#### SCD ECHO Resources - 04 10 2019

### Safe Outpatient Management of Sickle Cell Disease Related Chronic Pain

THOKOZENI LIPATO, MD
ASSISTANT PROFESSOR, GENERAL INTERNAL MEDICINE
VIRGINIA COMMONWEALTH UNIVERSITY

#### References

- A systemic review of validated risk measurement tools. British Journal of Anaesthesia, 119 (6): 1092–109 (2017)
- Intensity of Chronic Pain The Wrong Metric? N Engl J Med 2015; 373:2098-2099
- Field JJ. Five lessons learned about long-term pain management in adults with adults sickle cell disease. Hematology Am Soc Hematol Educ Program. 2017 Dec 8;2017(1):406-411

#### Resources



### Sublingual Buprenarphine Naticeane for Chronic Pain in At-Riek Patienta: Development and Pilot Test of a Clinical Protocol

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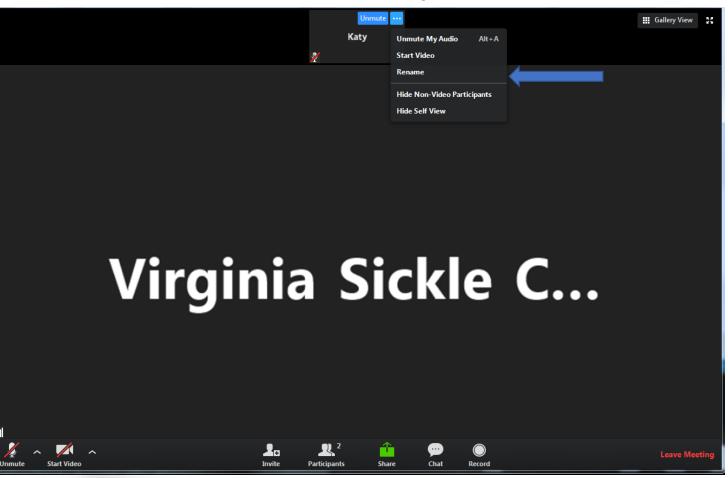
# Virginia Sickle Cell Disease ECHO\* Clinic

April 10<sup>th</sup>, 2019

\*ECHO: Extension of Community Healthcare Outcomes



### **Helpful Reminders**

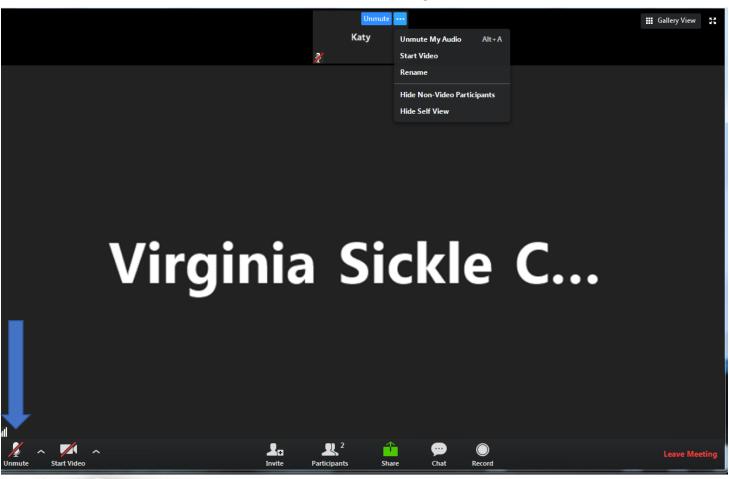




 Rename your Zoom screen, with your name and organization



### **Helpful Reminders**

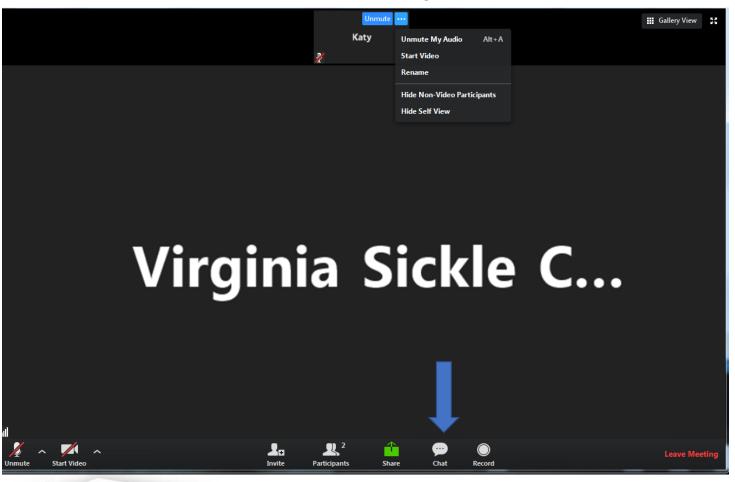




- You are all on mute please unmute to talk
- If joining by telephone audio only, \*6 to mute and unmute



### **Helpful Reminders**





- Please type your full name and organization into the chat box
- Use the chat function to speak with IT or ask questions



### VCU Sickle Cell Disease ECHO Clinics











- Monthly 2 hours tele-ECHO Clinics
- Every tele-ECHO clinic includes 2 case presentations and a didactic presentation
- Didactic presentations are developed and delivered by inter-professional experts in Sickle Cell Disease care and management

Website Link: <a href="http://vcuhealth.org/sicklecellecho">http://vcuhealth.org/sicklecellecho</a>





VCU Team			
Clinical Director	Wally R Smith, MD		
Administrative Medical Director ECHO Hub and Principal Investigator	Wally R Smith, MD		
Clinical Expert	India Y Sisler, MD Thokozeni Lipato, MD Jennifer Newlin, PA Mica Ferlis, NP		
Didactic Presentation	Thokozeni Lipato, MD		
Program Manager	Shirley Johnson, LSW		
IT Support	Daniel M Sop, M.Sc.Eng		
Administrative Assistant	Donna Casey		
Clinical Social Worker	Taylor Elliott, MSW		
Patient Navigators	Marla Brannon, BSW Stefani Vaughan-Sams		
Prior Authorization Specialist	Austin Hardy		

Virginia Commonwealth University







- Name
- Organization



### What to Expect

- . Case presentation #1 Elizabeth Yang, MD, PhD
  - i. Case summary
  - ii. Clarifying questions
  - iii. Recommendations
  - iv. Recap
- II. Didactic Presentation

Title: Safe Outpatient Management of Sickle

Cell Disease Related Chronic Pain Presenter: Thokozeni Lipato, MD

- III. Case presentation #2 Chelsea Rivenbark, NP
  - i. Case summary
  - ii. Clarifying questions
  - iii. Recommendations
  - iv. Recap
- IV. Closing and questions



Lets get started!

Case Presentation #1







### Case Presentation #1





- 12:50PM to 1:15pm [25 min]
  - Presentation: (5 min)
  - Case summary: Clinical Hub Lead(5 min)
  - Clarifying questions- Spokes (participants) 4 min:
  - Clarifying questions Hub (4 min):
  - Recommendations Spokes (participants) 2 min:
  - Recommendations Hub (2 min):
  - Recap Case /Recommendations- Hub (3 min):



Confidential

Participant ID 5 Page 1 of 5

#### Sickle Cell Disease Case Presentation Form

Virginia Sickle Cell Disease ECHO: De-Identified Case Study Submission

Thank you for submitting a case study!

Some benefits to submitting and presenting are...

-You will recieve valuable feedback regarding your case from our participating experts during the ECHO clinic

-A list of suggestions provided during the ECHO clinic will be sent to you as a reference after the clinic

-Your organization will be able to utilize suggestions and improve patient care!

- You will receive \$200 per case presented

DO NOT provide any patient specific information nor include any Protected Health Information.

Please complete the survey below.

Thank you!

Response was added on 04/08/2019 12:29pm		
Case Presenter First name	Elizabeth	
Case presenter last name	Yang	
Presenter Email:	eyang@psvcare.org	











# Didactic Presentation







# Safe Outpatient Management of Sickle Cell Disease Related Chronic Pain

THOKOZENI LIPATO, MD

ASSISTANT PROFESSOR, GENERAL INTERNAL MEDICINE

VIRGINIA COMMONWEALTH UNIVERSITY

# Objectives

- 1. Identify best opioid misuse/abuse screening tools to use for the management of chronic Sickle Cell Disease (SCD) pain
- 2. Determine whether or not chronic opioid therapy (COT) is successful in the management of chronic SCD pain
- 3. Discuss what can be done for patients with chronic SCD pain who are not stable on COT

# Assessment tools

#### Clear consensus

- Prior to initiating opioids for chronic pain assess for the risk of opioid abuse
- Reassess regularly for aberrant-drug related behavior after initiating chronic opioid therapy

No consensus on which assessment tool to use

## Table I: Opioid Therapy Risk-Assessment Screening Tools.

For the Opioid-Naïve	For the Opioid-Experienced			
<ul> <li>Self-Reported</li> <li>Drug Abuse Screening Test (DAST)</li> <li>Screener &amp; Opioid Assessment for Patients with Pain (SOAPP)</li> </ul>	Self-Reported     Current Opioid Misuse Measure (COMM)     Pain Medication Questionnaire (PMQ)     Prescription Drug Use Questionnaire (PDUQp)			
<ul> <li>Provider-Reported</li> <li>Opioid Risk Tool (ORT)</li> <li>Diagnosis, Intractability, Risk,</li> <li>&amp; Efficacy Score (DIRE)</li> </ul>	Provider-Reported     Prescription Drug Use Questionnaire     (PDUQ)			

### SOAPP-R

- 24 item tool
- Can be used to distinguish between low, moderate and high-risk groups

A systemic review of validated risk measurement tools. British Journal of Anaesthesia, 119 (6): 1092–109 (2017)

Table I: Opioid Therapy Risk-Assessment Screening Tools.

For the Opioid-Naïve	For the Opioid-Experienced
<ul> <li>Self-Reported</li> <li>Drug Abuse Screening Test (DAST)</li> <li>Screener &amp; Opioid Assessment for Patients with Pain (SOAPP)</li> </ul>	Self-Reported     Current Opioid Misuse Measure (COMM)     Pain Medication Questionnaire (PMQ)     Prescription Drug Use Questionnaire (PDUQp)
Provider-Reported  Opioid Risk Tool (ORT)  Diagnosis, Intractability, Risk,  Efficacy Score (DIRE)	Provider-Reported     Prescription Drug Use Questionnaire     (PDUQ)

### COMM

- 17-item tool
- Best tool to screen for current misuse of opioids.
- Use for regular monitoring

### PMQ

- 26-item tool
- Evaluates the risk of aberrant drug taking behavior
- Can be used to distinguish between low and high-risk groups.

A systemic review of validated risk measurement tools. British Journal of Anaesthesia, 119 (6): 1092–109 (2017)

Table I: Opioid Therapy Risk-Assessment Screening Tools.

For the Opioid-Naïve	For the Opioid-Experienced
Self-Reported     Drug Abuse Screening Test (DAST)     Screener & Opioid Assessment for Patients with Pain (SOAPP)	<ul> <li>Self-Reported</li> <li>Current Opioid Misuse Measure (COMM)</li> <li>Pain Medication Questionnaire (PMQ)</li> <li>Prescription Drug Use Questionnaire (PDUQp)</li> </ul>
Provider-Reported  Opioid Risk Tool (ORT)  Diagnosis, Intractability, Risk,  Efficacy Score (DIRE)	Provider-Reported     Prescription Drug Use Questionnaire     (PDUQ)

# When is COT successful?

Intensity of Chronic Pain — The Wrong Metric? N Engl J Med 2015; 373:2098-2099

Assessing pain-related functional impairment (Pre- and post- intervention)

Treat to a pain level and functional goal

### Assessing disease related impairment

PROMIS® (Patient-Reported Outcomes Measurement Information System)

- Person-centered measurement system
- Evaluates & monitors physical, mental, and social health.
- General population & with individuals living with chronic conditions

ASCQ-Me® (Adult Sickle Cell Quality of Life Measurement Information System)

- Patient-reported outcome measurement system
- Evaluates & monitors the physical, mental, and social well-being of adults with sickle cell disease (SCD).

# When is COT successful?

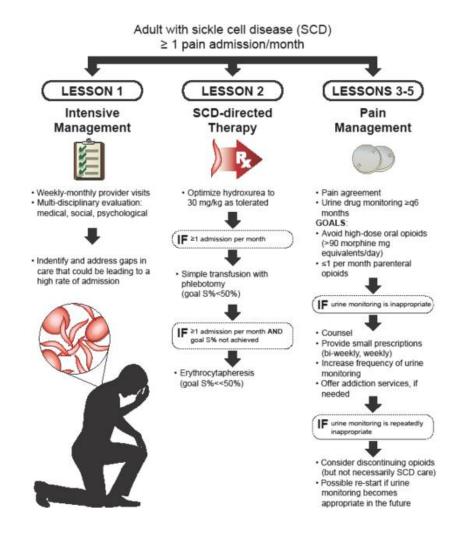
When it reduces emergency department visits & hospitalizations

# Caring for the *failing* patient

- Increase frequency of visits; shorten opioid prescription
- Close monitoring of opioids (i.e. pill counts)
- Increase frequency of UDS
- Behavioral health intervention (SW; psychologist)
- Patient navigators
- Opioid reduction, or tapered withdrawal
- Transition to safer pain regime (Tramadol; buprenorphine-naloxone)
- ED & Inpatient pain management plans tailored by outpatient team

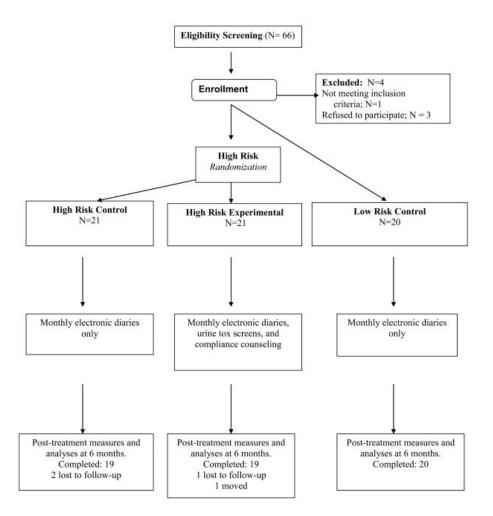
- 1. Focus resources on adults with high rates of utilization for pain
- Aggressively treat any underlying causes of pain, including SCD
- 3. Apply principles of chronic pain management to a clinic model
- Higher doses and larger quantities of oral opioids often do not help
- Parenteral opioids are not the optimal treatment of chronic pain

Field JJ. Five lessons learned about long-term pain management in adults with adults sickle cell disease. Hematology Am Soc Hematol Educ Program. 2017 Dec 8;2017(1):406-411



Substance misuse treatment for high-risk chronic pain patients on opioid therapy:
A randomized trial

PAIN 150 (2010) 390-400

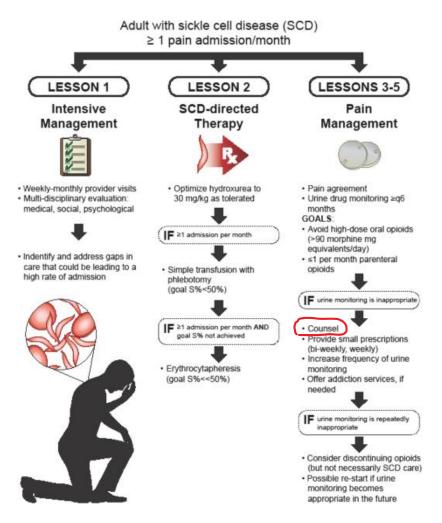


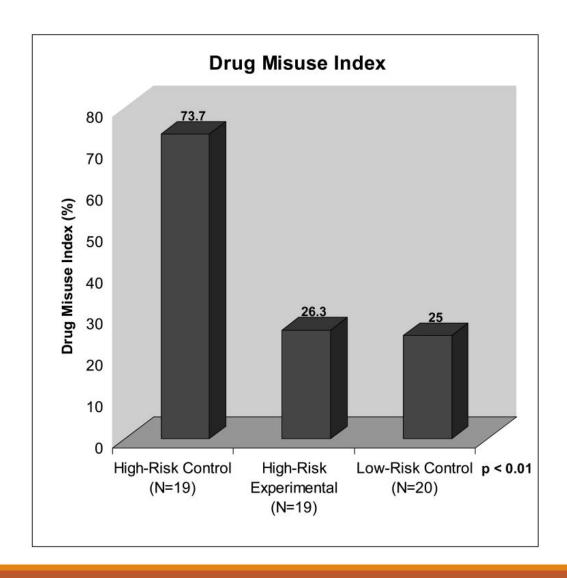
### High-risk experimental group

- 1. Completion of monthly electronic diaries
- Monthly urine screens for 6 months
- 3. Monthly completion of the Opioid Compliance Checklist
- 4. Monthly group education sessions (led by a psychiatrist [ADW] trained in pain and addiction medicine) with worksheet handouts on topics related to substance misuse
- 5. Participation in individual motivational compliance counseling (led by a clinical psychologist [RNJ] trained in pain and behavioral medicine) offering knowledge and training for substance misuse awareness and recovery

### **Opioid Compliance Checklist**

- 1. Taken your opioid medication other than the way it was prescribed?
- 2. Used more than one pharmacy to fill your opioid prescriptions?
- 3. Received opioid prescriptions from more than one provider?
- 4. Lost or misplaced your opioid medications?
- 5. Run out of your pain medication early?
- 6. Missed any scheduled medical appointments
- 7. Borrowed opioid medication from others?
- 8. Used any illegal or unauthorized substances?
- 9. Taken the highest possible degree of care of your prescription medication?
- 10. Taken any unauthorized substance that might be found in your urine
- 11. Been involved in any activity that may be dangerous to you or someone else if you felt drowsy or were not clear thinking?
- 12. Been completely honest about your personal drug use?







Published in final edited form as: J Opioid Manag. 2012; 8(6): 369–382. doi:10.5055/jom.2012.0137.

#### Sublingual Buprenorphine/Naloxone for Chronic Pain in At-Risk Patients: Development and Pilot Test of a Clinical Protocol

Andrew Rosenblum, PhD1, Ricardo A. Cruciani, MD, PhD23, Eric C Strain, MD4, Charles M. Cleland, PhD<sup>1,5</sup>, Herman Joseph, PhD<sup>1</sup>, Stephen Magura, PhD, CSW<sup>6</sup>, Lisa A Marsch, PhD<sup>1,7</sup>, Laura F McNicholas, MD, PhD<sup>8</sup>, Seddon R Savage, MD, MS<sup>9</sup>, Arun Sundaram, BA<sup>2</sup>, and Russell K. Portenov, MD2.3

Institute for Treatment and Services Research, National Development and Research Institutes,

\*Department of Pain Medicine and Palliative Care, Beth Israel Medical Center, New York, NY

<sup>3</sup>Departments of Neurology and Anesthesiology, Albert Einstein College of Medicine, Bronx, NY <sup>4</sup>Department of Psychiatry and Behavioral Sciences, Johns Hopkins University School of

<sup>5</sup>College of Nursing, New York University, New York, NY

<sup>6</sup>Evaluation Center, Western Michigan University, Kalamazoo, MI

<sup>7</sup>Center for Technology and Behavioral Health, Dartmouth Psychiatric Research Center, Department of Psychiatry, Dartmouth College, Lebanon, New Hampshire

<sup>8</sup>Department of Psychiatry, University of Pennsylvania, Philadelphia, PA

<sup>9</sup>Center on Addiction Recovery and Education, Dartmouth Medical School Hanover, New Hampshire

#### Abstract

Medicine, Baltimore, MD

Objective-Sublingual buprenorphine/naloxone (Bup/Nx) is approved for addiction treatment and may be useful for pain management, particularly in opioid-treated pain patients with nonadherence behaviors. The transition of opioid-treated pain patients to buprenorphine carries the risk of precipitated withdrawal and increased pain. This study convened pain and addiction specialists to develop and pilot a clinical protocol for safe transitioning to Bup/Nx.

Design-The protocol was revised three times based on outside expert review and pilot study bservations. The pilot was conducted with a prospective cohort of 12 patients with moderate to severe chronic pain, who were receiving long-term opioid therapy with any full µ-agonist drug, and had exhibited one or more aberrant drug-related behaviors. Patients were followed 6 months with the expectation that they would experience few adverse events and report lower

Results-The three patients on the highest baseline opioid dose (equivalent to 303-450 mg of oral morphine) and the three on the lowest doses ( 20 mg) had early adverse events (AEs) when switched to Bup/Nx and did not complete the trial. Of the remaining six, one withdrew due to AEs; one responded well, then withdrew; and four completed a three-month trial. A mixed effects





Pain Medicine 2014; 15: 2087-2094 Wiley Periodicals, Inc.

#### Conversion from High-Dose Full-Opioid Agonists to Sublingual Buprenorphine Reduces Pain Scores and Improves Quality of Life for Chronic Pain Patients

Danielle Daitch, MD,\* Jonathan Daitch, MD,\* S Daniel Novinson, MPH,\* Michael Frey, MD,\* S Carol Mitnick, ARNP,\* S and Joseph Pergolizzi, Jr MD\*\*, 11.23

\*Jefferson Medical College, Philadelphia. Pennsylvania; <sup>†</sup>Advanced Pain Management and Spine Specialists, Fort Myers, Florida; \*Park Center for Procedures ASC, Fort Myers, Florida; <sup>5</sup>University of California, San Francisco, California; <sup>5</sup>Virginia Commonwealth University, Richmond, Virginia. "Department of Anesthesiology, Georgetown University Medical School, Washington, DC: <sup>11</sup>Department of Medicine, Johns Hopkins University School of Medicine, Baltimore, Maryland: <sup>14</sup>Department of Pharmacology, Temple University School of Medicine, Philadelphia, Pennsylvania, USA

Reprint requests to: Jonathan Daltch, MD, Advanced Pain Management and Spine Specialists, 8255 College Parkway, Suite 200, Fort Myers, FL 33919. USA. Tel: 239-437-8000; Fax: (239) 437-8002; E-mail: idaltch@ampss.net.

Disclosure: Dr. Jonathan Daitch is a speaker for

Objective. This study aims to determine the effectiveness of converting patients from high doses of full-opioid agonists to sublingual (SL)

Design. An observational report of outcomes

Setting. An interventional pain management practice setting in the United States.

Subjects. Thirty-five chronic pain patients (age 24objects. Intry-we crrome pain pasteria (age ze-f6) were previously treated with high-dose opioid-agonist drugs and converted to St. buprenorphine. Patients' daily morphine equivalents ranged from 200 mg to 1,370 mg preconversion, with a mean

Methods. A retrospective chart analysis examined numerical pain levels and quality of life scores before and 2 months after conversion to SL

Results. After continuation of SL buprenorphine therapy for 2 months, the mean pain score decreased from 7.2 to 3.5 (% c.001), with 3 of the form of t

Conclusion. Average pain scores decreased from 7.2 to 3.5, and quality of life scores increased from 6.1 to 7.1 for 35 patients converted from high-dose full-opioid agonists to SL buprenorphine therapy for more than 60 days. Clinicians phine therapy for more than 60 days. Clinicians should consider bupenorphine St. conversion for all patients on high-dose opioids, particularly patients with severe pain (7-10) unrelieved by their current opioid regimen or patients for whom the clinician does not feel comfortable prescribing high-dose opioids.

Key Words, Buprenorphine; Sublingual Buprenor-phine; Opioid Conversion; Opioid-Induced Hyper-algesia; Analgesia; Opioid Tolerance

Analgesics that act at several sites along the pein path-way to diminish pain, opioids have been used to treat pain for thousands of years [1-3]. Today, some of the most commonly prescribed medications for severe pain



PAIN\* 154 (2013) 1442-1448



Buprenorphine/naloxone as a promising therapeutic option for opioid abusing patients with chronic pain: Reduction of pain, opioid withdrawal symptoms, and abuse liability of oral oxycodone

Perrine Roux a,b,c,d,\*, Maria A. Sullivan a, Julien Cohen b,c, Lionel Fugon b,c, Jermaine D. Jones a, Suzanne K. Vosburg a, Ziva D. Cooper a, Jeanne M. Manubay a, Shanthi Mogali a, Sandra D. Comer a

\*Division on Substance Abuse, New York State Psychiatric Institute, Department of Psychiatry, and College of Physicians and Surgeons, Columbia University, 1051 Riverside Dr., Unit 120, New York, NY 10032, USA

Suppose the compating interests that may be released to content are disclosed at the end of this article

ARTICLE INFO

Article history: Received 20 November 2012 Received in revised form 12 April 2013 Accepted 1 May 2013

ABSTRACT

A B S T R A C T

For studies have examined aboue of prescription opioids among individuals with chronic pain under buyersorphine/haborone BhopNo; maintenance. The current 7-week inpatient study assessed or all capacitosis of the property of the principals (in ~ 25) were transitioned from their preadmission prescribed opioid to BupPix. All of the participants (in ~ 25) were transitioned from their preadmission prescribed opioid to BupPix. All of the principants (in ~ 26) were transitioned property of the principants could self-administer oxyocodone orally (in 10, 20, 40 or 60 mg prescription opioids) or receive money during laberatory sessions. Dung choice (precentage) was the prinney dependent variable. Subjective ratings of clinical jaint and withdrawal symptoms also to place to conjust the principant sweet to place the conjustic regression analysis individually allowed to the principants were to place to conjustic regression analysis individually reduced while participants were observed between the active oxyocodone doses and placebo under each BupPix maintenance dose. However, Latexas associated with oxyocodone doses and placebo under each BupPix maintenance dose. However, Latexas associated with oxyocodone preference were forewer BupPix maintenance dose. However, Latexas associated with oxyocodone preference were over the BupPix maintenance dose. However, Latexas associated with oxyocodone preference were the overe BupPix maintenance dose new supplemental oxyocodone use. Importantly, adequate management of pain and withdrawal symptoms by BupPix may reduce oxyocodone preference in this population.

#### 1. Introduction

In the United States, the problem of nonmedical use of prescription opioids has emerged as a major public health issue [1]. Other countries, such as Australia, New Zealand [12] and Canada [33], also are concerned about the phenomenon of prescription opioid abuse. In the United States, oxycodone and hydrocodone are among the most commonly prescribed or regularly used opioids, as well as the most commonly diverted prescription opioids anal-gesics [5,25]. These data indicate that prescription opioids abuse has steadily increased among heroin and recreational polydrug users since 2000 [5]. An additional concern related to the increased

use of prescription opioids is opioid overdose, which increased in the United States from the mid-1990s to the present time [10,11] and recently became a leading cause of accidental death in the United States [4]. Thus, the risks of prescription opioids abuse and overdose make physicians reluctant to prescribe prescription opioids in general, and access to adequate pain management in opnois in general, and access or acceptance pain inflangement in drug users in particular is becoming increasingly difficult [20,39]. In those patients who are prescribed prescription opioids for pain relief, misuse may occur in pain patients with no history of opioid abuse who become dependent on the medications for their reinforcing properties, whether good drug effects or relief of anxiety or mood symptoms, or missue may occur in drug-seeking individ-uals with preexisting opioid abuse histories. Thus, balancing the need for effective pain relief and reducing the risks of opioid abuse and overdose remains a challenge for public health policy [7].

conding Author: Andrew Rosenblum, Institute for Treatment and Services Research (ITSR), National Development and th Institutes (NDRI), 71 West 23<sup>th</sup> Street: 8<sup>th</sup> Floor, New York, NY 10010. Phone: (212) 844-4528, Fax: (917) 438-0894,

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\*Université Aix Marseille, IRD, UMR-5912, Marseille, France
\*4085 PACA Déservatoire Régional de la Santé Provence Alpes Côte d'Azur, Marseille, France

<sup>\*</sup> Corresponding author at: INSERM U912 (SESSTIM), Marseille, France. Tel.: +33 4

E-mail address: perrine.roux@inserm.fr (P. Roux).

<sup>0304-3959/</sup>S36.00 Published by Elsevier B.V. on behalf of International Association for the Study of Pain. http://dx.doi.org/10.1016/j.pain.2013.05.004

### Case Presentation #2





- 1:40- 2:00pm [25 min]
  - Presentation: (5 min)
  - Case summary: Clinical Hub Lead(5 min)
  - Clarifying questions- Spokes (participants) 4 min:
  - Clarifying questions Hub (4 min):
  - Recommendations Spokes (participants) 2 min:
  - Recommendations Hub (2 min):
  - Recap Case /Recommendations- Hub (3 min):



Confidential

Participant ID 6 Page 1 of 6

#### Sickle Cell Disease Case Presentation Form

Virginia Sickle Cell Disease ECHO: De-Identified Case Study Submission

Thank you for submitting a case study!

Some benefits to submitting and presenting are...

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- You will receive \$200 per case presented

DO NOT provide any patient specific information nor include any Protected Health Information.

Please complete the survey below.

Thank you!

Response was added on 04/08/2019 5:09pm.	
Case Presenter First name	Chelsea
Case presenter last name	Rivenbark
Presenter Email:	chelsea.rivenbark@vidanthealth.com





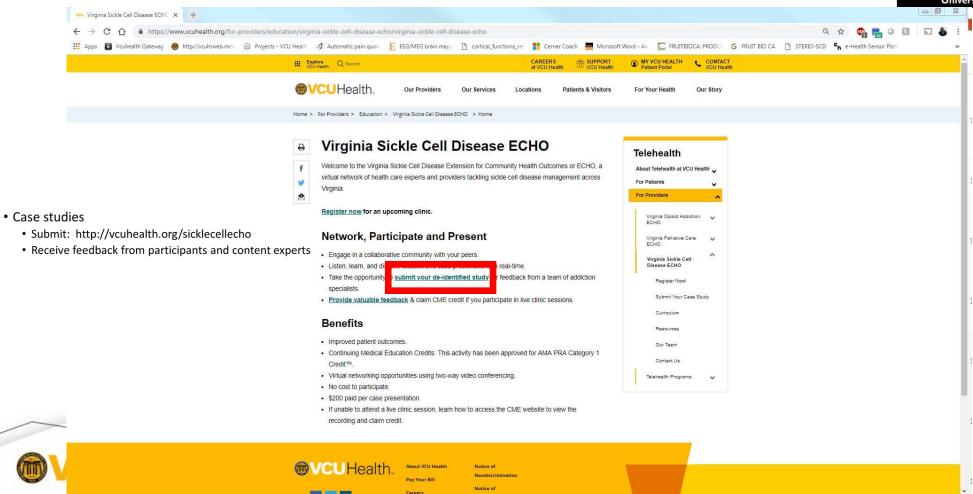






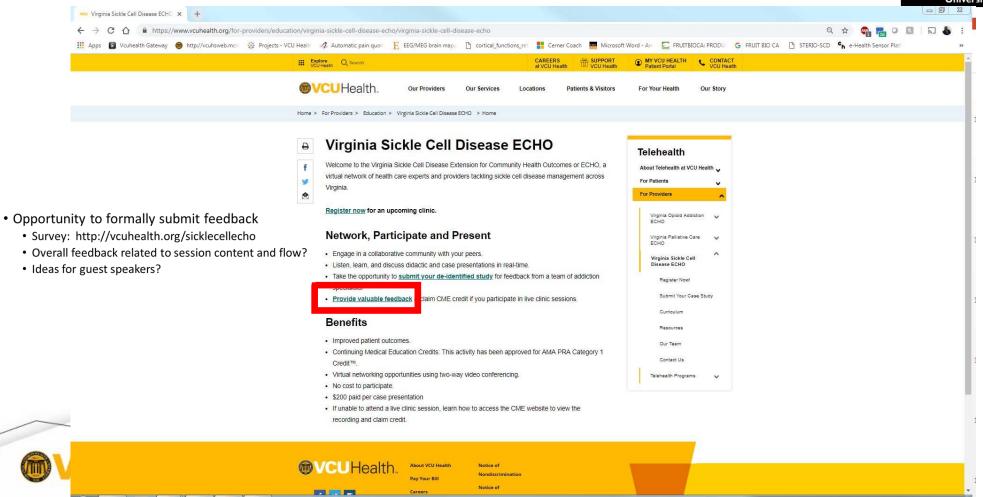
### Case Studies





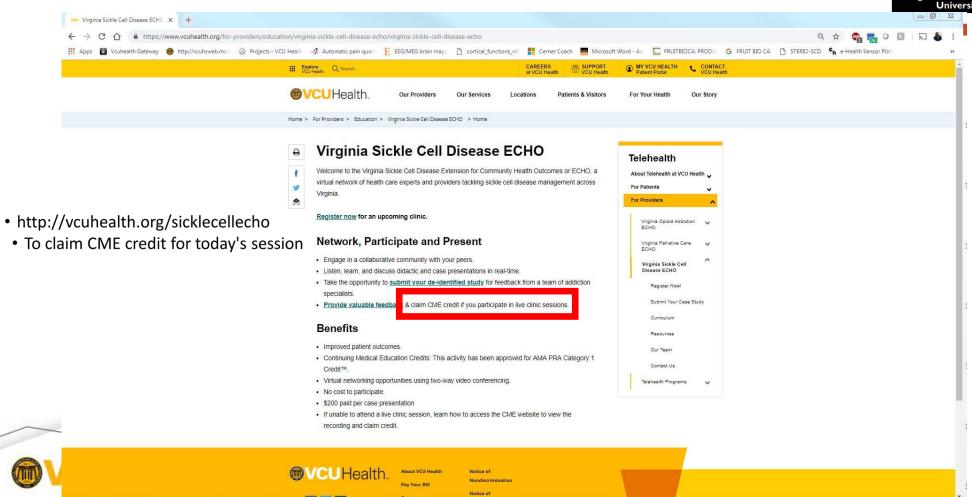
### Feedback





### Claim Your CME's





### Access Your Evaluation and Claim Your CME



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	the survey below.								
Thank you!									
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Doyo	u intend to make changes based o	n this presentation	,?						





### THANK YOU!

