



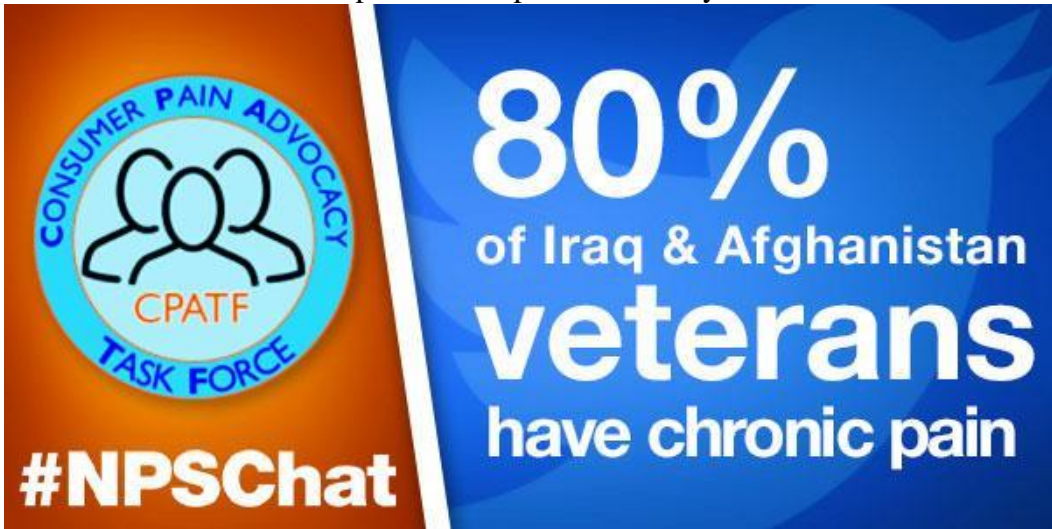
	Welcome
@powerofpain	Thank you for joining the Consumer Pain Advocacy Task Force #CPATF for today's #TwitterChat on #NationalPainStrategy #PainMonth15. Please use to stay involved in the conversation - the #CPATF chat on #NPS is being hosted by @powerofpain
@powerofpain	Please welcome our guest tweeters @LynnRWebsterMD @BobTwillman & @CPRAlliance_org talking all things National Pain Strategy for #CPATF
@powerofpain	@BobTwillman is Executive Dir of the American Academy of Pain Management, overseeing fed & state pain policy developments
@powerofpain	@LynnRWebsterMD is VP of Scientific Affairs, PRA HS Fmr Pres, American Academy of Pain Medicine Author of The Painful Truth
@powerofpain	@CPRAlliance_org (Christin Veasley) is a founding member of the Chronic Pain Research Alliance, whose mission is to advance scientific research on prevalent, neglected and poorly understood pain disorders that frequently co-occur and disproportionately affect women
@powerofpain	Let's get started #CPATF #PainMonth15
@BobTwillman	@powerofpain Good evening, everyone! Let's get started talking about the NPS!
@rsdcprfire	Thank you for joining the Consumer Pain Advocacy Task Force #CPATF for today's #TwitterChat on #NationalPainStrategy #PainMonth15
@AmerGastroAssn	The Gut Check: Know Your Medicine campaign (www.gutcheckfacts.org) is excited to join the about chronic pain!
	Q1. What is the National Pain Strategy report?
@BobTwillman	A1. The NPS was mandated as part of the Affordable Care Act (Obamacare) legislation. National Pain Strategy is an effort to outline a strategic approach to chronic #pain in the USA. The government's approach to pain research, treatment has historically been disorganized. We hope NPS fixes that. A1. The Department of Health and Human Services is in charge of developing the report and disseminating it.
@CPRAlliance_org	A1. It's the country's first strategic plan to advance pain care, research and education! Six federal agencies worked with nearly 100 volunteers to develop it.
@mkwinnette	A1. It contains short- and long-term goals, and identifies key partners and tools to measure progress in six areas.
@mkwinnette	A1. http://consumerpainadvocacy.org/nationalpainstrategy

	Q2. What did you like in the NPS draft report?
@BobTwillman	A2. The NPS draft report is comprehensive, covering a number of important areas related to good pain care.
@BobTwillman	A2. Draft report was developed by nearly 80 experts in all aspects of pain management. Dr. Mackey was co-chair of the panel developing the report.
@CPRAlliance_org	A2. The NPS report continues to focus on biopsychosocial model of pain and its treatment--vital if we are to progress. The NPS covers all areas that are needed to improve the lives of people with pain - pain care, education & awareness, research!
@PAINSPROJECT	A2. Sean Mackey gave a talk about the process for developing the report which is very enlightening. http://ow.ly/SOYf8
@RubysFibroDiary	Has the NPS been published since the public comment period ended?
@BobTwillman	@RubysFibroDiary The final report, after comment period, has not yet been published. No date has been set yet.
@BobTwillman	@ourCPC Absolutely--it won't be easy on either count, so we need to be strong advocates for implementation.
@powerofpain	<p>1 in 3 American adults live with chronic pain. #CPATF</p> 

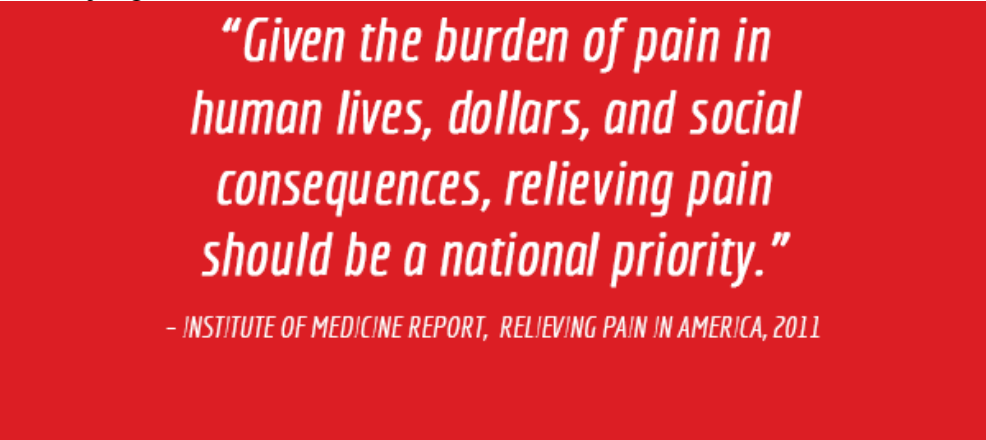

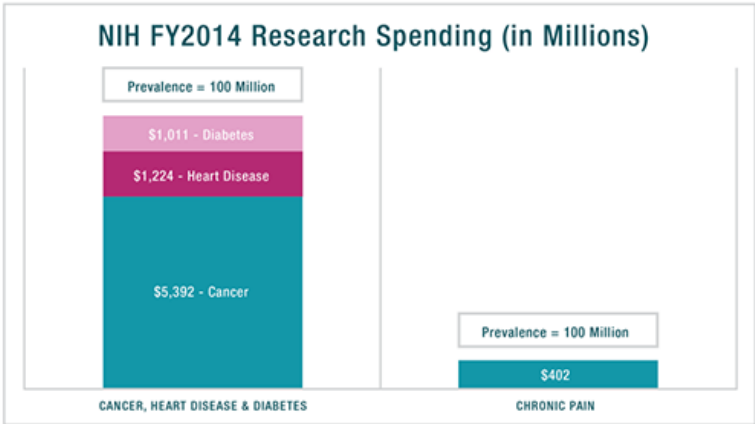
	Q3. Why is the NPS needed?																						
@CreakyJoints	A3: The NPS is needed because too many people live with #chronicpain. We can and will do better.																						
@mkwinnette	A3.Chronic pain is both an under-recognized and under-resourced public crisis with devastating personal and economic impact.																						
@mkwinnette	A3. Chronic pain is not one specialty, it takes a team!																						
@BobTwillman	A3. Pain research and treatment involves MANY different groups, in and out of government, so we need coordination. The NPS takes a comprehensive view of the needs and recommends ways to meet them.																						
@mkwinnette	Chronic pain is a real and complex disease that may exist by itself or be linked with other medical conditions.																						
@RubysFibroDiary	Will the Federal Pain Research Policy work closely with the NPS recommendations?																						
@BobTwillman	@RubysFibroDiary HHS received thousands of comments, so is working to respond and integrate them into the final report.																						
@ourCPC	@BobTwillman Has #NPS looked at companies or organizations that are already working to implement																						
@BobTwillman	@ourCPC I know HHS is interested in public-private partnerships, but I don't know how much outreach they have done.																						
@MIFibro	What steps can be taken to get funding & implementation for NPS?																						
@BobTwillman	@MIFibro We all need to be very vocal to ensure this isn't just a report that gathers dust on a shelf. This is going to take a LOT of work, and advocacy with Congress and other federal agencies like HHS.																						
@BobTwillman	@RubysFibroDiary Both are coming from the same group (IPRCC), so one would hope they are coordinated.																						
@RsdJulian	I have read the report but would like to know if it is having any affect with both our federal and state governments																						
@BobTwillman	@RsdJulian The final report still hasn't been published, so there hasn't been huge impact so far.																						
@BobTwillman	@RsdJulian However, there is an opportunity to start education policymakers NOW, rather than waiting for the final report.																						
@lisadbudzinski	http://www.iprcc.nih.gov/National_Pain_Strategy/NPS_Main.htm for the @centralpain																						
@lisadbudzinski	@BobTwillman @RubysFibroDiary and can be found here: http://iprcc.nih.gov/FPRS/FPRS.htm																						
@CPRAlliance_org	<p>Only 18% think of Americans think that pain is a major public health issue.</p> <table border="1"> <caption>Chronic Pain Ranks Below Many Other Conditions as Major Health Problem</caption> <thead> <tr> <th>Condition</th> <th>Percentage</th> </tr> </thead> <tbody> <tr> <td>Cancer</td> <td>59%</td> </tr> <tr> <td>Heart disease</td> <td>52%</td> </tr> <tr> <td>Diabetes</td> <td>52%</td> </tr> <tr> <td>Drug addiction</td> <td>47%</td> </tr> <tr> <td>Depression</td> <td>42%</td> </tr> <tr> <td>Alcoholism</td> <td>37%</td> </tr> <tr> <td>Alzheimer's disease</td> <td>34%</td> </tr> <tr> <td>Chronic pain</td> <td>18%</td> </tr> <tr> <td>Parkinson's disease</td> <td>15%</td> </tr> <tr> <td>Not sure</td> <td>15%</td> </tr> </tbody> </table> <p>Source: A ResearchAmerica poll of U.S. adults conducted in partnership with Zogby Analytics in March 2013.</p>	Condition	Percentage	Cancer	59%	Heart disease	52%	Diabetes	52%	Drug addiction	47%	Depression	42%	Alcoholism	37%	Alzheimer's disease	34%	Chronic pain	18%	Parkinson's disease	15%	Not sure	15%
Condition	Percentage																						
Cancer	59%																						
Heart disease	52%																						
Diabetes	52%																						
Drug addiction	47%																						
Depression	42%																						
Alcoholism	37%																						
Alzheimer's disease	34%																						
Chronic pain	18%																						
Parkinson's disease	15%																						
Not sure	15%																						

@SabrinaNumann	Chronic pain is a real and complex disease that may exist by itself or be linked with other medical conditions.
@exoeagle	What r u doing to make Dr's compassionate 2 the needs of individuals w chronic pain & not think them 2 be drug seekers
@BobTwillman	@exoeagle @powerofpain Achieving this requires improved education, which is one of the key components of the NPS.
@ourCPC	@exoeagle That is a hard one. @powerofpain is an amazing organization that supports #patients and #meded
@RubysFibroDiary	Have large employers shown interest in the NPS?
@CPRAlliance_org	@RubysFibroDiary The NPS hasn't been released yet, but we hope they will once it publishes! It costs America \$500 billion a year!
@ourCPC	@RubysFibroDiary yes :) we also talk to other companies that want to help the #pain community as well
@SabrinaNumann	Effective ...care requires access to... range of treatment options, including biomedical, behavioral health... complimentary treatment
@SabrinaNumann	Chronic pain is both an under-recognized and under-resourced public health crisis with devastating personal and economic impact
@CPRAlliance_org	Our government only invests a nickel per person in pain research. We need a strategic plan! <p style="text-align: center;">Federal government's investment in research</p>  <p style="text-align: center;">cpralliance.org</p>



	Q4. What will happen after the NPS is released?
@BobTwillman	A4. We also will need to ensure that SOMEONE takes primary responsibility for taking the ball and running with it. That's entirely up to us. We need to ensure that every member of Congress knows about it, for starters.
@LynnRWebsterMD	A4. We need all of the Gov't agency to be on board as well
@MIFibro	@BobTwillman Thanks and agreed. Wondering what a grassroots call to action could be for folks willing to do a LOT of work?
@BobTwillman	@MIFibro #CPATF is organized to address that very issue. Getting connected with them will enable us to get those messages out.
@CPRAlliance_org	@MIFibro @BobTwillman CPATF is organizing a large Capitol Hill Day. Please join us! www.ConsumerPainAdvocacy.org
@CPRAlliance_org	@MIFibro It's going to take a large advocacy effort, which is why the Consumer Pain Advocacy Task Force exists. Please join us!
@CPRAlliance_org	@MIFibro www.ConsumerPainAdvocacy.org
@CPRAlliance_org	@MIFibro @BobTwillman Yes! Please visit the CPATF's web site: www.ConsumerPainAdvocacy.org
@MIFibro	@BobTwillman are there established talking points for "voices" to speak in unison on NPS?
@ourCPC	@MIFibro The ball started rolling a while ago ;)
@gnsalminen	Everyone deserves the excellent caliber of care!! How can we get this to all who need it??
@BobTwillman	@gnsalminen We want to let people know it is coming, and to prime them for action as soon as the final report is released.
@gnsalminen	I hear about so many patients NOT being listened to by their doctors.
@mkwinnette	Having to wait 6 months for a Geneticist to be able to tell my children what all they may inherent is wrong.
@LynnRWebsterMD	Every person in pain must be made aware of the report and ask their providers and Congressman to support it.
@LynnRWebsterMD	I am calling for a national movement. Need 100,000 letters sent to Congressman within a year supporting NPS. Are you on board?
@lisadbudzinski	@ThePainComm @CPRAlliance_org Here's a link that breaks down some of the research: http://paindatabase.nih.gov/content/iprp-ontology
@lisadbudzinski	The link to the draft that is being discussed - http://iprcc.nih.gov/docs/DraftHHSNationalPainStrategy.pdf @centralpain
@exoeagle	@BobTwillman @powerofpain how will you improve education to practicing physicians and new physician?
@BobTwillman	@exoeagle @powerofpain We need increased inter-professional education in schools focused on multimodal multidisciplinary treatment.
@BobTwillman	@exoeagle @powerofpain Also will take LOTS of continuing education for clinicians already in practice.
@OneDayHeather	Is the NPS promoting advocacy for patients whose ins. company deny coverage 4 treatments? For ex: ketamine infusions @powerofpain
@CPRAlliance_org	@OneDayHeather @powerofpain Yes! There is an entire section with short, medium & long-term goals focused on reimbursement!
@MissionDriven4	For an individual w/ long term chronic pain who wants to be an advocate - what's the best way to get connected, take action?
@CPRAlliance_org	@MissionDriven4 Please contact us through our web site: www.ConsumerPainAdvocacy.org
@gnsalminen	We need doctors to follow Standards of Care!

@gnsalminen	A lot of them do not follow it!
@gnsalminen	What do the letters need to say?
@CPRAlliance_org	@gnsalminen Agreed! This is why the CPATF is calling for complete implementation of the NPS! We need everyone's support!
@ourCPC	@exoeagle At the @AmericanPainSoc annual meeting, #doctors said to take it to medical school. Change the teaching
@melissarvh	What specific actions might we take to ensure that happens?
@PAINSPROJECT	It's important for everyone to read the NPS Report and raise awareness so that change can start to happen
@RubysFibroDiary	Any idea when the CPATF Capitol Hill day will be?
@SPPAN1	@RubysFibroDiary Our #CPATF hill day will most likely be in early Spring 2016. More info will be on www.ConsumerPainAdvocacy.org soon
@cat_fish_dinner	I'm concerned a population based health strategy may inhibit personalized care. pain levels n impairment vary greatly
@CPRAlliance_org	@cat_fish_dinner The NPS very much addresses that individualized pain care is needed for and deserved by all!
@BobTwillman	@cat_fish_dinner Challenging, because true change only happens one patient/clinician interaction at a time.
@SabrinaNumann	Denying...care to people with chronic pain is unethical... can lead to unnecessary suffering, depression, disability... & suicide
@powerofpain	80% of Veterans have developed chronic pain in last 10 years. #CPATF 


	Q5. Why should we all care about the National Pain Strategy?
@BobTwillman	A5. If we're going to make progress it has to be guided by a strategic plan. We can't continue to do this haphazardly. Continuing to do what we have done will continue to give us the results we have. If we want change, we need a plan!
@LynnRWebsterMD	A5. Look we must insist on better payer coverage for integrative care. NPS calls for this. Payers must be part of the solution. Actually we start now to ask when the report will be made public. Call your congressman or woman tomorrow.
@barbyingle	A5. Because we need better and quicker access to care. This is one tool that can help
@BobTwillman	@LynnRWebsterMD Includes Medicare and Medicaid, VA/DOD, private payers-- everyone has to be part of it.
@gnsalminen	We should ALL care! Because what affects one of us, affects ALL of us!! Addiction and Opiate dependency ARE SEPARATE!! 3-5% of people with chronic pain become addicts Those of us on opiates are not the problem!
@gnsalminen	Bob Twillman, who are payers??
@BobTwillman	@gnsalminen We also need to remember that employers have a HUGE stake in making this work. They are payers, too.
@CPRAlliance_org	@gnsalminen The CPATF will have a letter template available on our web site to use and distribute widely. http://www.ConsumerPainAdvocacy.org
@FibroAndPain	How can we separate the disease of addiction from the disease of chronic pain?
@FibroAndPain	How can access to care for chronic pain issues be tied to Medicare essential health benefits?
@BobTwillman	@FibroAndPain Challenging, but the bottom line is that both groups are suffering.
@US_Pain	@FibroAndPain One way would be for pts to share their story, inform and educate lawmakers/society that the two are different conversations
@MIFibro	@ourCPC I was thrilled to see the ball rolling! Just querying this amazing group on next steps & how to advise public to help :)
@CPRAlliance_org	@MIFibro @ourCPC Please continue to revisit our web site for updates and calls to action. www.ConsumerPainAdvocacy.org
@CPRAlliance_org	@MIFibro Of primary importance is getting the federal government to allocate the funding we need for COMPLETE implementation!
@MIFibro	@CPRAlliance_org thanks for the invite and will do!
@crystalgrieves	@powerofpain @LynnRWebsterMD Will there be base template for these letters that could be drafted to use to have others send as well?
@crystalgrieves	@powerofpain imagine the possibility of lower costs if everyone is on the same page, testing for the same things over and over!
@barbyingle	@crystalgrieves Yes, there will be a template for the letters. Check back for updates at www.ConsumerPainAdvocacy.org
@exoeagle	@BobTwillman is this your model for current physicians?
@mkwinnette	There is not set standard. Different diagnosis from different doctors. Is our list really that long?
@BobTwillman	@mkwinnette @LynnRWebsterMD Prayers won't hurt, either! :-)
@mkwinnette	Chronic pain is the addiction! RT @FibroAndPain How can we separate the disease of addiction from the disease of chronic pain?

<p>@lisadbudzinski</p>	<p>@powerofpain Link to a description for the #NationalPainStrategy http://consumerpainadvocacy.org/take-action @centralpain</p>
<p>@PAINSPROJECT</p>	<p>We have to all stand together to raise awareness and change the way pain is perceived, judged and treated.</p> 
<p>@AmerGastroAssn</p>	<p>More Americans have chronic pain than the # of people w/ diabetes, coronary heart disease & cancer- combined</p> 
<p>@CPRAlliance_org</p>	<p>@AmerGastroAssn Yes! And the government spends 95% less on pain research than it does on these conditions!</p>  <p style="text-align: right;">cpralliance.org</p>

	Q6. Who is responsible for implementing the NPS after it's released?
@BobTwillman	A6. That's the problem--no one is identified as responsible. That's why we need to advocate for a point person.
@CPRAlliance_org	A6. Right now, no one, which is why we need everyone's support. We have to make sure the NPS doesn't sit on the shelf!
@CPRAlliance_org	A6. Right now, no one, which is why a large advocacy effort is needed. We need to hold Congress accountable! Let's make this hope a reality by uniting with a strong voice! www.ConsumerPainAdvocacy.org
@LynnRWebsterMD	A6. Actually we start now to ask when the report will be made public. Call your congressman or woman tomorrow. Plan now to write a letter to the editor of your paper, blog, or comment on a news article about the NPS when it is released.
@FibroandPain	Would including chronic pain in Essential Health Benefits be of benefit in the NPS initiative?
@BobTwillman	@FibroAndPain Need to use specific interventions that target each problem rather than broad interventions that hurt both.
@BobTwillman	@FibroAndPain That certainly should be a strategy that is part of the NPS.
@cat_fish_dinner	When care is decided by what's best for majority, access for individuals can be put at risk. What are the safeguards against that?
@BobTwillman	@cat_fish_dinner The NPS will say that we need to teach how to provide this type of individualized care to be successful.
@BobTwillman	@cat_fish_dinner The premise of the biopsychosocial model is that each individual needs an individual treatment plan.
@OneDayHeather	@BobTwillman, What is being done to conduct the needed research & case studies to validate ketamine to insurance companies?
@OneDayHeather	Can we post the needed the information online to send to Congress?
@BobTwillman	@OneDayHeather But it is SO important that we learn how to use tools we already have. As with many of these issues, research support is lacking for this.
@RubysFibroDiary	Is there a governing board which makes recommendations or requirements on what med schools teach?
@RubysFibroDiary	When released, will the NPS have hierarchies of implementation that need to be supported by this community?
@BobTwillman	@RubysFibroDiary ACGME--American Council of Graduate Medical Education.
@BobTwillman	Yes RT @RubysFibroDiary: When released, will the NPS have hierarchies of implementation that need to be supported by this community?
@CPRAlliance_org	@RubysFibroDiary Each of the six areas have short, medium and long-term objectives to implement in unison.
@melissarvh	@OneDayHeather @US_Pain @powerofpain shouldn't be difficult. Just need form letter & recipient list. Have done it w/other projects.
@BobTwillman	@RubysFibroDiary That remains to be seen. I think there will be parts of each section that are immediate implementation items.
@gnsalminen	They already are separate!! Should not be that hard to send this to Congress. What is an implementation team??
@CPRAlliance_org	@gnsalminen A committee that guides and ensures the NPS objectives are carried out!
@crystalgrieves	I had to pleasure to sit in on a call and give major feedback through the @US_Pain I hope for an implementation team. It's not near enough for research at all. Overlook by running a business only causes more for us to deal with.
@deanslewis	@AmerGastroAssn most of these diseases can cause chronic pain.

@forgrace	Fed Gov needs to embrace NPS. Until they say CP is public health problem #1, we will spin wheels
@lisadbudzinski	Everyone must be on the same page as @BobTwillman stated. http://painconsortium.nih.gov/federal_Pain_Activities/federal-pain-activities-index.html
@LynnRWebsterMD	@davidbecker19 Maybe but it does start a conversation. You can move it further once you have their ear. Run with it.
@LynnRWebsterMD	@tal7291 @FibroAndPain Loved you op ed. Write more.
@MIFibro	@CPRAlliance_org @BobTwillman Great news about Capitol Hill day!
@MIFibro	@davidbecker19 I have chronic pain & have been volunteering my time for 18 years to help people in chronic pain.
@ThePainComm	One of the biggest issues is getting access to care that works. If it's not covered, many can't afford the treatment they need.
@PAINSPROJECT	<p>Research is one of the key pieces of improving pain care</p>  <p style="text-align: right;">PAINSPROJECT</p>
@powerofpain	<p>Our federal government invests just \$0.05 per affected adult in pain research.</p> 

	Q7. What can someone do to get involved with the NPS now?
@CPRAlliance_org	A7. Visit www.ConsumerPainAdvocacy.org and join the effort! We need as many voices united and strong! Thank you!
@CPRAlliance_org	A7. A large united voice from every single American who has or knows someone about pain! www.ConsumerPainAdvocacy.org
@LynnRWebsterMD	A7. NPS gives us an opportunity to ask for a National Institute for Pain Medicine. I will be writing about this in an upcoming blog.
@LynnRWebsterMD	A7. Please commit to speak to 10 people about NPS when released. Make it a daily conversation & meet with your Senators
@PAINSPROJECT	A7: Stay informed. Sign up for alerts on www.ConsumerPainAdvocacy.org and its member organizations
@FibroAndPain	People are afraid of losing anonymity in speaking out about stigmatization and harmful practices. This has to stop!
@FibroAndPain	The NPS must address the access to care issues faced by minorities and underserved populations first.
@FibroAndPain	Would treating opioid addiction to the max help people with chronic pain have less stigmatization?
@BobTwillman	@FibroAndPain It has to help, at least in my estimation. And besides, that's also what we owe people with addictions.
@BobTwillman	@FibroAndPain True--they have it worse, but no one really has access to everything needed to provide the best care.
@OneDayHeather	@BobTwillman, I'm sad to hear that. Ketamine infusions have been offered for many years here & just recently getting ins. denials.
@BobTwillman	@OneDayHeather Yes, it is sad. Such an incredibly useful drug, and we don't know much of what it can do.
@CPRAlliance_org	@OneDayHeather @US_Pain @powerofpain Yes, visit www.ConsumerPainAdvocacy.org !
@cat_fish_dinner	@BobTwillman thanks. I hope it's applied that way. Seeing war on opioids is worrisome?
@lisadbudzinski	@melissarvh @OneDayHeather @US_Pain @powerofpain It would be great to have a form letter for delegates & advocates to send out.
@lisadbudzinski	Definition for biopsychosocial model: https://en.wikipedia.org/wiki/Biopsychosocial_model that is being mentioned for this chat.
@CPRAlliance_org	@lisadbudzinski @melissarvh @OneDayHeather @US_Pain @powerofpain This will be available on www.ConsumerPainAdvocacy.org Connect with us.
@MissionDrive	If we write blog posts, should we send or share them to a central place? Are there key talking points? What's the best way to get alerted when the NPS is released?
@CPRAlliance_org	@MissionDriven4 Yes & Sign up to receive alerts on www.ConsumerPainAdvocacy.org
@powerofpain	@MissionDriven4 - Follow along for updates at www.ConsumerPainAdvocacy.org
@crystalgrieves	@CPRAlliance_org @powerofpain @US_Pain I'm not stopping until my body does, I'll speak up as long as and loud as I can!
@crystalgrieves	Tough question, what has been the biggest roadblocks on this, and what else can others do to push through?
@crystalgrieves	@powerofpain @LynnRWebsterMD definitely on my to do list here in Maryland! I can't do it alone! We need more voices, mine is loud, but a group is powerful! #USPain #PAM15

@CPRAlliance_org	@crystalgrieves @powerofpain @US_Pain Thank you! We need everyone to speak up!
@melissarvh	@OneDayHeather @US_Pain @powerofpain Can even use site like Quorum. Example of our #Gastroparesis project: https://buildquorum.com/actions/1996
@melissarvh	@FibroAndPain I'm not afraid. :) I am willing to help.
@CPRAlliance_org	@melissarvh @OneDayHeather @US_Pain @powerofpain Not difficult. We need a unified voice and approach. Visit www.ConsumerPainAdvocacy.org
@powerofpain	Chronic pain costs the U.S. over \$500 billion every year #CPATF 

	Would you be interested in a future chat about NPS if yes, what? #CPATF
@mkwinnette	Treatments approved for different conditions. RT @powerofpain Would you be interested in a future chat about NPS if yes, what? #CPATF
@gnsalminen	We need Big Pharma to get into healing instead of just treating! I am sick of war on Opiates against Chronic Pain Patients!! I am willing to try and help.
@rsdcprfire	Yes. Q&A's on education to form solutions and partnerships. Chats provide opportunity for involvement.
	Wrap up
@powerofpain	Thanks to all participants & to our guest tweeters: @CPRAlliance_org @BobTwillman & @LynnRWebsterMD from #CPATF & @powerofpain
@powerofpain	Thank you to all participants in today's #CPATF #TwitterChat on the #NationalPainStrategy This has been an informative #PainMonth15
@powerofpain	Checkout @CPRAlliance_org www.cpralliance.org , @LynnRWebsterMD www.lynnwebstermd.com , @BobTwillman www.aapainmanage.org , & #CPATF www.ConsumerPainAdvocacy.org
@AAPainManage	Thanks to @powerofpain for hosting the about the #NationalPainStrategy! Very informative.
@BobTwillman	@OneDayHeather @powerofpain And thank you for being here!
@BobTwillman	@powerofpain Thank you for organizing--it's been great fun!
@CPRAlliance_org	Thanks for joining us in this important fight! Quality pain care for all! www.ConsumerPainAdvocacy.org
@LynnRWebsterMD	Thanks all for the conversation.
@US_Pain	@crystalgrieves Thanks for committing to do this!
@US_Pain	We wish to thank everyone who followed and participated in this evening's on Twitter! Working with... www.ConsumerPainAdvocacy.org
@crystalgrieves	@powerofpain @BobTwillman #PAM15 @ThePainComm @AAPainManage thank you all for your time!
@deanslewis	@powerofpain very informative, let's continue to work on helping others realize that it's ok to be open about their pain. We lose too many souls.
@deanslewis	@powerofpain let's work on helping others realize that it's ok to be open about their pain. We lose too many souls.
@forgrace	We need a potent voice in Congress to take NPS lead - one not beholden to special interest/industry. I know - "Good luck!"
@forgrace	Great chat everyone. We. Can. Do. This.
@gnsalminen	@crystalgrieves, if I can help you I will. <3 Love to ALL!! <3 Good night!!
@lisadbudzinski	@CentralPain is also on board to making a difference
@melissarvh	Thank you for hosting this most informative chat on #Chronicpain this evening. Looking forward to future discussions & projects!
@MIFibro	Yes, thanks to all organizers and presenters!
@mkwinnette	Thank you @powerofpain
@opiophobia	@BarbyIngle @kypaincare @powerofpain @BobTwillman @LynnRWebsterMD Thank all of you for giving us a voice, I didn't realize how BIG this is
@PAINSPROJECT	Thanks to all participants & to our guest tweeters: @CPRAlliance_org @BobTwillman & @LynnRWebsterMD, @powerofpain
@rsdcprfire	Thank you @powerofpain @BobTwillman and all individuals and advocacy groups/orgs who participated this evening.
@SabrinaNumann	Thank you to @powerofpain, @BobTwillman for this chat! Together we can make a difference! @FibroAndPain

@ThePainComm	Thanks to @powerofpain for hosting the about the #NationalPainStrategy! Very informative.
--------------	---