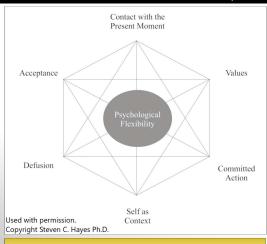


Acceptance and Commitment Therapy Delivered Via a Mobile Phone Messaging Robot Decreases Postoperative Opioid Utilization in Orthopedic Trauma Patients:

Randomized Control Trial

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INTRODUCTION AND PURPOSE

- Public health concerns regarding opioid medications persist and healthcare systems are currently seeking solutions to the ongoing epidemic
- Additional small daily amounts as low as 10 morphine milliequivalents (MME), and every week of continued opioid utilization represent an increased risk of eventual opioid misuse by patients.
- Patient reported outcomes (PROs) are important tools for determining the efficacy of healthcare treatments, assessing clinical research, and allow patients to quantify their health in a standardized fashion.
- The National Institutes of Health developed the Patient-Reported Outcome Measure Information System (PROMIS) tools to advance PROs by creating question banks that could be used for many major health issues.
- Acceptance and Commitment Therapy (ACT) is a contextual cognitive behavioral therapy that employs a pragmatic approach to help individuals decrease suffering and live according to self-identified personal values.
- The goal of ACT is to augment an individual's psychological flexibility, thus improving their life according to six core processes: Acceptance, Defusion, Contact with the Present Moment, Self-as-Context, Values, and Committed Action.
- Electronic communication methods, such as automated mobile phone messaging, for healthcare purposes have become more preferred by patients for delivering and receiving medical information.
- The purpose of this investigation was to evaluate the
 effectiveness of ACT delivered via an automated mobile
 messaging robot on (1) decreasing early postoperative
 opioid utilization and (2) pain-related patient reported
 outcomes (PROs) in the first two weeks following
 surgery for acute, traumatic orthopaedic injuries.

METHODS

- Adults presenting to a university hospital level 1 trauma center indicated for operative fixation of a traumatic upper or lower fracture were considered and approached for the study.
- Individuals who had a personal mobile phone, and were familiar with mobile messaging were eligible.
- Patients were randomized in a 1:1 ratio to either the intervention group, who received twice-daily mobile phone messages communicating an ACT-based intervention for two weeks after surgery, or the control group, who received no messages.
- The ACT-base intervention was developed in collaboration with a pain psychologist specializing in ACT for treatment of chronic pain.
- Baseline PROs were completed by all subjects. Two weeks after operative intervention, follow-up was obtained in the form of an opioid medication pill count and postoperative administration of PROs.
- Mean number of opioid tablets utilized by patients were calculated and compared between groups. Mean PRO scores were also compared between groups.

RESULTS

- 82 subjects enrolled in the study, 6 dropped from the study due to various issues, and the final study population was 76 patients with 38 in both the ACT and Control groups.
- Analysis of all patient demographics collected demonstrated no differences between study groups.
- Average subject age was 45.5 and 48.7 years old for the ACT and Control groups respectively.
- No differences between groups were found for injury type, disposition following discharge, method for reporting opioid medication consumption, preoperative opioid medication prescriptions, or preoperative PRO scores.
- The intervention group utilized an average of 26.1±21.4 opioid tablets while the control group utilized 41.1±22.0 tablets, resulting in 36.5% less tablets utilized by subjects receiving the mobile phone-based ACT intervention.
- Intervention group subjects reported a lower postoperative PROMIS Pain Intensity score of 45.9±7.2 compared to the control group's 49.7±8.8.

Opioid pain medication utilization by group during two-week study period											
	Opioid tablets dispensed			Opioid tablets consumed				Morphine milliequivalents consumed			
	ACT	Control		ACT	Control	%		ACT	Control	%	
	(N=38)	(N=38)	P-Value	(N=38)	(N=38)	Decrease	P-Value	(N=38)	(N=38)	Decrease	P-Value
Mean±	58.8±	61.6±		26.2±	41.1±			199.9±	307.0±		
SD	27.3	22.0		21.4	22.0			163.2	166.0		
Median	60.0	60.0	.62	21.0	43.5	36.5%	.004	157.5	307.5	34.9%	.006
(min-max)	(10-	(15-		(0-80)	(0-80)			(0-600)			
(IIIIII-IIIax)	146)	120)		(0-00)	(0-00)			(0-000)	(0-000)		

ACT: Acceptance and Commitment Therapy; Max: Maximum; Min: Minimum; SD: Standard deviation; % Decrease calculated by the formula $\left(\frac{A-B}{2}\right) \times 100 = Percent Decrease$

Mean PROMIS	score and change within	two-week study period by s	tudy aroun
IVICAITT TOWNS	Soore and onlinge within	Tivo week study period by s	tady group
PROMIS	Preoperative score	Postoperative score	Net so

PROMIS	Preopera	tive score	Posto	perative sco	ore	Net:	score chang	<u>e</u>
Instrument	ACT	Control	ACT	Control	P-Value	ACT	Control	P-Value
Pain Intensity 1Aa	5.4±2.9	6.2 ±2.6	3.4±2.2	4.1±2.4	.22	-2.0±2.9	-2.1±2.3	.79
Faill lillerisity IA-	(0-10)	(1-10)	(0-9)	(1-9)	.22	(-10-7)	(-9-2)	.19
Pain Intensity 3A	54.9±7.3	57.1±8.2	45.9±7.2	49.7±8.8	.04	-9.0±8.5	-7.4±7.7	.38
Faill lillerisity 3A	(36.3-71.8)	(40.2-71.8)	(30.7-64.1)	(30.7-67.4)	.04	(-25.5-10)	(-23.9-6.1)	.30
Pain Interference	63.6±11.4	66.1±8.4	56.6±9.4	60.6±8.2	.048	-7.1±13.7	-5.4±10.4	.55
8A	(40.7-77)	(40.7-77.0)	(40.7-72.0)	(40.7-77.0)	.040	(-36.3-24.8)	(-26.2-19.5)	.55
Emotional	56.5±11.4	56.5±9.2	51.5±10.4	52.3±10.6		-4.9±10.1	-4.2±9.4	
Distress-Anxiety	(37.1-80)		(37.1-75.4)		.76		(-20.3-16.5)	.74
Ιο Λ	(37.1-60)	(37.1-70.7)	(31.1-13.4)	(31.1-10.1)		(-33.7-12.3)	(-20.3-10.3)	1

Scores presented are raw numerical scores as no t-score conversion is available for the selected instrument. Data presented as mean t-score± standard deviation (range); ACT: Acceptance and Commitment Therapy; PROMIS: Patient-reported outcome measures information system

MOBILE PHONE MESSAGE EXAMPLE



CONCLUSIONS

- Delivering an ACT-based intervention via an automated mobile messaging robot in the acute postoperative period decreased opioid utilization in orthopaedic trauma patients in the first two weeks after their injury.
- Subjects in the ACT-based intervention group also reported lower pain intensity and pain interference after two weeks, though this likely did not represent a clinically important difference.
- Future studies may apply this intervention in other patient populations to assess its efficacy on a larger scale and may include assessment of pain and opioid use in a longer time frame after injury.

ACKNOWLEDGEMENTS

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