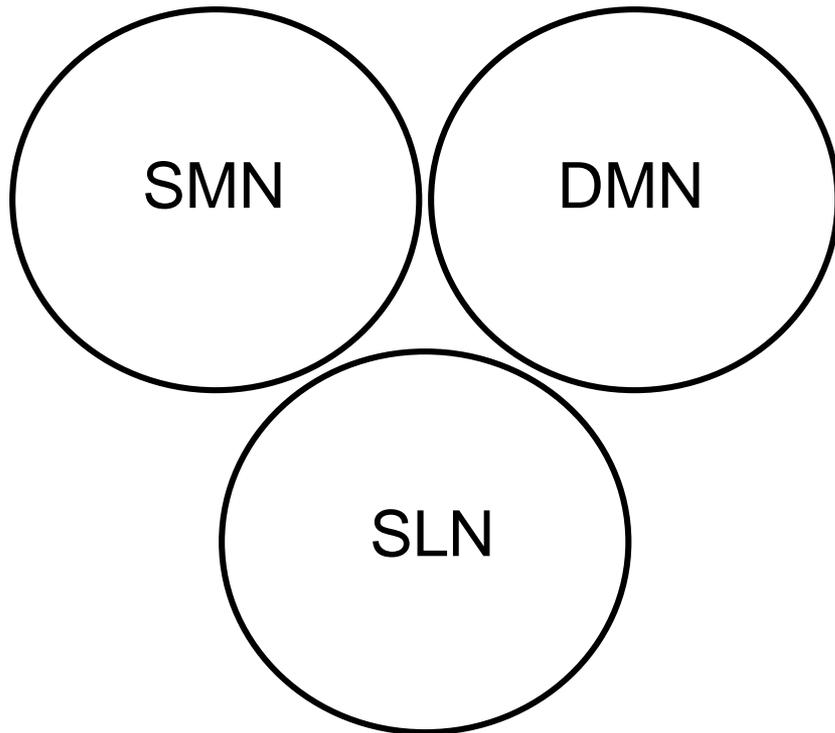
Antinociceptive System

Increased functional connectivity during mind wandering away from pain



Chronic Pain → Blurred Network Connectivity

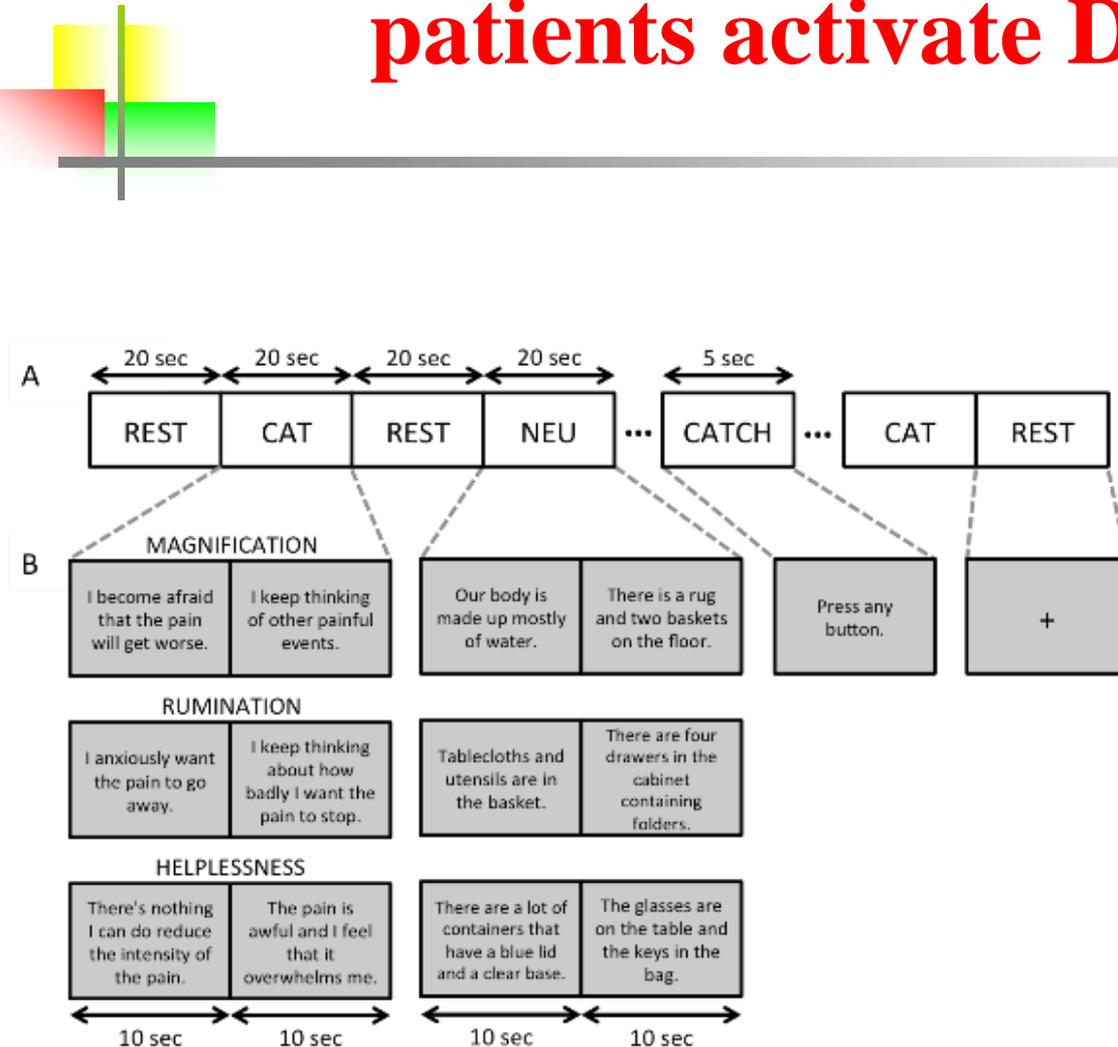
Healthy



Chronic Pain

Do pain-related catastrophizing and overlapping psychosocial processes contribute to this blurring?

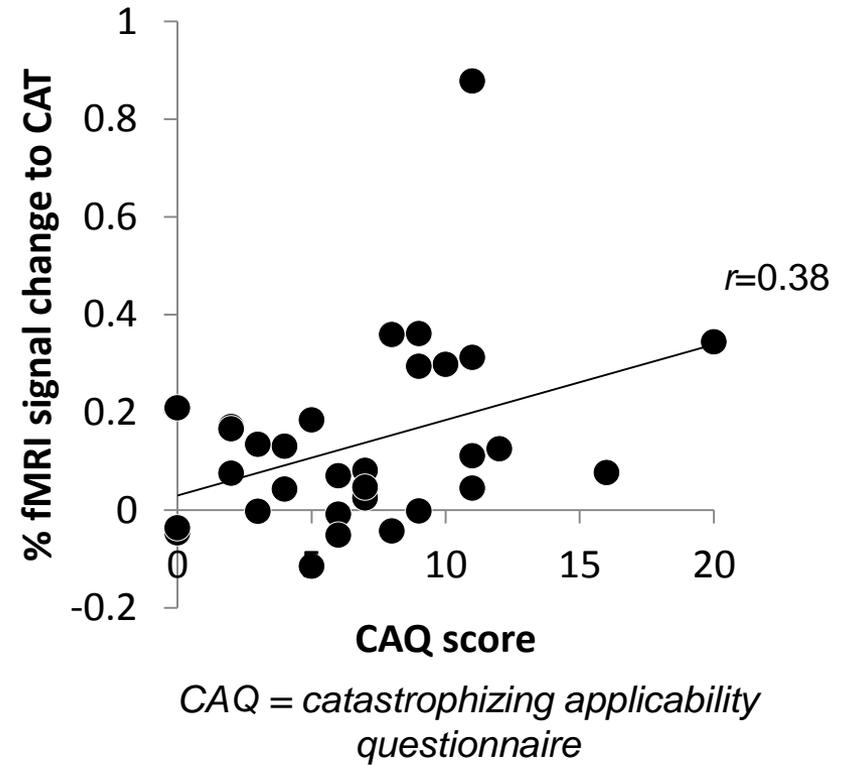
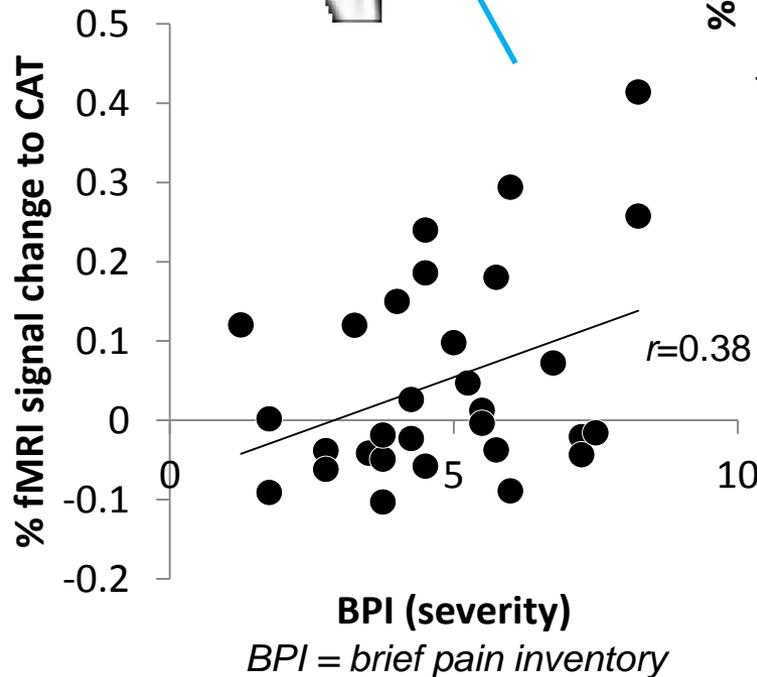
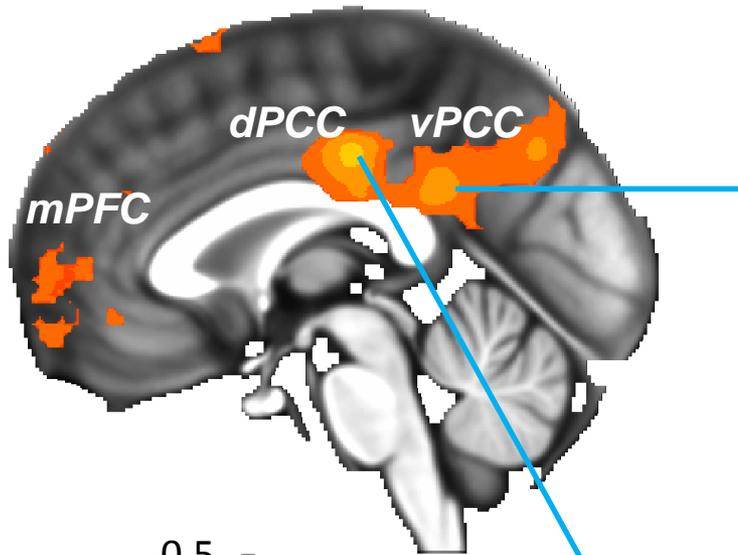
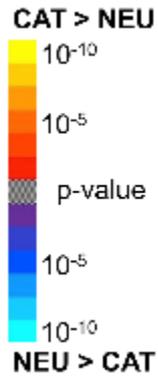
Catastrophizing cognitions in FM patients activate DMN circuits



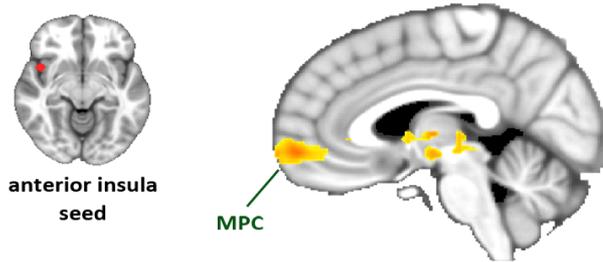
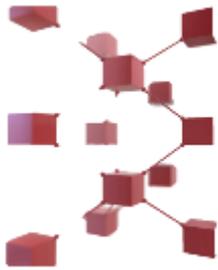
Objective. We designed a functional MRI-based pain catastrophizing task whereby patients with chronic pain engaged in catastrophizing-related cognitions compared to affectively neutral cognitions.

Methods. FM patients (n=31) reflected on how catastrophizing statements (CAT) impact their typical fibromyalgia pain experience. Response to CAT statements was compared to matched neutral statements (NEU).

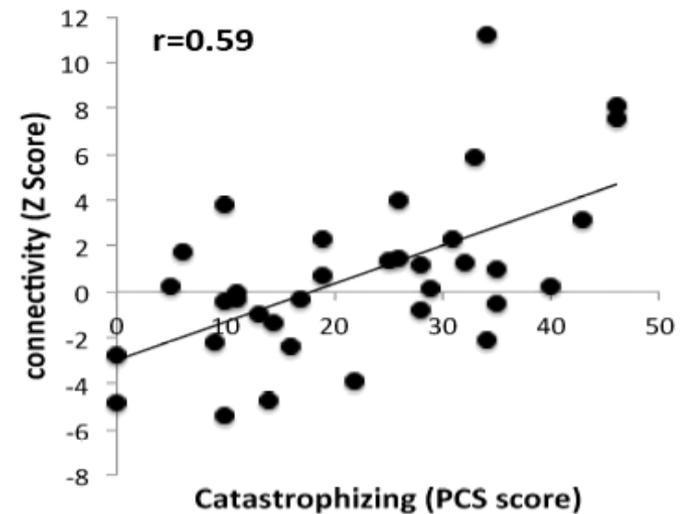
Neural circuitry supporting catastrophizing



Catastrophizing: Associations with connectivity between salience network (Insula) and DMN



ains-MPC Connectivity during PAIN vs. CAT

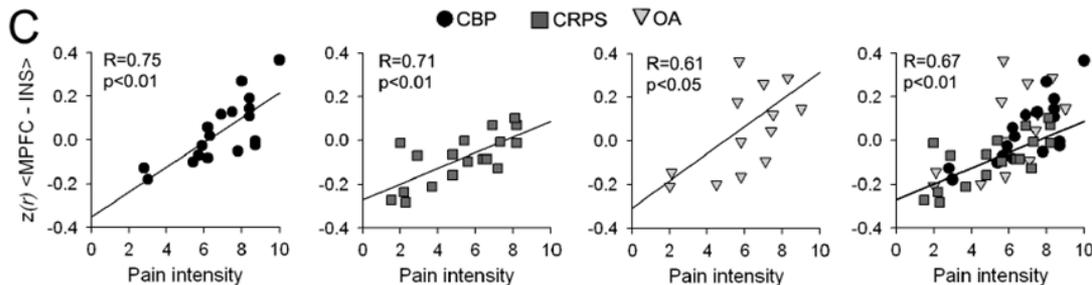


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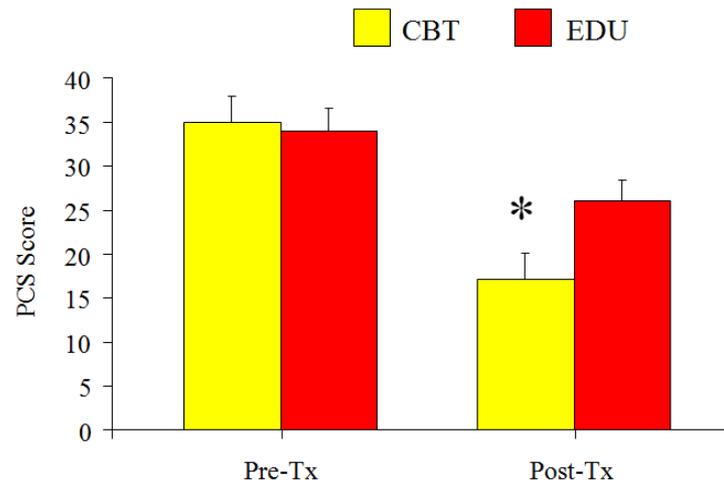
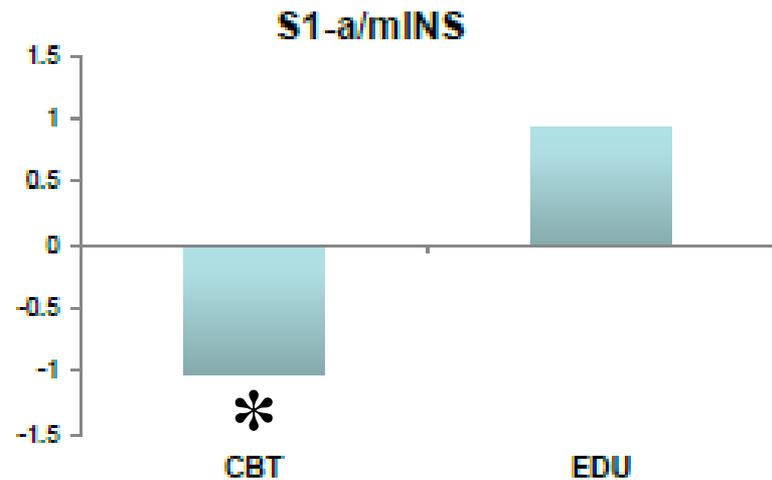
Functional Reorganization of the Default Mode Network across Chronic Pain Conditions

Marwan N. Baliki¹, Ali R. Mansour¹, Alex T. Baria¹, A. Vania Apkarian^{1,2,3*}

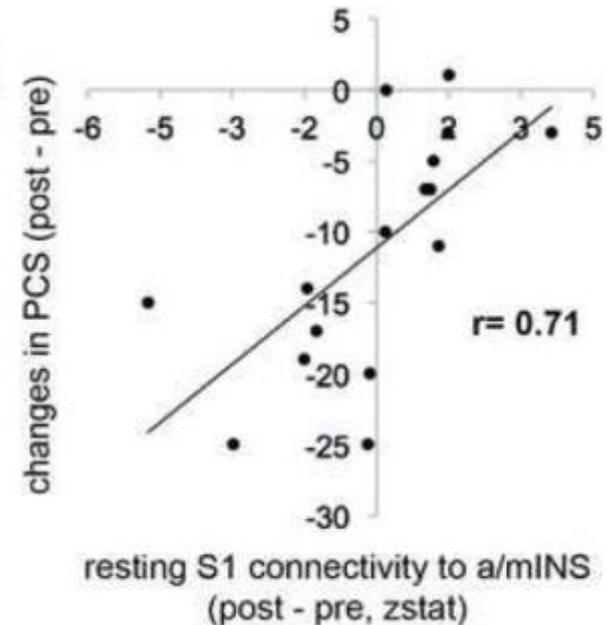
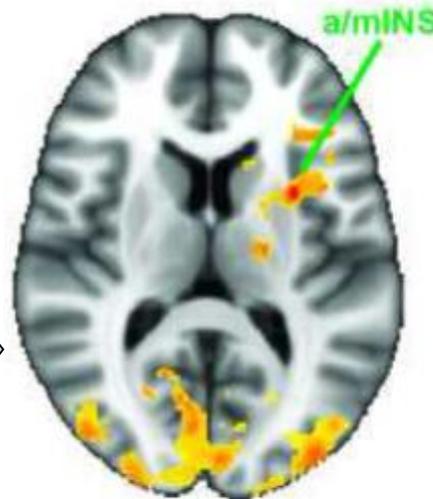
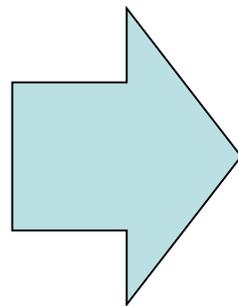


CBT reduces catastrophizing, which contributes to partial resolution of maladaptive connectivity (Lazaridou et al., 2017):

Patients in the CBT group show reductions in functional connectivity between S1 and anterior/medial insula at post-treatment:



And those who exhibit the largest reductions in catastrophizing also have the most substantial decreases in that connectivity . . .



CBT and Catastrophizing

Critical Review

How Can We Best Reduce Pain Catastrophizing in Adults With Chronic Noncancer Pain? A Systematic Review and Meta-Analysis

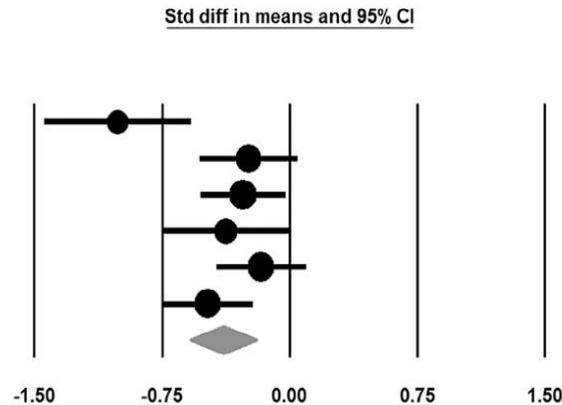


Robert Schütze,* Clare Rees,* Anne Smith,† Helen Slater,† Jared M. Campbell,‡,§ and Peter O'Sullivan†

Long-term effects at 6- to 12-months post-treatment:

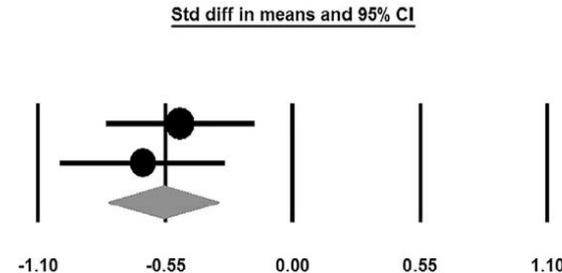
Cognitive Behavior Therapy

Study name	Std diff in means	p-Value	Sample size		
			Experimental	Control	Total
Alda 2011	-1.01	0.00	49	46	95
Amris 2014	-0.24	0.10	96	96	192
Broderick 2014	-0.28	0.03	129	127	256
Bromberg 2012	-0.38	0.05	46	74	120
Trudeau 2015	-0.17	0.20	113	115	228
Turner 2016	-0.48	0.00	112	113	225
	-0.39	0.00	545	571	1116



Multimodal treatment

Study name	Std diff in means	p-Value	Sample size		
			Experimental	Control	Total
Castel 2013	-0.49	0.00	81	74	155
Castel 2015	-0.65	0.00	69	61	130
	-0.56	0.00	150	135	285



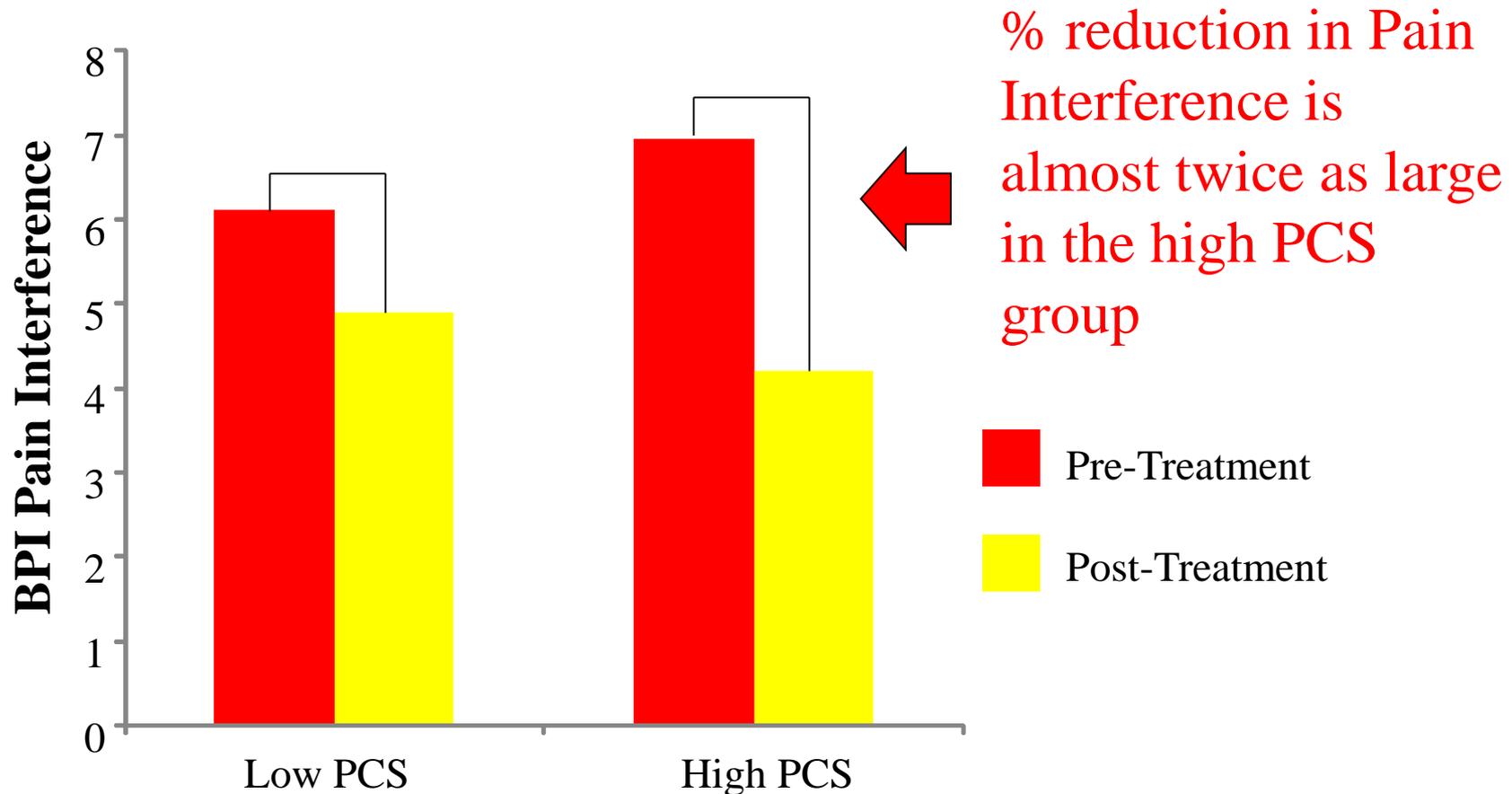
Favours Experimental

Favours Control

Significant, moderate-sized effects of CBT on reduction in catastrophizing. Some evidence that effects are largest in those with the highest baseline PCS scores.

CBT: Largest effects in High Catastrophizers

Ongoing BWH Trial



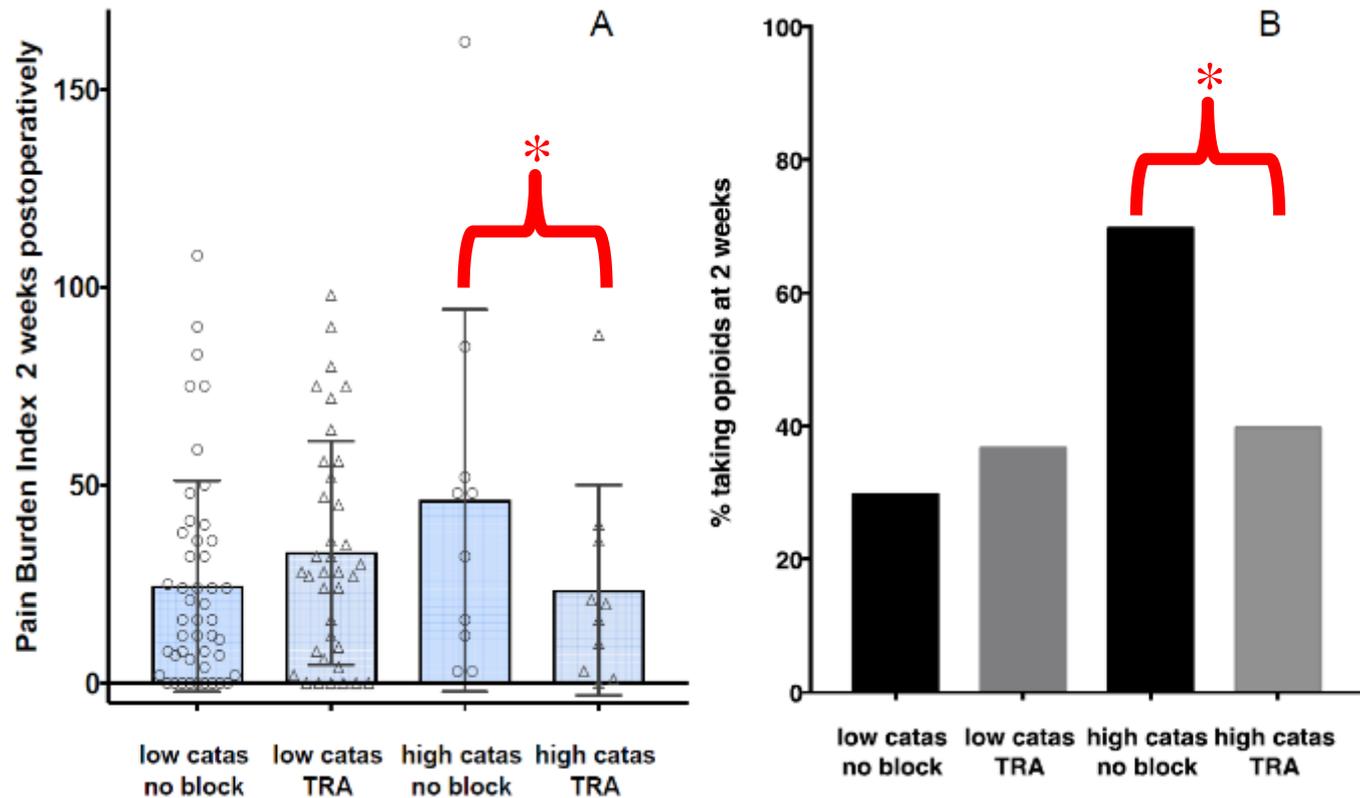
High catastrophizing may be a phenotype associated with enhanced benefit from certain interventions (and reduced benefit from others)

Significant interaction between catastrophizing and the application of regional anesthesia:

High catastrophizers get more benefit than low catastrophizers for pain and opioid outcomes at 2 weeks post-op:

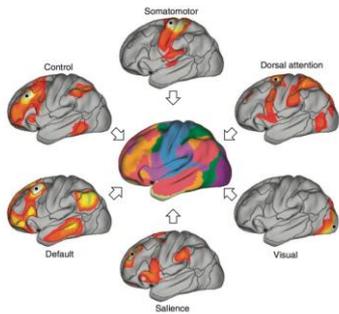
Schreiber et al., Under Review

Which Mastectomy Patients Benefit most from Regional Anesthesia? An Examination of the Influence of Catastrophizing



Conclusions

- Patients with complex, high-impact chronic pain have clusters of risk factors, many of them linked with cognitive/affective processes.
- Psychosocial phenotypes such as catastrophizing are reflective of brain/neurobiological phenotypes contributing to chronic pain.
- Non-pharmacologic interventions exert some of their benefits by acting on catastrophizing and related process variables.
- Eventually, such mechanism-based research should help us to move in the direction of “precision pain medicine” in which psychosocial and neurobiological phenotypes are employed to optimize treatment selection for individual patients.



Chronic Pain

